

Malawi Standard Treatment Guidelines (MSTG) 5th Edition 2015

Incorporating Malawi Essential Medicines List (MEML) 2015

Ministry of Health



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Ministry of Health

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Table of Contents

| Forev | vord | iii |
|-------|-------------------------------------|-----|
| Refer | ences | V |
| Ackno | owledgements | vii |
| Presc | ribing Guidelines | xi |
| 1.0 | Blood and Haematological Conditions | 1 |
| 2.0 | Cardiovascular diseases | 47 |
| 3.0 | Central nervous system | 62 |
| 4.0 | Ear nose and throat conditions | 95 |
| 5.0 | Emergencies | 106 |
| 6.0 | Endocrine disorders | 121 |
| 7.0 | Gastro-intestinal conditions | 134 |
| 8.0 | Hepatic disorders | 161 |
| 9.0 | Infectious diseases | 162 |
| 10.0 | Oncology | 225 |
| 11.0 | Musculoskeletal disorders | 246 |
| 12.0 | Sexual and Reproductive Health | 257 |
| 13.0 | Ophthalmic conditions | 304 |
| 14.0 | Oral maxillofacial conditions | 320 |
| 15.0 | Parasitic diseases | 333 |
| 16.0 | Respiratory conditions | 371 |
| 17.0 | Sexually transmitted infections | 393 |
| 18.0 | Skin conditions | 417 |
| 19.0 | Vaccinations | 437 |
| | ———— MSTG 2015 ————— | |

| 20.0 | Bites, burns and wounds | .439 |
|-------|-------------------------------------|------|
| 21.0 | Renal conditions | .454 |
| 22.0. | Poisoning | .465 |
| 23.0 | Nutritional Disorders | .475 |
| 24.0 | Pain management and Palliative care | .504 |
| 25.0 | Psychiatric condition | .520 |

Foreword

The purpose of the Malawi Standard Treatment Guidelines (MSTGs), which incorporates the Malawi Essential Medicines List (MEML) is to standardise prescribing patterns and practices, thereby also enabling a more consistent and uniform range of available medicines and medical supplies across all levels of the national health care system. This in turn helps in quantification, procurement and supply of national requirements of medicines and medical supplies.

The 5th Edition of the MSTGs which has new added chapters in fields such as oncology has, therefore, been developed to meet the needs of prescribers at different levels, in selecting and prescribing the right medicines, realising that "memory alone can sometimes be treacherous" and even the most thoroughly informed prescribers need a reminder "to make them masters of the situation".

It must be mentioned though that medicine is a very complex and dynamic field with emerging and re-emerging clinical conditions, hence as is always the case in clinical practice, professional knowledge and judgement is of paramount importance, and where necessary, this should be supplemented by specialised publications on specific clinical conditions as well as reference to relevant product literature.

It is the wish of the Ministry of Health that prescribers both in public and private health sector will make the most use of this 5th Edition of the Malawi Standard Treatment Guidelines in their time of duty.

C.V. Kang'ombe

SECRETARY FOR HEALTH

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The following are national guidelines or reference text which should be consulted for further information on specific areas or topics:

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Prescribing Guidelines

1. General points

Consider each of the following general points before writing a prescription:

1.1 Not all patients need a prescription for medicines. Non-medicine treatments and/or giving of simple advice may be more suitable in certain situations.

1.2 Good therapeutics practice depends on:

- Accurate diagnosis, based on thorough history-taking, necessary careful physical examination and, if required, supporting laboratory testing
- Knowledge of the medicines available
- Careful selection of the appropriate medicines
- Prescribing correctly the selected medicines
- Ensuring that the patient understands fully how to use each prescribed medicine properly
- 1.3 Try to resist patient demand to prescribe injections or other expensive dosage forms. e.g. capsules and oral liquids.

Always make an effort to explain to the patient that these may not represent the best form of treatment for the particular condition

- 1.4 In life threatening conditions, always prescribe the *most effective* medicine available irrespective of the cost or limited availability
- 1.5 In order to avoid possible confusion, always prescribe medicines by their generic name and not by the brand name e.g. diazepam (not Valium), paracetamol (not Panado) or abbreviations i.e. PCM
- 1.6 Avoid prescribing combination medicines unless they have a known significant therapeutic advantage over single ingredient preparations
- 1.7 When prescribing any medicine, always take into consideration factors such as:
 - Patient's age
 - Patient's sex
 - Patient's weight
 - The effect of other diseases present
 - Pregnancy
 - Breast-feeding
 - The likely degree of patient compliance with treatment
- 1.8 In all cases the likely benefit of any prescribed medication/s must be weighed against potential risks

- 1.9 Avoid overuse of symptomatic treatments for minor self-limiting conditions
- 1.10 Avoid multiple prescribing (polypharmacy), especially when the diagnosis is not clear

2. Prescribing of placebos

- 2.1 Avoid this whenever possible. Instead spend time reassuring and educating the patient
- 2.2 If it is absolutely necessary to prescribe a placebo, always choose a safe, cheap medicine which is not essential for the treatment of other important conditions, e.g. multivitamin tablets or vitamin B compound tablets
- 2.3 Never prescribe injections as placebo
- 2.4 Never prescribe tranquilizers e.g. diazepam, phenobarbitone, as placebos

3. Prescription writing

Note: Whenever possible, return all incomplete, inaccurate, illegible or unclear prescriptions to the prescriber for clarification, completion, or correction, before they are presented for dispensing

- 3.1 Write all prescriptions legibly in ink. Poor writing may lead to errors in interpretation by the dispenser which may have harmful and possibly disastrous consequences for the patient
- 3.2 Write the full name and address of the patient, and sign and date the prescription form
- 3.3 Write the name of the medicine or preparation using its full generic name.

 Do not use unofficial abbreviations, trade names, or obsolete names as these may cause confusion
- 3.4 Always state the strength of the preparation required where relevant
- 3.5 For solid dosage forms:
 - Quantities of one gram or more should be written as 1g, 2.5g, 10g, etc
 - Quantities of less than one gram but more than one milligram should be written as milligrams rather than fractions of a gram, e.g. 500mg and not 0.5g
- 3.6 Quantities less than one milligram should be expressed as micrograms (in full) and not as fractions of a milligram, e.g. 100 micrograms rather than 0.1 mg or 100mcg.

- 3.7 If decimals are used, always write a zero in front of the decimal point where there is no other figure, e.g. 0.5mL and not .5mL
- 3.8 Always state the full dose regimen, i.e.
 - Dose size
 - Dose frequency
 - Duration of treatment

The quantity to be dispersed will be deduced from this.

3.9 Avoid use of the direction "to be used/taken as required". Instead state a suitable dose frequency. In the few cases where 'as required' is appropriate, the actual quantity to be supplied should be stated

3.10 Avoid using unknown abbreviations. The following abbreviations can be used when writing a prescription:

List of Abbreviations

a.c before meal
b.d twice a day
IM Intramuscular
IV Intravenous
mane in the morning
n et m or n.m. night and morning

nocte at night

o.m. every morning
o.n. every night
p.o. by mouth
p.c. after meals

prn when

necessary

q4h
q6h
q8h
q.i.d
every 4 hours
every 6 hours
every 8 hours
4 times a day

stat immediately or at

once

sig label

t.d.s. 3 times a day

- 3.11 For oral liquids, doses should be stated in terms of 5mL spoonfuls for linctuses, elixirs, syrups and paediatric preparations, and in 10mL spoonfuls for adult mixtures
- 3.12 Doses other than 5mL or 10mL or multiples of these will be diluted to the nearest equivalent 5mL or 10mL quantity for dispensing
- 3.13 Total volumes of liquid preparations prescribed are usually selected from 50, 100, 300 or 500mL volumes
- 3.14 Total quantities of solid or semi-solid preparations prescribed are usually selected from 25, 50, 100, 200, 300, or 500g except where the product is supplied ready packed in a particular pack size, e.g. tetracycline eye ointment (3.5g)
- 3.15 Where relevant, always remember to include on the prescription any special instructions necessary for the correct use of a medicine or preparation, e.g. "before food" etc.

4. In-patient prescriptions

4.1 Write these prescriptions and records of dispensing and administration on inpatient treatment cards

- 4.2 Only use one card per patient at any one time
- 4.3 Clearly state a suitable dose frequency, or time of administration on medicines to be given 'as required'
- 4.4 Always state the route of administration for all medicines prescribed
- 4.5 When any changes or cancellations are made to a prescription card, or if treatment is to be stopped, clearly sign and date the card in the appropriate place
- 4.6 If the timing of a medicine dosage is critical, ensure that suitable arrangements are made for the medicine to be given at the specific time/s required

5. Guide to quantities to be supplied

5.1 Oral liquids

Adult mixtures (10 mL dose)

200mL (20 doses)

300mL (30 doses)

Elixirs, linctuses and paediatric mixtures (5mL dos

50mL (10 doses)

100mL (20 doses)

150 mL (30 doses

5.2 **Preparations used in body cavities**

E.g. ear drops, nasal drops

– MSTG 2015 –

5.3 External preparations

| Part of body | Semi-solid (g)* | Liquids (mL)** |
|--------------|-----------------|----------------|
| Face | 5-15 | 100 |
| Groin and | 15-25 | 100 |
| genitalia | 25-50 | 200 |
| Both hands | 50-100 | 200 |
| Scalp | 100-200 | 200 |
| Both arms | 200 | 500 |
| and legs | | |
| Whole body | | |
| | | |
| | | |

^{*} E.g. creams, pastes, ointments etc **e.g. lotions, applications, topical solutions, etc (for paints normally 10-25mL is supplied)

- 6. Prescriptions for controlled medicines
- 6.1 These medicines are controlled by the Laws of Malawi, The Pharmacy Medicines and Poisons Act, 1988. Consult the relevant sections of the Act for details of the appropriate legal requirements in **each case**
- 6.2 Medicines covered by the Act and which are also used in the MSTG are:
 - Morphine sulphate injection
 - Morphine sulphate solution
 - Pethidine hydrochloride injection
 - Morphine sulphate tablets
- 6.3 These medicines have potential for abuse which may be result in dependence. Carefully record all procedures involving them in the appropriate record books
- 6.4 Prescriptions for these medicines may only be written by registered medical practitioners

- 6.5 The following legal requirements must also be observed when writing such prescriptions:
 - a) The prescription must be in the prescriber's own handwriting
 - b) It must be signed and dated
 - c) The prescriber's address must be shown
 - d) The name and address of the patient must be stated
 - e) The total amount of the item to be supplied must be stated in words and figures
 - 6.6 It is an offence for the prescriber to issue and for the pharmacy/dispensary to dispense prescriptions for controlled medicines, unless the requirements of the law are fully complied with

Notes:

a) In certain exceptional circumstances, senior nurses in charge of departments, wards, or theatres, and midwives, may also obtain and administer certain controlled medicines as part of their work.

The relevant sections of the Act should be consulted for the details of the appropriate legal requirements in each case

b) Hospital in-patient prescriptions for controlled medicines should be prescribed on a separate prescription as well as written on treatment cards or case sheets and signed/dated by the person administering the medicine.

7. Adverse drug reactions (ADRs)

- 7.1 Nearly all medicines may produce unwanted or unexpected adverse effects, some of which may be life threatening e.g. anaphylactic shock, liver failure
- 7.2 Prescribers should immediately report any serious or unexpected adverse effects thought to be due to a medicine to:

The Registrar,
Pharmacy, Medicines and Poisons Board,
PO Box 30241, Lilongwe.
Tel: 01 755 165/166 Fax: 01755 204

- 7.3 Rules for prevention of ADRs
 - a) Never use a medicine unless there is a clear indication for its use
 - a. Only use medicines in pregnancy if absolutely essential

- b. Check if the patient has had any previous reactions to the medicine or to similar medicines
- Remember to reduce doses when necessary e.g. in the young, the elderly, and if liver or renal disease is present
- d. Always prescribe the minimum number of medicines possible
- e. Carefully explain the dose regimens to patients, especially those on multiple medicines, the elderly and anyone likely to misunderstand
- f. If possible, use medicines with which you are familiar
- g. Look out for ADRs when using new or unfamiliar medicines
- h. Warn patients about likely adverse effects and advise them on what to do if they occur
- i. Patients on certain prolonged treatments e.g. Anticoagulants, corticosteroids, Insulin etc. should carry a small card giving information about the treatment

8. Paediatric prescribing

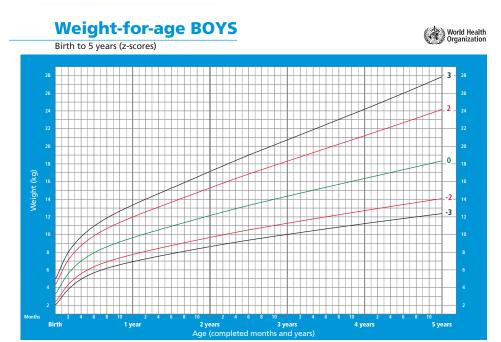
8.1 In these guidelines, paediatric medicine doses are usually given according to body weight and not age, and are therefore expressed as mg/kg etc.

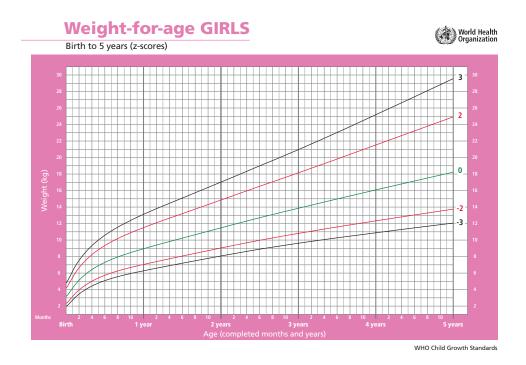
Prescribing guidelines

The main reason for this is that children of the same age may vary significantly in weight.

Thus it is safer and more accurate to prescribe drugs according to body weight. Moreover, this should encourage the good practice of weighing children whenever possible.

8.2 The following graphs shows weights of children aged 0 to 5 years and may be used to determine whether the child is well nourished, underweight as well as overweight for their specific ages.





Five lines are shown on each graph

- The middle green line shows weight for average children with a z score of zero
- The lower red line shows weights for children who are moderately underweight for their age with a z score of -2
- The lower black line shows weights for children who are severely underweight for their age with a z score of -3
- The upper red line shows weights for children who are moderately overweight for their age with a z score of +2
- The upper black line shows weights for children who are severely overweight for their age with a z score of +3
- 8.3 When a weighing scale is not available the following equ ation can be used to estimate the weight of the child: [Age (in years) + 4] x 2
- 8.4 Neonates have delayed hepatic and renal excretion of medicines and also unpredictable absorption of oral medication. Therefore give special consideration when prescribing for children less than 30 days old and especially premature infants.

9. Medicine interactions

- 9.1 Whenever prescribing a particular medicine, care should be taken to avoid problems of interactions with other medicines, whether these are:
 - also prescribed at the same time
 - previously prescribed by another prescriber for the same or another condition and currently being taken by the patient
 - purchased or otherwise obtained by the patient for the purpose of self-medication
- 9.2 Thus, before prescribing a medicine, always obtain details of any other medication currently being taken by the patient
- 9.3 Where a medicine interacts with alcohol (e.g. metronidazole, diazepam, anti-diabetic medicines, tricyclic antidepressants etc.) remember to counsel the patient to avoid taking alcoholic drinks during the course of treatment and for at least 48 hours after completion of the course

Presentation of Information

a. Arrangement of sections

Standard treatments have been grouped in sections according to either body systems (e.g. respiratory conditions, gastrointestinal conditions, etc.) or types of disorder (e.g. parasitic diseases, nutritional disorders, etc.) Use the table of contents, page ii, to locate the particular section required.

b. Indexing and Cross-referencing

All diseases, conditions, tables, etc. are included in an index. Extensive cross-references are given in the text, by section number and page number, to facilitate location of other references to the subject elsewhere in the guidelines. Use the Index on page 201-208 to quickly find the required subject.

Prescriber's guidance points

- These are given for most standard treatments and are key points to be considered before prescribing for a patient with a particular condition.
- Certain points as well as warnings are given added emphasis by inclusion in a boxed border.

Medicine administration

- Unless otherwise specified, the oral route is to be used. Even when a parental route is specified, with medicines which are well absorbed orally and which are available as an oral dosage-form, it is often possible to switch to oral administration once the patient has improved and is able to swallow/tolerate oral medication
- Additional guidance on medicine administration is given, where relevant, as bulleted points after dosage regimen.

Medicine names

 Medicines recommended for use are those on the current Malawi Essential Medicine List, 2014.
 Generic names are used and indicated in **bold type**. Where necessary, proprietary names are indicated in *italic* type.

Alternative medicines

- These are indicated where appropriate and available for alternative treatment of a particular condition. They should be used only if the recommended medicine is not available or is not suitable for a particular patient.
- In some cases (where indicated)
 alternative (i.e. 2nd line) medicines
 may be used when a satisfactory
 response has not been obtained
 with the recommended (1st line)
 medication.

Abbreviations

ARI Acute Respiratory Infections

ART Anti-Retroviral Therapy
BF Blood Film examination

BP Blood Pressure

COC Combined Oral Contraceptive

CSF Cerebrospinal Fluid

CVA Cerebrovascular Accident
CPAP Continuous Positive Airway

Pressure

CXR Chest X-ray

DIC Disseminated Intravascular

coagulopathy

DNS Dextrose Normal Saline

FBC Full blood count

FFP Fresh Frozen Plasma

g gram

Hb haemoglobin

HIV Human immunodeficiency Virus

IM IntramuscularIV Intravenous

IU International units

JVP Jugular venous pressure

Kg Kilogram

L Litre

LP Lumbar puncture

LRTI Lower Respiratory Tract Infection

mg milligram
mL milliliter
mmol millimole

MU mega (1 million) units

Abbreviations

NGT nasogastric tube PCV packed cell volume

s/c subcutaneous

STI sexually transmitted infections

TB Tuberculosis

TTP Thrombotic thrombocytopenic

purpura

URTI upper respiratory tract infection

Metric Units

1 kilogram (kg) = 1,000 grams (g)

1 g = 1,000 milligrams (mg)

1 mg = 1,000 micrograms

1 litre (L) = 1,000 millilitres

1 ml of water = 1 g

1% (m/v) = 10 mg/mL

Equivalents

1 litre = 1.8 pints

1 pint = 568.3 mL

1 kg = 2.2 pounds

1 lb = 453.4 g

1 ounce (oz) = 28.35 g

1.0 Blood and Haematological Conditions

1.1 Blood: Guidelines for Appropriate Use

Refer to the Ministry of Health Guidelines for Safe Blood Transfusion for further details on:

- Donor recruitment and selection,
- Blood collection,
- Storage procedures and records,
- Laboratory testing of donor and recipient's blood
- Refer to Guidelines for the Clinical Use of Blood and Blood Products in Malawi for Clinical aspects of blood transfusions and administration Transfusion reactions
- Blood transfusion, although having undoubted benefits, also carries serious risks including:
 - Possible transfusion of infections (e.g. HIV and hepatitis)
 - Immune-system related problems (e.g. Intravascular haemolysis)
 - Circulatory overload
- It is expensive and uses a scarce human resource, therefore only prescribe blood if:
 - Less hazardous therapy has been or will be ineffective, and
 - > The benefits outweigh the risks involved

- The decision to transfuse blood has been based on careful assessment of the patient which must indicate that it is necessary to save life or prevent major morbidity
- Except in the most exceptional lifethreatening situations, always transfuse blood which has been obtained from appropriately screened blood donors and/or appropriately screened for infectious agents.
- Ensure that compatibility testing is carried out on all blood to be transfused. In absolute emergencies, where there is no time for emergency cross-matching, uncrossmatched blood can be issued but a crossmatch should still be done while the transfusion is in progress.
- Observations of patient's vital signs should be done at the time of starting the transfusion, at 15 minutes, 1 hour, 4 hours and at 24 hours

1.2 Indications for transfusion of whole blood or red cell suspension

1.2.1 Severe Anaemia

Neonates

- Hb < 12 g/dl (PCV < 36%) in the first 24 hours of life.
- Hb < 12 g/dl (PCV <36%) in a neonate receiving mechanical ventilation.
- Hb 8-11 g/dl (PCV 24-33%) and oxygen dependent.
- Hb < 7 g/dl (PCV < 21%) when stable and off oxygen.

Children and infants

- If Hb < 4 g/dl or PCV <12
- If Hb < 6 g/dl or PCV <18 with any of the following:
 - Shock or clinically detectable dehydration
 - Impaired consciousness
 - Respiratory acidosis (deep laboured breathing)
 - Heart Failure
 - Requiring oxygen for any reason

Dose

- Transfuse 20ml/kg of whole blood or 10ml/kg of red cell suspension
 - A diuretic is usually not indicated because many of these children are usually hypovolemic with a low blood volume.
 - Check the respiratory rate and pulse rate every 15 minutes and if one of them rises, transfuse more slowly.
 - If there is evidence of fluid overload due to blood transfusion, give Frusemide (1-2mg/kg) up to a maximum total of 20mg.
 - If severe respiratory distress, consider oxygen therapy with or without CPAP as appropriate.
 - After the transfusion, if the Hb remains low, repeat the transfusion
 - In severely malnourished children fluid overload is common and difficult to recognize. Give 10mls/kg rather than 20ml/kg of whole blood once only and do not repeat the transfusion.
 - Figure 6 Give a daily iron-folate tablet or 1 Iron syrup for 14 days (Iron must not be given when acutely unwell, and in rehabilitation phase of malnutrition).

1.2.2 Non severe Anaemia in children

Children

Young children (less than 6 years) are anaemic if their Hb is less than 9 g/dl. Begin treatment unless the child has severe malnutrition, in which case, refer to malnutrition chapter.

- Give (home) treatment with iron (daily iron-folate tablet or dose of iron syrup) for 14 days
- Ask the parent to return with the child in 14 days. Treat for 3 months when possible, as it takes 2-4 weeks to correct anaemia and 1-3 months to build up iron stores
- If the child is ≥ 1 year and has not received Mebendazole in the previous 6 months, give one dose of Mebendazole (500mg) alternatively Albendazole (200mg or 400mg depending on age) for possible hookworm or whipworm infestation
- Advise the mother about good feeding practice

Pregnancy

Refer to Reproductive Health (Chapter 12)

Adults

- If Hb less than 5 g/dl
- If Hb less than 8 g/dl and there are clinical complications

Dose

 One unit of whole blood or one unit of red cell suspension will raise a patient's haemoglobin by 1-1.5g/dl

Pre-operative Surgery

• If Hb less than 8 g/dl

Red Flags

- Do not transfuse in megaloblastic anaemia (MCV≥110)
- If suboptimal rise or fall in haemoglobin level after transfusion and there are signs of haemolysis (such as jaundice, raised bilirubin level), refer the patient for specialist management)

Vitamin B12 deficiency

 Transfusion should be avoided unless the patient has symptomatic anaemia and even then, the minimum possible amount of blood or red cell suspension should be transfused e.g. one paediatric unit for an adult patient.

1.2.3 Acute Haemorrhage with shock (see section 1.5, table 1)

1.2.4 Intra-operative use (where necessary)

Note: Do not use whole blood or red cell suspension transfusion to expand blood volume

1.3 Platelets

Must be transfused immediately upon arrival. Platelets should never be stored in a refrigerator or blood bank or in the ward.

 Decision to transfuse should be based on a combination of clinical and laboratory findings rather than empirical platelet levels.

1.3.1 Indications for platelets use

- Bleeding due to thrombocytopenia as a result of defective platelet production such as aplastic anaemia or leukaemia
- Increased consumption e.g. DIC
- Dilutional effects e.g. in massive transfusion
 Note: All patients needing platelets
 need further investigations, please
 refer to Central Hospital.

Dose

- 1 unit per 10kg
- For infants under 10 kg, 5ml/kg

1.4 Fresh frozen plasma

- Contains all clotting factors
- Comes in volumes of 200-300 mls
- Fresh frozen plasma (FFP) should be thawed before use using water bath at 30-37 degrees (If water bath is not available, a plastic basin with lukewarm water or cold tap water can be used). Never use hot water

Note: Do not use whole blood, fresh frozen plasma or red cell suspension to expand blood volume

- Once thawed, FFP must be used immediately. FFP must never be refrozen.
- The patient needs further investigations; please refer to the central hospital.

1.4.1 Indications for use of fresh frozen plasma

- Replacement of single factor deficiencies (if single factor concentrates are not available)
- Immediate reversal of warfarin effect
- Vitamin K deficiency associated with active bleeding
- Acute disseminated intravascular coagulopathy
- Thrombotic thrombocytopenic purpura (TTP)
- May be used in massive transfusion or liver disease

Dose

15-20ml/kg

Note: No justification for use in hypovolaemia, nutritional support in protein losing states or/and plasma exchange except in TTP.

1.5 Acute Haemorrhage

- In massive haemorrhage i.e. from trauma it is difficult to estimate how much blood a patient has lost. However a good estimate can be made by calculating the patient's normal circulating volume versus vital signs and other organ function tests. See Table 1 below.
- Restoration of blood volume with suitable replacement fluids is more important than red cell replacement in the management of previously healthy patients who have lost under 30% of their blood volume
- The need for blood transfusion must be determined by:
 - The amount and speed of blood loss
 - The patient's vital signs

Table 1: Assessment of Blood Loss (For a 70 kg adult)

| | Stage 1 | Stage 2 | Stage 3 | Stage 4 |
|-----------------------------------|---------------|------------|---------------------------------|------------------------------|
| Blood loss (litres) | <0.75 | 0.75-1.5 | 1.5-2 | >2 |
| Pulse rate | <100 | >100 | >120 | >140 |
| ВР | Normal | Normal | 90/60 | <70 |
| Respira- tory Rate | <20 | >20 | >30 | >40 |
| Capillary Refill | <3 seconds | <3 seconds | >3 seconds | >3 seconds |
| Mental State | Normal | Anxious | Confuse d | Lethargic |
| Urine output/ Hour | >30 mls | 20-30 mls | <20 mls | <10 mls |
| Replaceme -nt fluid vol (L) | 2L | 2-4.5L | >5L plus 2 units blood | >6L plus 3 units Blood |

- Replacement fluids which may be used are:
 - ➤ Haemacel: Replace every 1 ml of blood lost with 1 ml of fluid
 - Sodium lactate compound
 (Ringers lactate) IV infusion
 or Normal saline IV infusion
 Replace 1 ml of blood lost
 with 3 mls of fluid.

Do not use dextrose 5% or Darrow's ½ strength in dextrose 5% as replacement fluids

- Maintain the airway and give oxygen by face mask first, especially for patients in stage 3 and 4. Make sure they are breathing adequately.
- Insert 2 large bore cannulae (gauge 14 or 16) and collect blood samples for full blood count (FBC), grouping and crossmatching.
- Give half of the calculated dose of replacement fluid in the first hour and give the other half over 3 hours.
- Always assess the effects of fluid therapy.
 Remember to give warm fluids and cover patients to avoid hypothermia.

 Aim at improving oxygen carrying capacity first before correcting anaemia.
 Remember to add maintenance fluids to the replacement fluid plus any on-going losses.

Note: Maintenance fluids can be calculated as follows:

- (i) Adults: Body weight x 1.5mls
- (ii) *Children*: May use the rule of 4.2.1 for children or refer to section on diarrhoea. (Section 7.5 page 42)
- Remember: deficit + maintenance + ongoing loss

1.6 Adverse Reactions to Transfusion

- Suspect an adverse reaction if any of the following occurs:
 - Severe pain at transfusion site or in the back, loin and/or chest
 - Rise in temperature of 1 C above the baseline
 - Increase in pulse rate of >20/minute above baseline
 - ➤ Fall in systolic BP >20 mm Hg
 - Urticaria
 - Rigors
 - > Haemoglobinuria
 - Shortness of breath
 - Wheezing
- Treatment depends on the severity of the transfusion reactions.

1.6.1 Mild reaction

Signs and symptoms: itchy rash

Possible cause: hypersensitivity (mild)

Treatment:

- Slow the transfusion
- Administer antihistamine IM/IV or PO
- Chlorpheniramine 0.1 mg/kg IM or IV for children; 10mg IM/IV for adult alternatively
 Chlorpheniramine 4mg PO or
- **Promethazine** 6.25-12.5mg for children aged 5-12 yrs and 25mg for adults.
- If no clinical improvement within 30 minutes or if signs and symptoms worsen, treat as Category 2.

1.6.2 Moderate Reaction

Signs and Symptoms:

 Anxiety, pruritus, palpitations, mild dyspnea, headache, rigors, fever, tachycardia

Possible causes:

- Hypersensitivity (moderate to severe)
- Febrile non-haemolytic transfusion reaction
- Contamination with pyrogens and / or bacteria.

Treatment

- Seek help immediately from the anaesthetic, emergency team or whoever is available and skilled to assist
- Stop the transfusion
- Replace the infusion set and keep IV line open with normal saline.
- Administer antihistamine IM/IV or PO
 - Chlorpheniramine 0.1 mg/kg IM or IV for children; 10mg IM/IV for adult or Chlorpheniramine 4mg PO.
 - Promethazine 6.25-12.5mg for children aged 5-12 yrs and Promethazine 25mg for adults.
- Give oral or rectal antipyretic (e.g.
 Paracetamol 10 mg/kg or 0.5g 1g in adults). Never give Aspirin.

- Give IV corticosteroids (e.g. Hydrocortisone 200mg IV stat) and bronchodilators (e.g. Aminophylline 100mg stat) if there are anaphylactoid features (e.g. bronchospasm, stridor).
- If there is clinical improvement, restart transfusion slowly with new blood unit and observe carefully.
- If no clinical improvement within 15 minutes or if signs and symptoms worsen, treat as Category 3.

1.6.3 Life-threatening Reaction

Signs and symptoms

 Anxiety, chest pain, pain near infusion site, respiratory distress, loin or back pain, headache, dyspnea, rigors, fever, restlessness, hypotension, tachycardia, haemoglobinuria, unexplained bleeding

Possible causes:

- Acute intravascular haemolysis
- Bacterial contamination and septic shock
- Fluid overload
- Anaphylaxis
- Transfusion Associated Acute Lung Injury (TRALI)

Treatment

 Seek help immediately from the anaesthetist, emergency team or whoever is available and skilled to assist.

- Stop the transfusion. Replace the infusion set and keep IV line open with normal saline.
- Infuse normal saline (initially 20- 30 ml/kg) to maintain systolic BP, if hypotensive, give over 5 minutes and elevate patient's legs.
- Maintain airway and give high flow of oxygen by mask.
- Give Adrenaline (as 1:1000 solutions) 0.01 mg/kg body weight by slow intramuscular injection.
- Give IV corticosteroids (Hydrocortisone 200mg IV stat) and bronchodilators (aminophylline 100mg stat) if there are anaphylactoid features (e.g. bronchospasm, stridor).
- Give a diuretic: e.g. Frusemide 1 mg/kg IV or equivalent (if there is fluid overload)
- Assess for bleeding from puncture sites or wounds. If there is clinical or laboratory evidence of a DIC treat accordingly.
- Maintain fluid balance accurately
- If bacteraemia is suspected (rigors, fever, collapse, no evidence of a haemolytic reaction), start broad-spectrum antibiotics IV and send blood product bag to the laboratory for culture of contents.

1.6.4 Alternatives to Blood Transfusion

- There is no fluid that has similar properties as blood which can be used instead of blood.
- IV Iron and Erythropoietin which can be used as substitutes to blood transfusion in chronic anaemia are currently not available in our health facilities.
- Acute anaemia due to bleeding or haemolysis cannot be adequately treated by giving these products.
- When available, cell salvage machines can be used in acute anaemia due to bleeding.

1.7 Anaemia

1.7.1 Sickle Cell Anaemia

Wide spectrum of disease severity

- Suspect if patient is chronically anaemic and/or received previous blood transfusions and also those with family history of siblings/relatives with multiple blood transfusions
- Suspect in children who have suffered from one or more of the presentations listed below. For sickle cell test, refer for further investigations and management

Sickle cell Disease presentations

| Presentation | Age | Management | Differential |
|---|---------------|---|---------------------------------|
| Hand foot syndrome- hand foot swelling early infancy | 0.5 – 2 yr | Analgesics and hydration | Septic arthritis |
| Stroke/ CNS complications | 5- 10yr | CT/MRI scan, treat seizures | Cerebral malaria/ abscess |
| Bone Infarct | 0.5- 10yr | Xray, analgesia, rest, ortho referral | Osteomelitis |
| Splenic sequestration | <3yr | O2, treat shock, Benzylpenicillin, Gentamycin, transfuse | Acute abdomen |
| Acute Chest | 0.5- 10yr | O2, Antibiotics, Xray | Pneumonia |
| Bowel infarct/Abdominal crisis | | O2, antibiotics, surgical rv | Acute abdomen/ cholelithiasis |
| Priapism- sudden painful onset of penis that fails to relax | 6- 20yrs | Surgical assessment if not relaxing for more than 4 hours | |
| Kidney infarct | | Urinary output, electrolytes and BP | Nephritic syndrome |
| Skin Ulcers | >10yrs | GV paint/ antibiotics ointments/ analgesics | Kwashiorkor |
| Eye disorders | >10yrs | Ophthalmologist review | Other causes of low vision |

Treatment

For severe anaemia, refer to section 1.2.1

- If shocked, consider splenic sequestration (20ml/kg normal saline – see shock protocol)
- Transfuse in acute chest syndrome or stroke
 - Consider BTF program/ Hydroxyurea 15mg/kg daily then increase by 5mg/kg every 12th week to max 35mg/kg daily

For painful and vasoocclusive crises

- All children presenting with sickle cell crises should be started on oxygen even if saturations are normal
- Per every 24hrs, give 1.5 times maintenance fluid IV
- Encourage lots of oral hydration
- Give analgesics (see palliative care section)

For Infection and fever

- Sepsis usually caused by S pneumonia, give
 Benzylpenicillin and Gentamicin
- Meningitis usually caused by S pneumonia, give Ceftriaxone for 2 weeks
- Acute Chest syndrome, give Ceftriaxone and Erythromycin
- Salmonella osteomyelitis: Ceftriaxone IV (2-6 weeks), then oral Ciprofloxacin

Always check for malaria parasites

Others

 Cholelithiasis/Cholecystitis (>10yrs) AXR, needs abdominal ultrasound and surgical referral

Discharge and follow up

- Ensure being followed up in hospital clinic for the following:
 - Malaria prophylaxis
 - Monthly Sulphadoxinepyrimethamine (SP)
 - <4yrs ½ tab</p>
 - 4-8yrs 1 tab
 - 9-14yrs 2tab
 - >14yrs 3tab
- Alternatively use weekly Chlroquine 5mg/kg
- Folic acid 1-5mg daily
- Benzathine Penicillin (>6months) monthly
 IM

> <30KG: 600000IU

>30KG: 1.2MU

- Pneumococcal vaccine (2 and 5yrs) if available
- Consider Hydroxyurea for those with frequent painful crises
- DO NOT GIVE FERROUS SULPHATE

- Educate patient and family to ensure early analgesics, and to promptly seek medical attention if
 - > severe pain
 - fast breathing
 - looking much more pale than usual
 - high temperature
 - vomiting and diarrhoea

1.7.2 Anaemia, Aplastic

 There is pancytopenia due to a hypoplastic bone marrow.

Signs and Symptoms

 Pallor, petechiae, purpura, and bleeding with frequent or severe infections.

Treatment

- If neutropenic and febrile.
- Stabilise patient, if necessary, with blood products before referral.
- Refer patient

Referral

 Discuss all cases of suspected aplastic anaemia with a haematologist/specialist.
 Stabilise patient, if necessary, with blood products before referral.

Note: Discuss all cases of suspected aplastic anaemia with a haematologist/specialist.

1.7.3 Anaemia, Chronic Disorder

- Anaemia due to chronic inflammation.
- This is characteristically a normochromic normocytic anaemia.
- Common causes of anaemia of chronic disorder include: malignancy, e.g. haematological or solid tumours, autoimmune disorders, e.g. rheumatoid arthritis, acute or chronic infections, e.g. HIV and TB, chronic kidney disease, and chronic rejection of solid-organ transplantation, etc.

Treatment

- Treat the underlying condition.
- Transfusion is seldom necessary.
- Do not treat with iron, folic acid or vitamin B12 unless there is a documented deficiency.

1.7.4 Anaemia, Haemolytic

Causes

- Anaemia due to destruction of red blood cells. Destruction may be due to:
- Extracellular factors such as auto-immunity or mechanical factors, e.g. disseminated intravascular coagulation (DIC), hypersplenism, medications.
- Abnormalities of the cell membrane, e.g. hereditary spherocytosis.
- Enzymes, e.g. G6PD deficiency.
- Haemoglobin, e.g. sickle cell anaemia, thalassaemia.

Investigations

- Evidence of haemolysis: anaemia, reticulocytosis, decreased haptoglobin, increased lactate dehydrogenase (LDH) and unconjugated hyperbilirubinaemia.
- Coombs' test (direct antiglobulin) is usually positive with autoimmune haemolysis.

General Measures

- Treat the underlying cause.
- Do not transfuse prior to appropriate investigations, unless anaemia is severe.
- Coombs-positive haemolytic anaemia may be technically difficult to cross match.

 In G6PD deficiency, avoid drugs known to cause haemolysis, including Aspirin, sulphonamides (including Cotrimoxazole), dapsone and primaquine. In patients with cold agglutinins all transfusions must be given through a blood warmer to avoid cold-induced haemolysis.

Note: Efficacy of transfusion is limited by the shortened red cell survival due to haemolysis.

Treatment

 Supplement with Folic acid, oral, 5 mg daily given to all patients (because of high red cell turnover)

Note: For Autoimmune haemolytic anaemia, give **Prednisolone**, oral, 1–2 mg/kg daily, initial dose. When a satisfactory response is obtained with recovery of the haemoglobin and a decrease in LDH serum concentrations, taper dose over a period of 4 weeks to 30 mg daily. Thereafter further reduction should be slower to prevent disease recurrence.

- Prednisone treatment can be stopped when the Coombs' reaction becomes negative.
- If inadequate response add **Azathioprine**, oral, 2.5 mg/kg daily. Titrate to Hb response. May be required for several months. Monitor for neutropenia.
- Patients who fail medicine treatment should be considered for splenectomy.

Referral

When there is no response to medicine treatment.

1.7.4 Anaemia, Iron Deficiency

- Anaemia due to iron deficiency.
- Common causes of iron deficiency are chronic blood loss or poor nutritional intake. This is usually hypochromic microcytic anaemia

Investigations

 Assess for a haematological response to Iron therapy.

General Measures

• Identify and treat the cause. Dietary adjustment.

Treatment

- Oral Iron supplementation: reticulocytosis begins on the 3rd or 4th day after therapy, peaks at approximately day ten and lasts between 12 and 21 days.
- Give 100–200 mg of elemental oral iron daily with a meal, e.g.: Ferrous sulphate compound, oral, BPC 170 mg daily with food. (The expected haemoglobin rise is approximately 2 g/dL every 3 weeks.)

- Continue treatment for 6 months after the haemoglobin has returned to normal in order to replenish the iron stores adequately.
- Prophylaxis: **Ferrous Sulphate** compound 65 mg elemental iron oral, BPC 170 mg daily with meals (e.g. during pregnancy).

Note: If there is failure to respond to iron therapy, then consider the following causes: non-adherence, continued blood loss, wrong diagnosis, malabsorption, and mixed deficiency, concurrent folate or vitamin B12 deficiency.

Note: The use of parenteral iron may be associated with anaphylaxis. Parenteral iron is only indicated when oral iron is ineffective, e.g. malabsorption or patients on haemodialysis and erythropoietin therapy, or not tolerated. In people who require repeated therapy, the intravenous route is preferred.

- Where a once-off dose is required, give intramuscularly. Minimum required dose is 250 mg of iron per gram of Hb below normal. Use in consultation with a haematologist/specialist.
- Iron sucrose, IV.
 - Total dose = weight (kg) x [11 g/dL actual Hb (g/dL)] x 2.4 + 200 mg.
 - Maximum daily dose: 200 mg.
 - Administer over 30 minutes in 200 mL Sodium Chloride 0.9%.

- Repeat every second day until the total dose is given.
- Ensure that the correct formulation is given as some preparations can be given IM, or IV only, or both.
- Resuscitation equipment should be ready to manage anaphylaxis.
- Blood transfusion (see above section 1.2)

1.7.4 Anaemia, Megaloblastic

Anaemia caused by a deficiency of folate and/or vitamin B12.

Investigations

- Elevated MCV (mean corpuscular volume) and MCH (mean corpuscular haemoglobin).
- Macro-ovalocytes on blood smear; polysegmentation of neutrophils, thrombocytopenia with giant platelets.
- Decreased serum vitamin B12 or red blood cell folate. Pancytopenia in severe cases.
- Intrinsic factor antibodies in vitamin B12 deficiency, and anti-parietal cell antibodies in pernicious anaemia.

General Measures

 Dietary modifications to ensure adequate intake of folate and Vitamin B12. Identify and treat the underlying cause, e.g. antibiotics for intestinal overgrowth with bacteria.

Treatment

- Start with Folic Acid and Vitamin B12. Take blood samples for RBC, folate and vitamin B12 levels before starting treatment.
- Monitor serum potassium and replace if necessary.
- Give vitamin B12 and folic acid together until the test results are available as giving folic acid alone in patients with a B12 deficiency may precipitate a permanent neurological deficit.
- Adjust management according to results.
- Folic acid deficiency: Folic acid, oral, 5 mg daily until haemoglobin returns to normal.
- Prolonged treatment may be required for malabsorption states.
- Vitamin B12 deficiency: Vitamin B12, IM. 1
 mg daily for 7 days, then weekly for a
 further 4 doses. Follow with 1 mg every
 third month for life in patients with
 pernicious anaemia, except in patients with
 clearly modifiable nutritional deficiency.

Note: Response to treatment is associated with an increase in strength and improved sense of well-being. Reticulocytosis begins 3–5 days after therapy and peaks at about day 7. The anaemia is corrected within 1–2 months. The white cell count and platelets normalise in 7–10 days. As there is an increase in red blood cell production, short-term iron and folic acid supplementation is also recommended.

- Consider the following if there is failure to respond: co-existing folate and/or iron deficiency, infection, hypothyroidism, myelodysplasia, incorrect diagnosis, and drug-induced, e.g. hydroxyurea, Stavudine and Zidovudine.
- Prophylaxis. Vitamin B12 is indicated for patients after total gastrectomy or ileal resection. Give vitamin B12, IM, 1 mg every third month for life (4 times a year).
- Indications for folic acid: chronic inherited haemolytic anaemias, e.g. sickle cell anaemia, thalassaemia; myeloproliferative disorders; exfoliative skin disorders; increased demands, e.g. pregnancy, chronic haemodialysis. Give Folic acid, oral, 5 mg daily.

1.7.6 Febrile Neutropenia

Description

- Febrile neutropenia is defined as an absolute neutrophil count of < 0.5 x 10g/L with a temperature of greater than 38°C for > 1 hour.
- This is a medical emergency as these patients can rapidly develop features of severe sepsis (multi-organ failure and/or hypotension).

General Measures

- Treat the underlying cause of neutropenia, if applicable. Withdraw any drug that may cause neutropenia.
- Take blood cultures before starting antimicrobial therapy. Once culture results are available, adjust treatment to the most appropriate narrow spectrum agent.

Treatment

 For patients with febrile neutropenia within 48 hours of admission: 3rd generation cephalosporin, e.g.: Ceftriaxone, IV, 1 g daily. PLUS Gentamicin, IV, 6 mg/kg daily.

- If IV line infection is suspected as the cause at any stage: ADD Vancomycin, IV, 20 mg/kg/dose 12 hourly. Monitor trough levels after the third dose. Adjust dose to maintain a trough level of 15–20 micromol/L.
- If fever develops after 48 hours of admission choice will depend on local susceptibility patterns. One or more of the following antibiotics/classes must be available: Piperacillin/tazobactam, IV, 4.5 g 8 hourly or cefepime, IV, 1 g 12 hourly. OR Carbapenem with activity against Pseudomonas, e.g.: Meropenem, IV, 1 g 8 hourly or Imipenem, IV, 500 mg 6 hourly. Note: Ertapenem is not recommended because it is not effective for pseudomonas species which are important pathogens in this setting.
- If no response after 5–7 days ADD
 Amphotericin B, IV, 1 mg/kg daily in dextrose 5 % over 4 hours. Ensure adequate hydration to minimise nephrotoxicity.
 Regular, e.g. 3 times a week, monitoring of potassium, magnesium and renal function is essential.
- Duration of therapy: If neutrophil count increases to > 0.5 x 10g/L, continue for 2 days after fever has settled. If neutrophil count remains < 0.5 x 10g/L, continue for 7 days after fever has settled.

Referral/Consultation

• All cases – consult with haematologist/oncologist.

1.7.7 Myelodysplastic Syndromes

Description

 A group of disorders characterised by refractory cytopenias due to bone marrow failure. Anaemia is very common and there is a risk of developing acute leukaemia.

Investigations

- Evidence of cytopenia, with normal B12 and folate levels and substantial morphological dysplasia on the blood smear.
- Bone marrow examination confirms dysplasia of the blood elements and the presence of cytogenetic abnormalities.

Treatment

 Transfusion should ideally be with leucodepleted red cells to delay immunisation, as these patients require frequent transfusions. Bone marrow transplantation can be curative in selected patients if neutropenic and febrile.

Referral

 All patients for further investigation and management.

Bleeding Disorders

General Principles

A bleeding tendency may result from: a coagulation defect (congenital/acquired), a vessel wall defect, or a platelet defect (quantitative/qualitative).

A careful and detailed history, thorough examination and review of relevant laboratory investigations will allow differentiation between these three categories, as the management of each of these groups differs significantly.

Early consultation with a haematologist or a clinician with expertise in the handling of such patients is advisable.

Patients with a chronic bleeding tendency should be advised to wear a medic alert bracelet which clearly mentions the type of disorder he/she suffers from, e.g. Severe Haemophilia A, Factor VIII <1%, no inhibitors.

1.7.8 Haemophilia A And B, Von Williebrands's Disease

Description

Haemophilia A, haemophilia B and von Willebrand's disease are chronic bleeding disorders caused, respectively, by a lack of clotting factor VIII, clotting factor IX and von Willebrand factor (VWF, a carrier protein for factor VIII). Presentation depends on severity of the condition (see classification below). Complications include haemarthrosis with later chronic arthropathy, intracranial haemorrhage, soft tissue and muscle haematomas. Pain/tingling in a joint suggests bleeding into the joint in a known haemophiliac.

Subclassification (factor VIII and IX deficiency):

| CLASS | %OF | SIGNS |
|----------|--------|---------------------------|
| | NORMAL | |
| Mild | 5–25% | Occasional bleeds |
| Moderate | 2–5% | |
| Severe | < 1–2% | Trauma/spontaneous bleeds |

Investigations

- Prolonged partial thromboplastin time (PTT).
- Factor VIII or factor IX concentration < 25% of normal activity. Prolonged bleeding time (Von Willebrand's).
- Patient with factor VIII deficiency should be tested annually for factor VIII inhibitor.

General Measures

- Haemophilia register.
- Ideally, patients should attend a specialised haemophilia centre with a dedicated multidisciplinary health care team.
- Medic alert bracelet.
- Dental care (consult a haematologist to plan the procedure). Avoid contact sport.
- Acute bleeds into joints: apply ice packs; bed rest; rest the affected joint/limb until pain free and no further bleeding; no weight bearing; splint (no circumferential casting).

Treatment

 For mild to moderate pain: Paracetamol, oral, 1 g 4–6 hourly when required to a maximum of 4 doses per 24 hours. If needed ADD Tramadol, oral, 50mg, 6 hourly.

- For severe pain give Morphine, IV, 10 mg 4 hourly.
- Exercise great caution when taking blood specimens. Taking blood from femoral veins is absolutely contra-indicated. Avoid IM injections. Avoid aspirin and NSAIDS.
- Refer to haematologist/specialist for management including when contemplating dental extraction.

Referral

- All cases with suspected haemophilia (prolonged PTT and normal INR) to a haemophilia treatment centre, for assessment, genetic counselling and planning of management.
- Patients with proven antibodies against factor VIII.
- For further replacement, complex situations and complications consult a Haematologist.

1.7.9 Immune Thrombocytopenic Purpura (ITP)

Description

 A common bleeding disorder due to immune destruction of platelets. To diagnose ITP, isolated thrombocytopenia is present (rest of the complete blood count, including an examination of the peripheral blood smear, is entirely normal). Clinically apparent associated conditions, drugs (e.g. penicillins, cephalosporins, quinine, rifampicin and heparin), or other agents that may cause thrombocytopenia are NOT present. Patients with suspected ITP should be tested for SLE and for HIV infection.

Investigations

- Thrombocytopenia with normal white cell count and red cell series. Anaemia may be present due to blood loss.
- Peripheral blood smear to exclude RBC fragments. Smear may show large platelets.
- Do INR and PTT, which should be normal in ITP.
- If there is a poor response to treatment do a bone marrow biopsy.

General Measures

- Avoid: medication that affects platelet function, e.g. NSAIDs and aspirin; platelet transfusions unless life-threatening bleeds; dental procedures in acute phase, and IM injections.
- Reassure the patient that resolution usually occurs in acute ITP. Medic alert bracelet.
- Platelet transfusions may be given if surgery is required or in life-threatening bleeding.

Treatment

- Acute ITP: Prednisolone, oral, 2 mg/kg daily. Taper dose once response is achieved, usually within 10–14 days. Therapy may be required for a few months before prednisone is eventually discontinued. Also indicated for HIV-associated immune thrombocytopenia. Also start combination antiretroviral therapy urgently in these patients.
- Platelet transfusions: platelet transfusions are only indicated in acute active bleeding uncontrolled by other means or before procedures. In an adult, 1 mega-unit of single donor, leucocyte depleted platelets is usually sufficient to control the bleeding initially.
 Platelet transfusions have limited benefit in this condition as platelets are rapidly destroyed by the immune system.

Referral

 All cases not responding to steroids and, in the case of HIV patients, not responding to ART – discuss with haematologist.

1.8.0 Thrombotic Thrombocytopenic Purpura-Haemolytic Uraemic Syndrome (TTP-HUS)

Description

- Acute syndromes with abnormalities in multiple organ systems with evidence of micro-angiopathic haemolytic anaemia and thrombocytopenia. This condition presents with: anaemia; thrombocytopenia, often with purpura but not usually severe bleeding; acute renal insufficiency that may be associated with anuria and may require acute dialysis; neurologic abnormalities, and fever.
- TTP-HUS is associated with HIV infection and patients should be tested for HIV. TTP-HUS should be distinguished from disseminated intravascular coagulation (DIC) and severe pre-eclampsia where the coagulation profile is deranged.

Treatment

- In HIV-associated thrombotic thrombocytopenia, start combination antiretroviral therapy urgently.
- Fresh frozen plasma, IV infusion, 30 mL/kg in 3–4 divided doses. The use of platelet transfusions should be discussed with a specialist.

Referral

• All patients – discuss with a haematologist.

1.8.1 Acquired Coagulation Defects

1.8.2 Disseminated Intravascular Coagulation (DIC)

Management

- Identify and treat the underlying cause.
- If the patient is bleeding, replace haemostatic factors with cryoprecipitate or fresh frozen plasma.
- If the patient is not actively bleeding and platelet count > 20 000, then platelet transfusion is not necessary.
- Replacement therapy for thrombocytopenia should consist of 1 apheresis single donor unit / megaunit (expected platelet count increment 30–50 x 10⁹/L) or 6 random donor units (expected increment 50–60 x 10⁹/L), ideally aiming to raise the platelet count > 50 x 10⁹/L.
- In chronic DIC, or in the absence of bleeding, platelet transfusions should not be given merely to correct the thrombocytopenia.
- For hypofibrinogenaemia: Cryoprecipitate,
 8–10 units.

- For depletion of other coagulation factors:
 Fresh frozen plasma, 2–4 units, i.e. 15–20 mL/kg as initial dose. Volume: ±280 mL/unit.
- Repeat replacement therapy 8 hourly or less frequently, with adjustment according to the clinical picture and laboratory parameters.
- Perform frequent estimation of the platelet count and coagulation screening tests.

1.8.3 Disseminated Intravascular Coagulation (DIC)

1.8.4 Venous Thrombo- Embolism

Description

- Venous thrombosis should be seen as a spectrum from calf deep venous thrombosis to pulmonary thrombo-embolism. All patients should be seen as high risk.
- Differential diagnosis include: cellulitis, superficial thrombophlebitis, chronic venous insufficiency, lymphoedema, popliteal (Baker's) cyst, internal derangement of the knee, and calf muscle pull or tear
- Diagnosis is primarily clinical and confirmed with imaging studies, e.g. Doppler.

General Measures

- Acute management: in pulmonary embolism, cardiovascular resuscitation may be necessary and surgery may be undertaken for intractable disease.
- Note: Superficial thrombosis does not require anticoagulation. Distal venous thrombosis in the lower limbs, i.e. involving tibial veins only, need not be treated with anticoagulants.
- Monitor patients with repeat ultrasound if anticoagulants are not used.
 Ultrasonography should be repeated after a week but may be omitted if D-dimer is negative.
- Prophylaxis: Advice on prophylaxis should be emphasised. Eliminate all predisposing factors. Prevent deep vein thrombosis.

Treatment

 Acute treatment: unfractionated heparin initially, plus simultaneous warfarin. After 4–6 days, heparin is usually stopped and oral warfarin continued when a therapeutic INR level is reached. Note: Heparin and warfarin therapy should overlap for at least 5 days.

1. Blood and haematological conditions

 For proximal venous thrombosis and/or pulmonary embolism: Unfractionated heparin, SC, 333 units/kg as an initial dose. Follow 12 hours later by 250 units/kg/dose 12 hourly.

| Units of unfractionated heparin | | | Volume of heparin in mL (25 000 units/mL) | | |
|---------------------------------|----------------------|---------------------------------|---|-------------------|------------------------------|
| Weight (kg) | Loading dose (units) | 12 hourly dose (units) | | Loading dose (mL) | 12 hourly dose (mL) |
| 35 kg | 11 000 units | 8 750 units | | 0.44 mL | 0.35 mL |
| 40 kg | 13 000 units | 10 000 units | | 0.52 mL | 0.4 mL |
| 45 kg | 15 000 units | 11 250 units | | 0.6 mL | 0.45 mL |
| 50 kg | 17 000 units | 12 500 units | | 0.67 mL | 0.5 mL |
| 55 kg | 18 000 units | 13 750 units | | 0.73 mL | 0.55 mL |

1. Blood and haematological conditions

| 60 kg | 20 000 | 15 000 | 0.8 mL | 0.6 mL |
|-------|-----------------|--------------------|---------|---------|
| | units | units | | |
| 65 kg | 22 000 units | 16 250 units | 0.87 mL | 0.65 mL |
| 70 kg | 23 000 units | 17 500 units | 0.93 mL | 0.7 mL |
| 75 kg | 25 000 units | 18 750 units | 1 mL | 0.75 mL |
| 80 kg | 27 000 units | 20 000 units | 1.07 mL | 0.8 mL |
| 85 kg | 28 000 units | 21 250 units | 1.13 mL | 0.85 mL |
| 90 kg | 30 000 units | 22 500 units | 1.2 mL | 0.9 mL |

NB: Evidence indicates that PTT monitoring is not necessary with weight based dosing. However in morbid obesity and renal failure (eGFR < 30 mL/minute) unfractionated heparin should be used with PTT monitoring to maintain the PTT at 1.5 to 2.5 times the control. PTT should be taken 4 hours after SC dose.

OR

- Low molecular weight heparin, e.g. enoxaparin, SC, 1 mg/kg 12 hourly.
- Do not use LMWH in morbid obesity and renal failure (eGFR <30 mL/minute).
- Follow with: Warfarin, oral, 5 mg daily.
 Adjust dose to keep INR within therapeutic range. Continue warfarin for 3 months if there was a transient precipitating cause.
 Continue life-long if there is a non-transient precipitating cause or if repeated episodes.
 Contraindications for warfarin: first trimester and the last month of pregnancy. In these instances, replace with heparin.
- Most patients can be managed successfully with therapeutic anticoagulation.
 Thrombolytic therapy is indicated only in patients with angiographically confirmed early pulmonary embolism where haemodynamic stability cannot be achieved. Discuss with a specialist.
- Prophylaxis: prophylaxis is indicated for most medical and surgical patients. Low molecular weight heparin, e.g.: Dalteparin, SC, 5 000 units daily. OR unfractionated heparin, SC, 5 000 units 12 hourly.

1. Blood and haematological conditions

 Although the risk of bleeding is small, in the following patients prophylaxis should only be used under exceptional circumstances: active bleeding; intraocular, intracranial or spinal surgery; lumbar puncture or epidural anaesthesia within 12 hours; renal insufficiency; coagulopathy; or uncontrolled hypertension.

1.8.5 Heparin induced thrombocytopenia

- A severe immune-mediated drug reaction occurring in 1–5% of patients receiving heparin (unfractionated or low molecular weight heparin) therapy. It presents with thrombocytopenia and thrombosis. Diagnosis needs a high index of suspicion and should be considered if a patient has a 50% drop in platelet count within 5–10 days after initiating heparin therapy. Confirmation is done by positive antibody testing.
- Stop heparin and refer all patients.

Referral

• Refer all patients with Heparin-induced thrombocytopenia.

2.0 Cardiovascular diseases

2.1 Acute Heart Failure (Pulmonary Edema)

Signs and symptoms:

 Dyspnoea, cough {often with frothy, pink-tinged sputum}, tachypnoea, signs of increased respiratory effort and diffuse rales or crackles.

General measures:

- Prop up patient to sitting position
- Restrict fluids

Treatment:

- Oxygen therapy
- Drain pleural effusions if present.
- If BP >120/80, give sublingual
 Nitroglycerin for pulmonary vascular dilation
- Alternatively Digoxin if low systolic BP

Adults:

- Frusemide 40-80 mg slow IV (over 5 mins). Repeat if required
- Intravenous Morphine 2.5 mg -5-10mg (Be cautious of patients with low blood pressure) and Metoclopramide 10mg IV.
- Repeat both if required.

Alternatively:

Second Line action (refer patient to next level of care)

- Depends on systolic blood pressure
 - Nitrates if SBP >100mmHg
 - Dopamine /Epinephrine if SBP 70-100 mmHg and with signs and symptoms of shock

Children:

- Give Morphine 0.1-0.2mg/kg slow IV (over 5 mins) Repeat every 4 hours if required
- Give **Frusemide** 1-2mg/kg IV, po.
- Specific treatment should be given according to the cause e.g. hypertension

2.2 Congestive Heart Failure

- Defined as a clinical syndrome in which patients have typical symptoms and signs resulting from abnormalities of Ventricular function
- Heart failure is a syndrome not a final diagnosis therefore it is very important to establish the cause

Signs and symptoms:

 Shortness of breath, fatigue,
 Orthopnoea, Paroxysmal nocturnal dyspnea, ankle and body swelling

General measures

- Patient and family education on heart failure
- Restriction of fluids
- Reduce salt intake
- Moderate exercises
- Stop smoking and alcohol intake
- Clinicians to screen and treat other cormobidities

Symptomatic relief:

- Give Frusemide 40-160 mg in divided doses
- Give **Digoxin** (0.125 mg daily) is recommended in patients with heart failure and rapid atrial fibrillation

Symptomatic relief and mortality benefit:

- ACE inhibitors: Enalapril, start at low doses 2.5-5mg and escalate to 10 mg twice a day.
- Spironolactone: 25 mg daily dose for patients with persistent symptoms of heart failure (NYHA II-IV)

Note: ARB (Angiotensin Receptor Blocker) or Hydralazine + Nitrate can be an alternative if a patient cannot tolerate ACE inhibitors or has acute renal failure.

2.3 Congenital Heart Disease

Important points in history and exam:

 Cyanosis- Clinican should determine: when was it noted, was it present all the time, or what brings a cyanotic spell, does child squat?

Other cardiac symptoms:

- Breathlessness on feeding, sweating, failing to thrive, easily fatigued
- Dysmorphic, clubbed, cyanosed, pulse rate and volume, Bp, apex beat location, hepatomegaly and oedema, bulging chest

Indications for admission:

- Cyanosis in any child with acute cyanosis (except if longstanding and has already been investigated)
- Any evidence of acute cardiac failures. If child has a heart murmur, then may be referred to a hospital clinic

Cyanotic congenital heart disease presenting in newborn as cyanosis:

- Consider if blue baby doesn't pink up with oxygen, may or may not have signs of heart failure nor murmur. May be difficult to distinguish from persistent pulmonary hypertension
- Investigate with chest x ray and or cardiac echo, parental counselling re condition. Baby might have persistent pulmonary hypertension give oxygen in this case.

Cyanotic congential heart disease presenting as cyanosis in older child

 Usually Tetralogy of Fallot, associated with hypercyanotic spells

Management of spell

- Place child in knee chest position or encourage squatting in older child.
- Give oxygen
- Give IM or IV Morphine 0.1mg/kg or oral Propanolol 0.5mg/kg statistically

Long term management

Propanolol 1mg/kg BD to prevent spells

Congenital heart disease causing heart failure

Examples include Large
 <ventricular septal defect,
 AtrioVentricular septal defect,
 large persistent ductus arteriosus

Management

- Manage cardiac failure as per protocol
- Refer to hospital clinic
- If child has not developed pulmonary hypertension may benefit from surgery
- Generally, antibiotic prophylaxis against bacterial endocarditis when undergoing dental treat or any surgery (see infective cardiac disease)

2.4 Hypertension

- Diagnosis is based on a raised blood pressure measured while patient is at rest on at least 3 separate readings.
- Hypertension is generally asymptomatic.
- Essential hypertension is unusual in children and young adults and an underlying cause should be excluded at hospital level

Causes include:

- Renal disease (glomerulonephritis, Hamolytic uraemic syndrome, renal failure, cystic renal disease)
- Renovascular (renal artery stenosis)
- Cardiovascular (coarctation of the aorta)
- Endocrine (cushings, neuroblastoma)

Children:

- Refer all children with hypertension to a doctor for management
- In children, hypertension is defined statistically because BP levels vary with age and outcome. Based data are not available for this population. Hypertension is defined as systolic and /or diastolic pressure levels greater than the 95th percentile for age and gender on at least 3 occasions
- The upper limit for normal systolic Bp in children greater than One year may be calculated as follows:
 - \triangleright (Age in years x 3) +100
 - ➤ Diastolic BP is 2/3 of systolic BP

2. Cardiovascular diseases

 The table below shows normative blood pressure levels (systolic/diastolic) in children up to age 5 years. Blood pressures above the 95th percentile indicate hypertension

| Age | Mean BP levels | 95 th Percentile |
|----------|----------------|-----------------------------|
| 1-3 days | 64/41 (50) | 78/52 (62) |
| 1mo -2yr | 95/58 (72) | 110/71 (86) |
| 2-5yr | 101/57 (74) | 115/68 (85) |

 Remember to use the correct cuff size when measuring BP. It should cover 2/3 of the upper arm

Classification of Hypertension

| Type of Hypertension | Systolic Blood | Diastolic Blood Pressure | |
|----------------------|-------------------|--------------------------|--|
| | Pressure | | |
| Mild | 140-159 | 90-99 | |
| Moderate | 160-179 | 100-109 | |
| Severe | >180 | >110 | |

General measures

- Reduce salt intake
- Stop smoking
- Regular exercise
- Lose weight
- Avoid excessive alcohol consumption

Note: Consider medicine treatment for mild hypertension only if the above general measures are unsuccessful

Treatment:

- Explain to the patient that treatment must be regular (every day), closely monitored and generally has to be taken for life
- Use the following stepped treatment approach with the medicines in this order unless there are specific contraindications, comorbidities or side- effects

Stepped anti-hypertensive treatment approach (adults):

- Step 1: Hydrochlorothiazide 25 mg each morning, increasing the dose is not advised Explain to the patient that treatment must be regular (every day), closely monitored and generally has to be taken for life
 - Alternatively, give Bendrofluazide 2.5 mg each morning

Note: Avoid in pregnancy and breastfeeding

Step 2: Give Hydrochlorothiazide
 25mg once daily and Amlodipine 5 10mg once daily

- Where Amlodipine is not available
 Nifedipine 10-20mg slow release
 tablets twice daily can be used.
- Step 3: Give Hydrochlorothiazide
 25mg once daily, Amlodipine 5-10
 mg once daily and Enalapril 10 20mg once daily
 - Where Enalapril is not available Captopril 12.5-50mg every 8 hours can be used.

Note:

- (i) Best to start with a lower dose of Enalapril 5mg and increase to 10mg after observation of the BP response over a few days.
- (ii) Avoid Enalapril and Captopril in pregnancy and breast-feeding
 - Step 4: Give Hydrochlorothiazide
 25mg once daily and Amlodipine 510mg once daily, Enalapril 10-20mg
 once daily and Atenolol 50-100mg
 once daily
 - Where Atenolol is not available
 Propranolol 40mg 80mg every 8
 hours can be used.
 - Step 5: Refer to Medical Specialist

Note:

- (i) Side-effects may outweigh benefits
- (ii) In patients with severe
 hypertension or complications
 (heart failure, renal failure) start
 medicine treatment immediately
- (iii) In patients without co-morbidity, aim for a BP of around 140/90
- (iv) For patients taking ART, because of the interactions between Calcium Channel Blockers, and NNRTIs, please consider Enalapril or Atenolol before a Calcium Channel Blocker.

2.4.1 Emergency hypertension treatment

Signs and symptoms:

- Encephalopathy, convulsions, retinal hemorrhages or blindness.
- Reduce the blood pressure in a controlled manner to avoid impaired auto- regulation of cerebral blood flow.
- Only use parenteral therapy in:
 - hypertensive heart failure
 - hypertensive encephalopathy
 - > malignant hypertension
 - > eclampsia
 - hypertension and dissecting aneurysm of the aorta

2. Cardiovascular diseases

Note: Intravenous rapid lowering of blood pressure has several risks and should be done under close monitoring only, preferably in a high or intensive care setting. It is only indicated in hypertensive emergencies mentioned above.

Treatment:

Adults:

- Give **Hydralazine** 5-10 mg IM
- Repeat up to every 1 hour as necessary
- If heart failure: add Frusemide 40 mg
 IV stat

Sub-lingual **nifedipine** (10 mg) should be avoided due to the unpredictable response of the blood pressure, unless parenteral drugs are unavailable.

Children:

- For fluid overload: Give Frusemide 1 mg/kg bolus IV or IM
- hypertensive encephalopathy: Give
 Hydralazine 0.15 mg/kg slow IV
 - Repeat every 30-90 minutes as required
 - Maximum dose: 1.7-3.6 mg/kg in 24 hours
- Long term management of hypertension would depend on the cause hence these patients need to be referred for proper management.

2.5 Ischiemic Heart Disease

 Condition in which there is inadequate blood and oxygen supply to any portion of the myocardium

General measures

- Minimize risk factors by:
 - Weight reduction (if obese)
 - Control of hypertension
 - Control of diabetes
 - Stop smoking
 - Address other factors such as:
 - High blood cholesterol
 - Stressful lifestyle
 - Excessive alcohol intake
- Encourage regular moderate exercise

2.5.1 Stable Angina (infrequent attacks)

 Central chest pain (squeezing, heavy discomfort) with radiation to the left arm on exertion or at rest lasting 2-5 minutes, crescendo and decrescendo pattern

Treatment

• Give Aspirin 150 mg daily

Acute relief of Angina

- Give Glyceryl Trinitrate 0.5 mg sublingually as required.
- Maximum 3 tablets per 15 minutes
- Deteriorates on storage: keep tablets in original container for no more than
- 3 months after opening
- Alernatively use Isosorbide Dinitrate 5-10mg sublingually as required instead of Glyceryl Trinitrate

Long term management

- Check correct doses
- Atenolol 50 mg bd
- Amlodipine 5-10mg daily/Nifedipine 10-20 mg daily replace or be cautiously added to Atenolol
- If pain continues despite the above treatment refer to Medical Specialist

Red flags

- UNSTABLE ANGINA, NON ST SEGMENT MYOCARDIAL INFARCTION
 - Central chest pain as above lasting more than 10 minutes and has a crescendo pattern
- ACUTE MYOCARDIAL INFARCTION (ST SEGMENT ELEVATION)
 - Typical angina pain plus most patients being restless, anxious, pale and with cold extremities

Treatment

- Nitrates and Morphine 2-5mg IV for pain control if there is no hypotension
- URGENTLY DISCUSS THESE PATIENTS WITH MEDICAL SPECIALIST!

2.6. Peripheral Arterial Disease

- If there are poor pulses and delayed capillary refill, or claudication or non healing wounds, consider adding Aspirin and Statin
- Patients with critical limb Ischemia should be referred to high level of care to avoid death from septic shock

3.0 Central Nervous System Conditions

3.1 Seizures and Epilepsy

3.1.1 Seizures

- Sudden abnormal function of the body, often with loss of consciousness, and excess of muscular activity, or sometimes loss of it, or an abnormal sensation.
- Ensure airway is clear and patient is not hurting himself. Turn patient in a recovery position. Don't insert any object between the teeth
- Monitor blood sugar. If hypoglycemia is suspected, give 1 ml/kg 50% Dextrose or 5 ml/kg 10% Dextrose.

Treatment:

Adults:

- Give Diazepam 5-10 mg IV slowly.
 Repeat once after 10 minutes.
- If convulsions continue for another 10 minutes or are repeated more than 3 times without patient gaining consciousness between seizures, treat as status epilepticus. If repeated seizures, consider antiepileptic therapy.

3. Central nervous system conditions

- Look for treatable causes and provoking factors (malaria, infection, tumour, alcohol).
- Diazepam IM absorbs slowly and unreliably: IV or rectal routes are preferable

3.1.1.1 Generalized seizures in children

Treatment

 Give Sodium Valproate 20-40 mg/kg/day in 2 to 3 divided doses

Alternatively

or

- Give **Phenobarbitone** 5-8mg/kg daily
- Give Carbamazepine 2.5mg/kg per dose twice daily,
- Increase the dose weekly by 5mg/kg until 20mg/kg is reached.

3.1.1.2 Partial seizures in children

Treatment

Give Carbamazepine 5mg/kg/day
 in 2 divided doses (2.5mg/kg bd)

Alternatively

 Give Sodium Valproate 20-40 mg/kg/day in 2 to 3 divided doses

3.1.1.3 Petit mal

Treatment

 Give Ethosuximide 15 mg/kg at night as a single dose increased gradually if necessary to 50 mg/kg daily in 2 divided doses

3.1.2 Status Epilepticus

- Continuous seizure activity or seizures without recovery of consciousness for > 30 minutes
 - Always an emergency, mortality is high.
 - Clear airway, insert ivline, position patient in a recovery position.
 Don't insert any object between the teeth.

Treatment Adults:

- Give Diazepam 5-10 mg IV. Repeat every 10 minutes until the patient stops convulsing. If patient not controlled give continuous diazepam IV infusion with careful attention of respiratory depression.
- Give a loading dose of anti-epileptic medicines:
 - Give **Phenytoin** 15 mg/kg (600-1200 mg) IV. Dilute with 100 ml normal saline and give slowly, no more than 100 mg/minute.
- If still fitting after 10 minutes, then give Phenobarbitone 10 mg/kg
 (400-600 mg) IV: dilute with water for injection 1:10 and give slowly, no more than 100 mg/minute. Or give 200 mg IM in each buttock
- If status continues, give
 Paraldehyde 5 ml deep IM in a buttock, and repeat 5 ml IM in alternate buttock. Paraldehyde can also be given through the rectum using a syringe with needle removed.
- Ensure that the dose is given promptly and therefore remains in the syringe for only a short time (Paraldehyde dissolves plastic)
- Check blood sugar. Give glucose, if suspicious of hypoglycemia

3. Central nervous system conditions

- Give Thiamine 100mg IV or IM once daily before giving glucose if patient suffers from alcoholism. Continue for 3 days.
- If still no improvement, consider general anaesthesia (in ICU setting preferably).
- If patient improves, start antiepileptic treatment and continue until cause of status epilepticus is treated.

3.1.3 Epilepsy

- Repeated seizures due to a disorder of the brain cells.
- Look for treatable causes (infections, neuro-cysticercosis, tumour)
- Counsel patient: no bathing alone, careful with fire, driving and climbing.
- Young female patients should be advised to plan their pregnancy.
 When they wish to get pregnant folic acid once daily should be started and continued through the pregnancy.
- Doses may be reduced to the lowest level that still prevents convulsions.

- Treatment should not be stopped because of pregnancy: it is more dangerous for the mother and foetus to have uncontrollable seizures than to continue the anti-epileptic medicine.
- If patient has more than 2 seizures in a year of unknown cause, consider starting antiepileptic therapy.
- Always start with small dose.
- Increase dose gradually over weeks or months.
- Use maximum dose of one medicine before adding another.
- Treatment should never be stopped suddenly due to risk of status epilepticus, but rather taperedoff over weeks or months.

Treatment

• Give **Phenobarbitone** sodium 60-180 mg at night

Alternatively

 Give Carbamazepine 100 -200mg 1-2 times daily. Increase by 100 -200 mg weekly until dose is 800 mg -1200mg per day.

or

 Give Sodium Valproate 600 - 2000mg daily divided in 2 doses.

or

 Give Phenytoin 150 - 300mg daily divided in 1-2 doses. Can be increased to 500mg daily.

3.2 Stroke

• There are 2 types of stroke: Ischaemic/Embolic and Haemorrhagic. Without imaging (not commonly available) you cannot distinguish. Ischaemic is more common. (Sudden reduction of blood flow to the brain resulting in a neurolgical deficit corresponding to that area of the brain's function.)

Common Symptoms Clinical features

 Facial drooping, slurred speech, dragging a leg, arm/leg weakness on one side.

Acute Treatment

- All strokes are likely to benefit from statin therapy. Commonly used statin is Simvastatin and Atorvastatin as second line therefore it will be ideal to put Simvastatin 10-20 mg nocte
- Ischaemic stroke benefits from Aspirin:
 300mg chewed x1 then 75mg od
- Ischaemic: allow BP to remain moderately elevated for 1-2 weeks (140-180/90-100) then treat to normal.
- Haemorrhagic: reduce BP faster (140/190) and avoid Aspirin
- Assess safety/ ability to swallow without aspirating (NG tube if needed)

If signs of increased intracranial pressure (vomit, lower level of consciousness) should be given **Mannitol**.

Chronic Treatment

- Rehabilitation/ physiotherapy
- Treat underlying conditions with reference to their sections, such as: Syphilis, HIV, Diabetes, Hypertension, Hyperlipidemia (simvastatin), atrial fibrillation (aspirin or warfarin)

Note:

- Conditions that may present similar to Stroke include Subdural haematoma brain masses, meningitis, and enchepratitis.
- Recurrent stroke when already taking
 Aspirin should be referred to a specialist.
- Sudden non-convulsive loss of neurological function due to ischemic or hemorrhagic vascular event.
- Remember brain infection as a differential diagnosis in HIV infected patients
- Risk factors include hypertension, diabetes, smoking, genetic disorders, atherosclerosis, cardiac disease, atrial fibrillation, HIV and high cholesterol.
- Look for treatable cause and counsel the patient.
- 85% of the strokes are ischaemic.

Treatment

- Give long term Aspirin 75mg once daily
 Note: Aspirin is not advised in intra-cerebral or subarachnoid hemorrhage.
 - Give IV fluids to correct any dehydration, avoid IV glucose.
 - Treat hyperglycemia
 - Treat any fever: look and treat for infections (aspiration pneumonia, urinary tract infection common).
 - Give Paracetamol 1g every 8 hours orally to reduce fever.

- Keep patient half seated if increased intracranial pressure is suspected (e.g. in drowsy or unconscious patients).
- Don't start new antihypertensives during the first 10 days.
- If hypertension is repeatedly higher than 180/120 mmHg, start
- Give Hydrochlorthiazide 25mg once daily orally
- Start physiotherapy on day 1.
- Start mobilisation as soon as it is possible. Consider referral to an institutional rehabilitation unit.

3.2.1 Subarachnoidhaemorrhage

Description

Bleeding into the subarachnoid space, most commonly due to the rupture of a vascular aneurysm. Patients frequently present with an acute onset of severe headache and may have additional neurological symptoms and signs. Diagnosis is confirmed preferably by neurological imaging and, when this is not available, urgently do lumbar puncture, demonstrating xanthochromia.

General Measures

Maintain normal hydration and electrolyte status. Control blood pressure.

Treatment

Analgesia if level of consciousness is not impaired:

 Paracetamol, oral, 1 g 4–6 hourly when required to a maximum of 4 doses per 24 hours.

If no response:

 Morphine, IV, 1–2 mg/minute to a maximum total dose of 10 mg. Dilute 10 mg up to 10 mL in sodium chloride solution 0.9%. This may be repeated 4 hourly.

Avoid NSAIDs.

In patients with grades 1 to 3 impairment of consciousness level while waiting for transfer to neurosurgical facility and in consultation with neurosurgeon:

• Give **Nimodipine**, oral, 60 mg 4 hourly for 21 days.

Referral

- All patients with minimal impairment of consciousness level for possible angiography and appropriate neurosurgical management. Patients initially deemed unsuitable for further investigation, may be referred at a later stage, should their condition improve.
- For neurological imaging: patients in whom the diagnosis has to be confirmed radiologically and where a lumbar puncture may be considered hazardous.

3.3 Headaches and Facial Pain Syndromes

3.3.1 Migraine

Description

Episodic headache, usually focal in nature, which may occur with or without an aura. It is usually accompanied by nausea and vomiting. Several variants of migraine also occur.

General Measures

Reassure patient that this is a benign condition. Attempt to identify any precipitating factors or food allergies from the history (although this is usually unrewarding), and try to diminish patterns of tension.

Treatment

Acute treatment

Initiate therapy during the attack or at the onset of the headache.

Analgesics, e.g.:

 Paracetamol, oral, 1 g 4–6 hourly when required to a maximum of 4 doses per 24 hours.

OR

NSAIDs, e.g.:

 Ibuprofen, oral, 800 mg immediately then 8 hourly, if needed.

If severe and not responding to therapy above:

Morphine, IM, 10 mg as a single dose.

For nausea:

• Give **Metoclopramide**, oral/IM, 10 mg 8 hourly.

Prophylaxis

Regular, daily, prophylactic therapy is advised if:

- attacks are frequent, i.e. more than 2–3 per month, or
- severe, causing a significant amount of disability, or attacks are long lasting.
- Also consider for patients who tolerate therapy for acute attacks poorly.

Give:

Amitriptyline, oral, 10–25 mg at bedtime.
 Titrate dose up to adequate response. More than 75–150 mg as a single bedtime dose is seldom required.

OR

• Propranolol, oral, 20–80 mg q12h. Note: The evidence for using atenolol for this indication is limited.

OR

 Carbamazepine, oral. Start with 100 mg q12h. Increase every two weeks up to a maximum of 400 mg 12 hourly.

Note: Only about half of patients will respond to one of these agents and this response may take 1 to 2 months to occur.

Referral

- Patients with unexplained neurological signs, additional risk factors for an alternate diagnosis, such as immune deficiency, or an atypical short history require brain imaging.
- Sudden onset of a first severe headache, even if it resembles migraine, as this may indicate serious organic pathology, such as subarachnoid haemorrhage.
- Acute migraine, not responding to treatment.
- Recurrent migraine not controlled with prophylactic therapy.

3.3.2 Cluster Headache

Description

Repetitive episodes of excruciating headache typically of short duration (up to 2 hours) in clusters for weeks to months at a time. Typically the headache is of sudden onset, unilateral during the specific cluster, and quickly reaches a climax. Associated redness of the eye with lacrimation and rhinorrhoea occurs.

Treatment

Oxygen inhalation may abort some episodes.

Analgesics are ineffective in this indication.

To induce rapid remission in patients with episodic cluster headache:

Prednisone, oral, 40 mg daily for 5–10 days.
 Tapering is not necessary when the above duration is used.

OR

Verapamil, oral, 40 to 80 mg 8 hourly.

Referral

 When the is inadequate response to treatment.

3.3.3 Trigeminal Neuralgia

Description

Severe, very short lived stabs of facial pain in the sensory trigeminal distribution. It is important in the diagnostic workup to exclude intracranial mass lesions, which may impinge on the trigeminal nerve.

Treatment

 Carbamazepine, oral, 100 mg 2–3 times daily, initial dose. Increase dose slowly.
 Doses of up to 1, 200 mg daily may be required. After exacerbation, reduce to maintenance dose of 400–800 mg daily.

Referral

- For neuro-imaging, if not available locally.
- If there is poor response to single drug therapy.

3.3.4 Tension Headache

Description

Headache over the back of the head, but sometimes over the entire head, described as a tight band around the head, usually worse in the afternoon.

3. Central nervous system conditions

General Measures

Consider use of relaxation techniques. The importance of this diagnosis is the exclusion of other, more sinister conditions. Exclude analgesia overuse headache.

Treatment

• Amitriptyline, oral, 10–75 mg at night.

Referral

- When there is atypical pain, suggestive of alternate diagnosis.
- When there is poor response to therapy.

3.4 Infectious and Parasitic Conditions

3.4.1 Viral Meningoencephalitis

Description

Patients present with headache, fever and mild meningism. Lumbar puncture typically shows mildly elevated protein, normal glucose and mildly raised cells (< 500), mainly lymphocytes (early on polymorphs may predominate). Most cases do not require specific therapy, other than analgesia.

Treatment

Analgesia, i.e.:

 Paracetamol, oral, 1 g 4–6 hourly when required to a maximum of 4 doses per 24 hours.

AND

Tramadol, oral, 50 mg 6 hourly.

OR

- Morphine, IV, 1–2 mg/minute to a maximum total dose of 10 mg. Dilute 10 mg up to 10 mL in sodium chloride solution 0.9%. This may be repeated 4 hourly.
 - Beware of respiratory depression in patients with reduced level of consciousness.

Clinical features are fever, change in behaviour and seizures, which may be either focal or generalised.

Evidence of mucocutaneous involvement is usually not present. Lumbar puncture shows the above features of viral meningoencephalitis, but in this condition may be additionally haemorrhagic in nature. A temporal focus on EEG or neuro-imaging is strongly supportive of the diagnosis. A positive HSV PCR test on CSF is diagnostic.

Aciclovir, IV, 10 mg/kg 8 hourly for 21 days. Start therapy as early as possible, i.e. before results are available. If PCR is negative, stop treatment.

Treat seizures appropriately with **Phenytoin** or **Carbamazepine**.

It is important to initiate therapy and then refer to centre where neuro- imaging is available.

Referral

 For neuro-imaging: patients not responding or worsening in condition, i.e. decrease in consciousness and cranial nerve palsies, despite appropriate therapy.

3. Central nervous system conditions

- This is especially urgent in patients with tuberculous meningitis, who may develop hydrocephalus and require an urgent shunting procedure.
- Patients with shunts.

3.4.2 Brain Abscess

Diagnosis

Patient may present with focal neurological signs and signs of infection. Neurological signs may not always be prominent. Neuro-imaging usually confirms diagnosis. Patients may have concomitant infection of ears, paranasal sinuses or lower respiratory tract.

Treatment

Empiric antibiotic therapy

• Ceftriaxone, IV, 2 g 12 hourly.

PLUS

 Metronidazole, oral, 400 mg 8 hourly or IV, 500 mg 8 hourly. Adjust according to antimicrobial sensitivity after surgical drainage.

Referral

 All, as patients require urgent neurosurgery opinion and treatment.

3.4.3 Neurocysticercosis

Diagnosis

Patients may present with seizures and/or focal neurological deficit. Typical cystic lesions are seen on neuro-imaging.

General Measures

Health education.

Surgery for treatable ventricular blockage or spinal or intraocular cysts.

Drug Treatment

For active or viable cysts only:

Albendazole, oral, twice daily for 8 days. >
60 kg: 400 mg. < 60 kg: 7.5 mg/kg to a
maximum of 800 mg daily. Do not use in
pregnancy.

Progressive recovery may occur for a period of up to one year. The presence of viable cysts does not require repeating antihelminthic treatment.

Drug-induced damage to cysticerci may precipitate an acute inflammatory reaction, the intensity of which is related to the number of viable cysts and may cause cerebral oedema. This reaction is minimised by adding corticosteroids to the antihelminthic treatment, e.g.: **Prednisone**, oral, 60 mg daily for 8 days.

Anticonvulsants, if required.

3.5 Movement Disorders

Description

Abnormalities of movement/initiation of movement, divided into those with reduction of movement (hypokinesia or bradykinesia), or those with excessive movements (hyperkinesia).

Referral

- To differentiate functional from organic disorders.
- Tardive dyskinesia.
- All complicated cases, i.e. patients with Parkinsonism, not responding to small doses of carbidopa/levodopa.
- Patients with Parkinsonism developing disease-, drug- or autonomic nervous system complications.
- Patients with myoclonus or chorea, not responding to therapy.

3.5.1 Parkison's Disease

Description

Parkinsonism is a syndrome characterised by tremor, rigidity, bradykinesia and postural disturbances. It may be primary, i.e. Parkinson's disease, or secondary, i.e. drug-induced or due to uncommon disorders that may initially resemble Parkinson's disease.

The objective of treatment is to:

- minimise disabling symptoms,
- prevent complications and avoid serious drug-induced side effects, and
- exclude secondary forms.

General Measures

Educate the patient. General supportive therapy and advice about lifestyle modification, physiotherapy and occupational therapy.

Treatment

Note: Set therapeutic targets so that the patient is functioning as well as possible.

Primary Parkinsonism Bradykinesia, rigidity and postural disturbance:

 Carbidopa/levodopa, 25/100 mg, oral, ½ tablet 8 hourly. Increase dose in consultation with a specialist.

If optimal control has not been achieved, consider an alternative diagnosis or changing to a drug containing a higher dose of levodopa:

Carbidopa/levodopa 25/250 mg. Specialist initiated.

3. Central nervous system conditions

Drug-induced Parkinsonism

Anticholinergics have a very small role in this setting and should be used with caution.
Anticholinergic agent, e.g.: **Orphenadrine**, oral, 50 mg 8 hourly.

Tremor only:

Consider anticholinergic agent, e.g.:

- Orphenadrine, oral, 50mg 8 hourly.
 Increase gradually according to clinical response or maximum dose of 400mg daily
- Usual dose: 150–250 mg daily.

Acute dystonic reaction

Usually follows administration of dopamine antagonistic drug, e.g. **Metoclopramide** and **Phenothiazines**. Anticholinergic agent, e.g.:Biperiden, IM/IV, 2 mg. Repeat as necessary.

Referral

- If there is no improvement or poor control with treatment.
- Increasing on/off phenomenon.
- Dyskinesias.

3.5.2 Essential Tremor

General Measures

Exclude and manage alternate causes, such as drugs, thyrotoxicosis, hyperadrenergic states and psychiatric disorders.

Occasionally a patient may present with essential tremor and an additional neurological condition, which may make the diagnosis difficult.

Medicine Treatment

If tremor is severe and interfering with normal daily activity: Give ß-blocker, e.g.:

 Propranolol, oral, 60–320mg daily in divided doses.

3.5.3 Myoclonus

Description

Irregular, involuntary movements due to muscle jerks, which may be due to myoclonic seizures, but may follow injuries to the brain and are thus not always of an ictal nature.

Referral

• All patients where the diagnosis is unclear.

3.5.4 Chorea

Description

Involuntary random, irregular movements. Aetiology is classified as:

- primary Huntington's chorea, benign hereditary chorea and others; or
- secondary due to Sydenham's chorea, vascular pathology, metabolic, endocrine and infective conditions, amongst others.

Treatment

• Give **Haloperidol**, oral, 0.5– 5 mg 2–3 times daily.

To be prescribed by a specialist only.

3.5.5 Neuropathy

Description

Defective functioning of nerves, which may involve both peripheral nerves (peripheral neuropathy) and cranial nerves. Different patterns are noted, i.e. polyneuropathy, mononeuritis multiplex and mononeuropathy, each of which may be caused by axonal degeneration or demyelination or a combination of the above.

3. Central nervous system conditions

Clinical features may be predominantly of a sensory, sensorimotor or motor nature. Important causes of neuropathy include:

- alcohol,
- diabetes,
- HIV infection,
- thiamine deficiency,
- acute inflammatory demyelinating polyradiculoneuropathy (Guillain-Barrè), and
- chronic inflammatory demyelinating polyradiculoneuropathy (CIDP).

General Measures

Observe rate of progression. If the disease is progressing fairly rapidly, i.e. deterioration noted over 5-7 days, admit patient and monitor ventilatory status carefully with spirometry, as intubation and ventilatory support may be required. Remove the cause where possible, i.e. drug- or alcohol-induced neuropathy, control diabetes mellitus, etc. Specialised nursing care and dedicated physiotherapy may be indicated.

If not managed appropriately, chronic cases may develop contractures, weakness affecting gait, develop chronic bedsores and become wheel chair- bound.

Treatment

Most cases respond to management of the underlying disease process or removal of the aetiological agent.

Neuropathic pain (i.e. pain due to a disease or injury of the central or peripheral nervous system)

• Give **Amitriptyline**, oral, 25–75 mg daily.

OR

 Give Carbamazepine, oral, 200–1200 mg daily in divided doses.

Isoniazid-induced polyneuropathy

• Give **Pyridoxine**, oral 75 mg daily for 3 weeks. Follow with 25–50mg daily.

Post-herpes zoster neuropathy (**Note:** Aciclovir is not beneficial in treating this condition).

• Give **Amitriptyline**, oral, 25–75 mg daily.

AND/OR

 Give Carbamazepine, oral 200–1200 mg daily dose in divided doses. Beware of possible drug interactions in patients on ART.

Bells' palsy Note:

Exclude herpes zoster
Start within 4 days of onset of symptoms:

 Give Prednisone, oral, 60 mg daily for 7– 10 days.

Referral

- Electrophysiological studies may be needed in the diagnostic assessment, although many common causes do not warrant specialist investigations, e.g. polyneuropathies due to diabetes mellitus, HIV, isoniazid, hydralazine, dapsone, antiretrovirals (stavudine and didanosine), amiodarone and alcohol. These cases may initially be managed locally, with referral of non-responding or atypical cases.
- Gullain-Barré Syndrome: referral criteria are progressive, extensive paralysis with impending respiratory failure, bulbar palsy and swallowing problems, and aspiration, as well as for diagnostic confirmation.

3.5.6 Acute Myelopathy

Description

Patients present with a sudden onset of paraparesis, with associated sensory loss, i.e. a sensory level may be found. Incontinence and autonomic instability may be present.

There are numerous causes for this condition and it is important to exclude neoplastic and infectious conditions, i.e. granulomas and abscesses, causing external compression of the spinal cord. Lesions, such as intervertebral disk prolapse, and mass lesions below the spinal cord may present with cauda equina syndrome. These cases usually have asymmetrical weakness, but may have saddle anaesthesia and sphincter involvement alone. Incontinence is a marker of severity.

Note: Do not perform a lumbar puncture, until obstructive lesions of the spinal cord have been excluded clinically or radiologically.

Referral

All patients for urgent imaging.

3.5.7 Multiple Sclerosis

Description

A demyelinating disease of the central nervous system, characterised by episodes of unifocal or multifocal neurological dysfunction. Diagnosis is confirmed by imaging. The CSF may show oligoclonal bands and raised IgG index.

Recovery between acute flares of illness is common, although a general stepwise degeneration in baseline is usually found.

Consult with neurologist/specialist for diagnosis and treatment.

Referral

All patients.

3.5.8 Oedema, Cerebral

Description

Swelling of brain parenchymal tissue, due to vasogenic, cytotoxic and osmotic causes. Only the vasogenic causes, such as brain tumours and inflammation, respond to corticosteroids.

Consider mannitol for brain oedema in traumatic brain injury causing raised intracranial pressure, pending neurosurgical intervention.

3.5.9 Brain Oedema Due to Tumours and Inflammation

General Measures
Supportive management.

Treatment

Treat the underlying cause. This is especially important with brain oedema associated with systemic conditions, such as electrolyte disturbances and organ failure. Patients with primary brain tumours or brain metastases should be considered for specific treatment of the tumour, which includes surgery and/or radiotherapy.

Give Dexamethasone, IV, 4 mg 6 hourly, initially.

OR

 Give Betamethasone, oral/IV, 4 mg 6 hourly. Discontinue if no response has occurred after 48 hours. Taper dose according to response and duration of therapy.

3.5.10 Brain Oedema Due to Traumatic Injury

General Measures

Refer patient for neurosurgical opinion, if indicated. Supportive management.

Note: DVT prophylaxis with heparin may be contraindicated owing to risk of increased bleeding.

The following measures should be used in patients with raised intracranial pressure:

- head elevation and position,
- airway and ventilation control,
- sedation and analgesia,
- control of fever,
- control of hypertension, and
- prevention of seizures.

Currently, no evidence supports the use of hyperventilation in this setting.

Treatment

For raised intracranial pressure, pending neurosurgical procedure only:

Give Mannitol 15–25%, IV, 0.25–1
g/kg administered over 30–60
minutes. Monitor neurological
response and urine output. Do not
repeat more than 6–8 hourly.
Beware of hypovolaemia and
electrolyte disturbances, especially
hypokalaemia.

Currently no evidence exists to support the use of hypertonic saline infusion.

Corticosteroids used in this setting have a harmful effect.

4.0 Ear Nose and Throat Conditions

4.1 Mastoiditis Secondary to Acute or Chronic Otits Media

Symptoms and signs:

- Swelling behind or above the ear.
- Fever, Pinna pushed forward, tender

Treatment: Adults:

- Give Ampicillin 1g every 8 hourly for 5 days plus
- Give Flucloxacillin 500mg IM or IV,
 6 hourly for 5 days plus
- Give Metronidazole 500mg IV, 8 hourly for 5 days
- Give **Analgesics** as necessary

Alternatively:

Give Ceftriaxone 2g daily for 5 days

Children:

• Give **Ampicillin** 25 – 50mg/kg IM or IV q8h for 5 days

4. Ear Nose and Throat Conditions

Red Flags:

- Watch for complications of brain involvement (meningitis or brain abscess).
- Surgical drainage may be necessary (If fluctuant)
- Refer patient to hospital (if no response to drugs)

4.2 Otitis

4.2.1 Otitis Externa

 This is an inflammation of the skin lining the external auditory canal.
 May be a furuncle or diffuse.

Symptoms and signs:

• Ear pain, hearing loss

Treatment:

(a) Furuncle

- Give Analgesia
- Give Flucloxacillin 500mg qid for 5 days

Alternatively

Give Cloxacillin 500mg qid for 5 days

 Make a wick of ribbon gauze impregnated with
 Hydrocortisone or Betamethasone cream and gently insert in the ear for 2 to 3 days.

(b) Diffuse Otitis Externa

- Give **Analgesia** when necessary
- Dry mop the ear
- Give Acetic acid ear drops 2% in Alcohol q6h for 5 days

4.2.2 Acute Otitis Media (Children)

 Local medicine treatment is ineffective and should be avoided

Acute otitis media is often viral in origin and needs only a simple analgesic for pain

Symptoms and Signs:

 Fever in about 50% of patients, sudden persistent ear pain or pus discharge for < 2 weeks

Treatment:

 Give Amoxycillin 15 mg/kg tds for 5 days

Alternatively:

- Give Erythromycin 6.25 mg/kg tds or Azithromycin 10mg/kg stat the 5mg/kg od for total maximum 5 days for patients with penicillin allergy
- Give Analgesia as required
- Pus discharge from the ear for over 2 weeks

Symptoms and signs:

- Persistent pus discharge, hearing loss
- If the eardrum has been ruptured for over 2 weeks, secondary infection with multiple organisms usually occurs
- Common in immunosuppressed patients
- This makes oral antibiotic therapy much less effective.

Treatment:

 Ensure ear is always dry by dry wicking with cotton wool.

Note: A chronically draining ear can only heal if it is **dry**. Drying the ear is time consuming for both the health worker and the mother but it is the only effective measure.

 The mainstay of treatment is topical therapy with Acetic acid ear drops 2% in alcohol q6h for 5 days or where available Ciprofloxacin/Chloramphenicol ear drops.

- Demonstrate/explain carefully to the patient (or guardian in the case of a child) how to dry the ear by wicking (see below)
- Refer for further assessment if no improvement after 3-4 weeks therapy
- Dry the ear by wicking
 - Roll a piece of clean absorbent cloth or cotton wool into a wick and insert carefully into the patient's ear
 - Leave for one minute
 - Remove and replace with a clean wick
 - Watch the patient/guardian repeat this until the wick is dry when removed
 - The patient/guardian should dry the ear by wicking at home at least four times daily until the wick stays dry
 - If bleeding occurs, temporarily stop drying the ear
 - Do not leave anything in the ear between treatments
 - Commercially made ear buds should be avoided in cleaning the ear
 - The patient should avoid swimming or otherwise getting the inside of the ear wet

- Re-assess weekly to ensure that patient/mother is drying the ear correctly
- Check for mastoiditis

Note: TB is an important cause of a chronically discharging ear in Malawi, condition is common in HIV infected children

4.3 Nose Conditions

4.3.1 Epistaxis

- Bleeding from the nose
- Bleeding can be bilateral or unilateral, or posterior and anterior
- Causes include trauma, repeated nose pickings, infections such as rhinosinusitis, systemic causes such as hypertension, bleeding disorders, anaemia and leukamia etc.

Treatment:

• Pinch the nose alar (wings) for 5 to 10 minutes. Let the patient lean forward and breathe through the mouth.

Alternatively:

- Apply cold pack or ice block to the forehead
- Use ribbon gauze impregnated with liquid paraffin

or

- Apply nasal packs soaked in Adrenaline
 Note: Avoid use of adrenaline in hypertensive patients
- If bleeding continues, refer to hospital

4.3.2 Vestibulitis

- Diffuse infection of the skin of the anterior nares and may occur due to frequent trauma such as occurs in constant nose picking.
- Persistent nasal discharge leads to excoriation and infection of the skin of the nasal vestibule

Treatment:

- Give Analgesia
- Give Amoxycillin 500mg tds for 5 days and
- Give Liquid Paraffin 2 drops each nose 3 times a day

4.3.3 Sinusitis

 Inflammation of one or more sinuses that occurs most often after a viral nasal infection or allergic rhinitis.

4.3.3.1 Bacterial Sinusitis

Symptoms and Signs:

 Purulent nasal discharge, persistent or intermittent, pain and tenderness over one or more sinuses, nasal obstruction, postnasal discharge, occasional fever.

Note: (1) Sinusitis is uncommon in children under five years as sinuses are not fully developed

(2) Unilateral foul smelling nasal

discharge is a foreign body until proven otherwise

Treatment:

Adults:

- Give Oxymetazoline 0.05% 2 drops twice a day for not more than one week
- Give **Cetrizine** 10mg daily for 3-5 days
- Give Amoxycillin 500mg tds for 5 days
- Steam inhalations using Menthol are advised

Children:

 Give Oxymetazoline 0.025% 2 drops twice a day for not more than one week Give Amoxycillin 25 mg/kg/dose in exacerbations of chronic sinusitis and HIV positive children who are on Cotrimoxazole prophylaxis.

Alternatively if penicillin hypersensitivity:

Give Erythromycin 12.5 mg/kg/dose
 qid or Azithromycin 10mg/kg stat then
 5mg /kg od for 7 days

If pain or fever:

 Give Analgesic/Antipyretic treatment as required

4.3.4 Allergic Rhinitis

 Recurrent inflammation of the nasal mucosa due to hypersensitivity to inhaled allergens e.g. pollen, house dust, grasses and animal proteins.

Symptoms/Signs:

- Blocked stuffy nose, watery nasal discharge, frequent sneezing often accompanied by nasal itching and irritation, conjunctival itching and watering, edematous pale gray nasal mucosa, mouth breathing, snoring at night.
- Exclude other causes such as infections, vasomotor rhinitis, over use of decongestants drops, side effects of antihypertensives and antidepressants.

Treatment:

- Allergen avoidance
- Give Cetrizine 10mg daily for 3-5 days plus
- Give Beclomethasone nasal sprays

4.3.5 Pharyngitis

 Viral pharyngitisis a painful red throat without purulence. Respiratory viruses are a major cause.

Symptoms/Sign:

- Sore throat and fever
- Diffuse congestion of the pharyngeal wall, uvula and adjacent tissues.

Treatment:

- Antibiotics are not indicated
- Homemade salt mouth washes or gargles for 1 minute twice daily

4.3.6 Tonsillitis

 Acute inflammation of the tonsils. The main organism implicated in the causation is beta-hemolytic streptococcal.

Symptoms and Signs:

 Sore throat, difficulty and pain on swallowing, inflamed tonsils, multiple white spots on the tonsillar surface, and sudden onset of fever.

4. Ear Nose and Throat Conditions

Treatment:

- Warm salt gargles
- Give **Amoxicillin** 500mg tds for 7 days

Alternatively:

- Give Erythromycin 500mg, every 8 hours in Penicillin allergy
- Give Analgesia see Section 24.1 on pain relief

4.4 ENT Emergencies

4.4.1 Foreign Body Inhalation

Symptoms and Signs:

- Drooling in saliva especially in children
- Signs of upper airway obstruction such as Difficulties in breathing, stridor, use of accessory muscles to breathe, painful swallowing.

Treatment:

- Refer for surgical review
- Retropharyngeal Abscess to be under here

5.0 Emergencies

Note: Initial Trauma Management: For all emergencies and trauma, the following format should be applied in management (ABDCE):

- A Airway
 - Jaw thrust, Guedel airway, immobilize C-Spine
- B Breathing
 - RR, O2 Therapy, Respiratory exam
- C Circulation
 - BP and HR, IV access (use green or grey cannula in adults)
- D Disability
 - AVPU or GCS (see table below)
- E Exposure
 - Examine the rest of the body (Head to toe)
 - Don't forget to check Glucose if able to
 - For suspected Fractures, aim to immobilize, sterile compress and give analgesics before referral.

RED FLAG:

- Open fractures, fractures of the long bones or pelvis.
- Risk of bleeding
- TTV for open fractures

| GCS TABLE | 1 | 2 | æ | 4 | 2 | 9 |
|--------------|-------------------|---------------------------------|---------------------------|-----------------------------|-------------------------|-------------------|
| Motor | Unresponsive | Extension to pain | Flexion to pain | Withdraws from pain | Localizes pain | Obeys commands |
| Verbal | No sounds | Incompre- hensible sounds | Words but no conversation | Confused | Comprehensive speech | |
| Eye | No eye opening | Opens to Pain | Opens to voice | Opens sponta- neously | | |

5.1 Shock

 Acute circulation failure resulting in inadequate tissue perfusion and cellular hypoxia, generally with a low blood pressure.

Causes

- Hypovolemic (haemorrhage, cholera, severe vomiting, diabetic ketoacidosis)
 - Cold, clammy skin; weak pulse, tachycardia
- Cardiogenic (myocardial infarction)
 - Signs of heart failure
- Obstructive (pericardial tamponade, tension pneumothorax)
 - Raised JVP, pulsus paradoxus
 - 4 distributive (sepsis, anaphylaxis)
 - Fever, warm peripheries

5.1.1 Anaphylaxis (anaphylactic shock)

- This is a medical emergency in which seconds count
- Taking appropriate measures immediately may save a life
- It requires prompt treatment for laryngeal oedema, bronchospasm and hypotension

5. Emergencies

- It is most commonly precipitated by drugs, especially after parenteral administration including:
 - Antibiotics
 - Aspirin and other Non-Steroidal Antiinflammatory Drugs (NSAIDS)
 - Antiarrhythmics
 - > Heparin
 - Neuromuscular blocking drugs
 - > Injectable iron
 - Vaccines
- It may also be caused by:
 - Insect stings (especially wasps and bees)
 - Blood products and blood transfusions
 - Certain foods e.g. eggs, cow's milk, nuts

General Measures:

- Adrenaline IM is a priority
- Determine and remove cause
- Lie the patient down
- Keep the patient warm
- Elevate the patient's legs
- Maintain airway
- Give 100% oxygen if available

Treatment:

- Give Adrenaline 0.01 ml/kg
- Repeat as required (several times if necessary) every 10 minutes according to BP and pulse until improvement occurs
- Give Sodium Chloride 0.9 % 20 mL/kg by IV infusion over 60 minutes
- Start rapidly then adjust according to BP
- An antihistamine is a useful additional treatment given after adrenaline and continued for 24-48 hours to prevent relapse
- Give Promethazine 25-50 mg by deep IM or, in emergencies, slow IV, as a solution containing 2.5 mg/mL in water for injection
 - > Adults: max 100 mg
 - Children:
 - o 6-12 years: 6.25-12.5 mg
 - o 1-5 years: 5 mg
- Repeat dose every 8 hours
- An IV Corticosteroid is of secondary value in initial management of anaphylaxis as its action is delayed but should be given in severe cases to prevent further deterioration:

Adults:

Give Hydrocortisone 200 mg by slow IV push

5. Emergencies

Children:

< 1 year: 25 mg 1-5 years: 50 mg 6-12 years: 100mg

- May be repeated as necessary
- Monitor pulse, BP, bronchospasm and general response/condition every few minutes
- If there is continuing deterioration or no improvement the following may be necessary:
- Give **Aminophylline** IV as for asthma if bronchospasm persists (see Section 16.2.1)
- Ventilation and/or tracheotomy
 If acidosis is severe (blood pH<7.1) after
 20 minutes

Treatment

- Give Sodium Bicarbonate 50 mmol by slow IV (appr. 100 mL of 4% solution)
- Monitor plasma pH

5.2 Management of Emergencies and Trauma in Children

- Triage all sick children soon when they arrive in hospital into three categories in order to identify:
 - a) Those with emergency signs

 Emergency signs include: obstructed
 breathing, severe respiratory distress,
 central cyanosis, signs of shock, coma,
 convulsions, severe dehydration
 - b) Those with priority signs *Priority* signs{"3TPR MOB"}, tiny baby, very hot or very cold temperature, trauma or other urgent surgical condition, severe pallor, poisoning, pain, respiratory distress, restlessness, referral {urgent}, malnutrition, oedema of the feet, burns
 - c) Those who are non urgent cases

5.2.1 Emergency Management:

- Assess the airway and breathing (A,B)
 - Does the child's breathing appear obstructed? If so open as follows
 - If no trauma do chin lift, head tilt.
 - If suspected trauma- do jaw thrust.

- Manage airway in choking by using back slaps, chest thrusts or Heimlich manoeuvre in an older child.
- If there is severe respiratory distress i.e. tachypnoea, recessions, nasal flaring, cyanosis etc.
 - Oxygen therapy
 - Ventilatory support if not breathing and there are signs of life – use bag and mask.
- Assess circulation and level of consciousness
 - Check if the child's hand is cold if so
 - Check capillary refill time

 (apply pressure to the nailbed for 5 seconds and determine the time from the moment of release until total recovery of the pink colour) normal is <3 seconds.
 - If in shock manage as follows:
 - Manage shock, coma and convulsions
 If unconscious
 - Put on oxygen
 - Check blood sugar
 - Place in the recovery position.
- Assess and manage severe dehydration in a child with diarhoea

5.3 Management of convulsions in children

- Establish the cause and when convulsions are controlled, treat accordingly
 - In neonates convulsions are usually due to hypoglycemia, hypoxia or infection e.g. meningitis
- Always exclude hypoglycemia as cause especially in children. If blood sugar can not be checked, assume hypoglycemia and treat accordingly
 - See Section 6.1.1 for treatment of this in children
 - Hypoglycaemia is defined as <2.5 mmol/l or 3 mmol/l in severely malnourished children.
- For persistent convulsions, see Section 5.2 General measures:
 - Ensure airway is clear
 - Do not place any object which might be swallowed in the mouth of a convulsing child
 - Protect patient from injury and put in lateral position

Flow chart for controlling convulsions:

- To control the fit.
 - A- Airway: place in recovery position
 - B- Breathing: support with oxygen if necessary
 - C- Circulation: treat shock
 - D- Don't
 - E- Ever
 - F- Forget

5. Emergencies

G- Glucose (correct **Hypoglycaemia** with 2ml/kg of 25% **Dextrose** or 2.5ml/kg of 20% **Dextrose** or 5ml/kg of 10% **Dextrose**)

Diazepam

0.5mg/kg rectal or 0.25mg/kg IV

If the fit has been going on more than 5 minutes

Diazepam 0.5mg/kg rectal or 0.25mg/kg IV

 \downarrow

If fit still ongoing after 10 minutes or has recurred \downarrow

If fit still ongoing after 10 minutes or has recurred $\ \downarrow$

If fit still ongoing after further 10 minutes or

Paraldehyde 0.2ml/kg IM or 0.4ml/kg rectal

Diazepam

0.5mg/kg rectal or 0.25mg/kg

has recurred, inform senior health worker



If fit still ongoing after further 5 - 10 minutes or in status epilepticus



Note:

- It is not recommended to give Diazepam intramusculary
- Rectal administration may be quicker and easier than IV in fitting child. Use a syringe with the needle removed.
- For paraldehyde use a glass syringe preferably. If not available, use a plastic syringe instead but ensure that the dose is given promptly and therefore remains in the syringe for a short time (paraldehyde dissolves plastic)

 For neonates, use Phenobarbitone 15-20mg/kg loading dose and maintenance dose of 2.5-5mg/kg once daily.

5.4 Diabetic Ketoacidosis (DKA)

- Persons at extra risk: known insulindependent diabetics with poor compliance, intercurrent infection, failure to administer insulin when ill and not eating
- Investigate immediately blood sugar; urine dipstick for glucose and ketones, electrolytes and urea if possible
- Hypoglycaemia, subdural haematoma (elderly), stroke, malaria, meningitis, sepsis may also precipitate DKA
- If blood sugar levels cannot be obtained, it may be difficult to distinguish clinically between hypoglycaemic and hyperglycaemic coma; in that case
 - give 50 ml 50% dextrose stat: in case of hypoglycaemia, it will wake the patient up; in case of hyperglycaemia, it will do no harm

Treatment:

 Fluids – use large bore cannula; if possible set up 2 cannulae

- In case of hypovolaemia i.e. blood pressure < 90 mm Hg or postural drop of blood pressure > 20 mm Hg
 - Give 4 litres Sodium chloride 0.9%
 in the first hour
 - In case of no hypovolaemia
 i.e. good urine output, normal
 blood pressure
 - Give 2 litres Sodium chloride 0.9%
 in the first hour
 - In both cases after one hour give 3 litres /24 hours + amount lost in urine (see sliding scale for type of fluid)
 - If still dehydrated after first hour, give 2 litres sodium chloride per hour and review
 - Potassium chloride 10 mmol per litre (0.75mg) in each litre until IV fluids are stopped

Note: Do not give **Potassium Chloride** with the first litre of Sodium Chloride instead.

- Give 10U Soluble Insulin IV stat, thereafter give Insulin IV according to sliding scale every 2 hours
- Check blood sugar every 2 hours
 - Give IV soluble Insulin and only change to sc if patient is well hydrated

5.5 Hyperosmolar non-ketotic coma (HONK)

 Persons at risk: elderly with non-insulin dependent diabetes

Signs and Symptoms:

 Confused / reduced consciousness (any reason e.g. stroke or infection)

Treatment:

- Fluid replacement is more important than insulin even if blood sugar is high; however fluids should not be given too rapidly to avoid large electrolyte shifts.
 - Give 2 litres Sodium Chloride 0.9% in the first hour, then 1 litre every hour. Adjust to slower rate if elderly patient with risk of heart failure.
 - Change to **Dextrose 5%** when blood sugar approaches normal levels.
- Give 10U **Soluble Insulin** IM stat; this is usually enough.
 - Do not try to lower the blood sugar rapidly at all cost by giving high doses of insulin.
- Add Potassium Chloride as indicated in Section 5.2
- Adjust / individualize insulin treatment when fully conscious and eating

5. Emergencies

Sliding scale for insulin dosage based on blood sugar taken 2 hourly (see table below)

| Blood Glucose | Dose soluble insulin (IM) | Type of fluid |
|--------------------------------------|------------------------------------|----------------------|
| HHH (very high) | 10 units | Sodium chloride 0.9% |
| >400mg/dL (22mmol/L) | 10 units | Sodium chloride 0.9% |
| 300 – 399 mg/dL (16.5 – 22mmol/L) | 10 units | Sodium chloride 0.9% |
| 200 – 299mg/dL (11 – 16.5mmol/L) | 5 units | Sodim chloride 0.9% |
| <200mg /dL (11mm | 5 units | Dextrose 5% |

6.0 Endocrine Disorders

6.1 Diabetes Mellitus

 The diagnosis of diabetes is based on 2 abnormal blood sugar measurements (FBS > 7 mmol/l, 126 mg/dl or RBS >11mmol/200 mg/dl or HbA1C >6.5%) in an asymptomatic patient or 1 abnormal measurement if the patient has symptoms of hyperglycaemia

General Measures:

- Establish treatment aims: some patients require strict glycaemic control with near normal glucose values targeted, for others symptom control and avoiding severe side effects of treatment may be the maximum achievable.
- Check BP regularly and aim for BP <130/80 mmHg
- Discourage smoking
- Educate about foot care and screen annually for foot problems (neuropathy or peripheral vascular disease)
- Screen annually for decrease in visual acuities, look for cataracts

Refer for specialist opinion if:

- Pregnant diabetic
- Acutely ill diabetic, particularly if vomiting or decreased Glasgow
- Coma Score (GCS)
- Treatable complications e.g. cataracts

Note: Suspect Diabetes Mellitus in an adult with foot ulcer not healing for long time; vaginal candida; blurred vision even without typical symptoms and signs of Diabetes Mellitus. All diabetic clinics should have access to HbA1C test.

Treatment:

- Give Aspirin 75mg daily to hypertensive diabetics aged over 50
- If a diabetic is admitted unconscious always consider the possibility of hypoglycaemiaadminister 100ml 20% or 50ml 50%
 Dextrose/Glucose IV even if a blood glucose measurement is not available.

Note: Do not rash to refer the unconscious Diabetic mellitus before you consider managing hypoglycaemia for hypoglycaemia can cause sudden death than hyperglycaemia.

6.1.1 Diabetes Type 1 (Insulin dependent)

- Most children will have type 1 diabetes.
- Children with diabetes should be referred for proper management and treatment.
- Consider hydration while waiting for transport for referral
- These children need to be followed up every 3 months to monitor blood sugars and long term complications of diabetes

6.1.1.1 Adults with no ketoacidosis or other acute complication

Treatment:

• Give Lente/Protophane Insulin 2 doses daily

Note: For proper management of diabetes mellitus refer to Clinical handbooks of QECH College of Medicine and KCH Intern Logbook as well as Pediatric handbook

- To decide the starting dose of Insulin:
 - The total daily number of Insulin units will be approximately half the patients body weight, e.g. for a 60 Kg person give 30 units Insulin/day divided into 2 doses
 - Then give 2/3 daily dose half an hour before breakfast, give 1/3 daily dose half an hour before evening meal, preferably 12 hours apart

- Adjust Insulin dose according to fasting blood sugar (FBS) or 2 hours post-prandial blood sugar or symptoms of hypo- or hyperglycaemia
- Give Metformin 500mg twice daily can be added to Insulin treatment to improve glycaemic control and curb weight gain in adult diabetics

Note: Insulin requirements can go *up* when a patient is acutely ill, even if they are not eating. NEVER stop Insulin in a type 1 diabetic.

Diet:

- Increase fibre intake
- Reduce refined sugar intake
- Insulin treated patients require 3 meals a day containing complex Carbohydrate to avoid risk of hypoglycaemia
- Advise patients to eat more before unaccustomed exercise

6.1.1.2 Children with Diabetic Ketoacidosis

Signs and symptoms:

 Vomiting, polyuria, dehydration, ketonuria and acidosis. The blood sugar will be high >15mmol/l

Treatment:

- Address airway and breathing
- IV fluids are the most important resuscitation measure
- Give Normal Saline or Ringers Lactate
- Give 10mls/kg bolus and repeat to a maximum of 30mls/kg to correct shock if present
- Ongoing fluid requirement =
 (Maintenance) plus (Deficit) minus
 (shock bolus)
- CORRECT OVER 48HRS TO AVOID CEREBRAL OEDEMA

Note: Child is usually approx. 7.5 to 10 % dehydrated. Deficit is calculated as % body weight loss. Maintenance is calculated as per shown below

Maintenance requirements are as follows:

- First 10 kg body weight 100mls /kg /day
- Next 10 kg body weight 50mls/kg/day
- Each kg thereafter 20mls/kg /day.

For example:

- Comatose child weighing 20kg on admission in shock in DKA X 10ml/kg bolus needed to correct shock = 2 X 200 = 400mls
- Maintenance is 1.5L/ day (1000mls +500mls)

- Deficit= 20kg X 7.5% = 1.5L (one litre weighs1kg)
- Requirement over 48 hours
- Maintencance (1.5 +1.5L) +deficit(1.5L)
 minus bolus (400) 4.1L +/48hours =
 85ml/hr
- Add KCl to IV fluids when patient urinates and peripheral circulation has improved.
- Change to oral K+ supplements when patient is able to feed.
- ECG monitoring if potassium is <2.8 or >6mmol/L

Give Insulin

- Should be short acting, soluble
- Start insulin one hour after starting IV fluids
- start with small subcut dose of 0.1u/kg.
 Recheck blood glucose after an hour.
- If glucose is unchanged or increased, repeat subcut dose of 0.1u/kg. Repeat hourly until blood glucose starts falling.

Sliding Scale:

- Blood glucose (mmol/L)
- >20 : 0.5u/kg
- 15 19.9 : 0.4u/kg
- 10 14.9 : 0.3 u/kg
- 5 9.9 : 0.2 u/kg
- - 4.9 : 0.1u/kg only if on a glucose drip
- <2 : omit Insulin and give Dextrose or food

Ongoing Management:

- Change IV fluid to 1/2 strength Darrows or 5 % Dextrose if blood glucose < 15mmol/L
- IV fluids must be continued until child is drinking well, tolerating oral feeds and has ketone free urine
- Monitor level of consciousness.
 Deteriorating neurological state may indicate cerebral oedema. Ensure airway is protected
- Consider NGT on free drainage if child is unconscious.
- Check each urine passed for glucose and ketones as a guide to recovery
- Maintenace Insulin requirements
 - Once child is drinking and eating
 - Calculate total daily dose of insulin once the child is stable. This is usually 0.5 to 1u/kg/day, but should be based on the Insulin requirement of the previous 24hours.
- If only short acting Insulin is available:
 - Continue tds regime prior to meals according to requirement
- If long acting is available:
 - ➢ BD regime, 2/3 of the total dose should be given before breakfast and 1/3 before dinner
 - ➤ Proportion for long acting and short acting should be about 2:1 to 3:1
- Educate patient and family on diet:
 - Importance of regular meals

- Avoid refined sugars eg SOBO, bananas, cakes and biscuits
- Encourage complex carbohydrates eg cereals and a high fibre diet
- Educate patient and family on Insulin:
 - Keep in a cool place e.g clay pot if no refrigerator
 - Rotation of injection sites
 - How to give injections

6.1.2 Hypoglycaemia

Symptoms and signs:

- Sweating
- Trembling
- Tachycardia
- Drowsiness
- Confusion

Treatment:

- Glucose in form of refined sugars eg SOBO
- Follow up
 - Attend monthly clinics at hospitalblood glucose and/or urine should be checked
 - Check injection sites
 - Ask about nocturia.
 - HbA1c can be measured
 - Diabetic patients should have the following annually: fundoscopy, urine microalbuminuria and thyroid function tests if available.

6.1.3 Diabetes type 2 (Non Insulin-Dependent)

- Adjustment of diet and/or weight reduction (if obese) and increased exercise may control blood glucose without the need for drug therapy.
- Wherever possible give a 4-6 week trial of diet before introducing oral hypoglycaemic agents. If the above is unsuccessful, then:

Treatment:

- Give Metformin 500mg twice daily, increased to a maximum of 1g twice daily.
- Note: (1) Metformin is the drug of choice in type 2 diabetes, particularly in obese patients.
 - (2) It is contra-indicated in renal insufficiency, and severe respiratory and cardiac disease due to risk of lactic acidosis.
- If glycaemic control still poor, add
 Glibenclamide 5mg daily, increasing to a maximum of 10 mg twice daily

Note: 20% of type 2 diabetics eventually require Insulin treatment - use principles as in type 1diabetes to initiate treatment. Use **Lente** 0.3U/kg bodyweight to start with.

6.2 Thyroid Disorders

6.2.1 Hyperthyroidism

Causes:

Graves disease, toxic multinodular goiter, toxic solitary nodule

Signs and symptoms:

- Fatigue, nervousness or anxiety, weight loss, palpitations, heat insensitivity, tachycardia, warm moist hands thyromegaly and tremor.
- Management should be supervised by a doctor
- Refer to tertiary level

Treatment:

Adults:

- Give Propranolol 40-120mg tds to control symptoms, especially tachycardia
- Give Carbimazole 40mg od for approximately
 2 months then reduce dose to 10mg od
- In Graves disease continue for 18 months then stop (in large percentage hyperthyroidism will be resolved)
- In other causes continue Carbimazole and refer for surgery

Children:

- Give Carbimazole 0.5 mg/kg od po
- Give **Atenolol** 1-2 mg/kg od orally

Refer for surgery if:

- Relapsed Graves disease after carbimazole treatment
- Toxic nodule or toxic multinodular goitre

Note: If a patient develops a fever or sore throat while taking Carbimazole, neutropenia should be urgently excluded. If present then stop Carbimazole and treat with antibiotics.

6.2.2 Hypothyroidism (Myxoedema)

- Management should be supervised by a doctor
- Refer to tertiary level

Treatment:

- Give Thyroxine 100-150mcg od for life
- Elderly patients start with 25mcg then increase by 25 - 50mcg every 2 weeks up to 100mcg

6.2.2.1 Congenital Hypothyroidism

- Congenital hypothyroidism is one of the common treatable causes of preventable mental retardation in children.
- Congenital hypothyroidism must be treated as early as possible to avoid intellectual impairment.

Signs and symptoms:

 Prolonged jaundice, feeding difficulties, hypotonia, wide open fontannelles, oedema, constipation, enlarged tongue, dry skin, bradycardia, lethargy etc.

Treatment:

- Give Levothyroxine 10-15 mcg/kg od orally for neonates and infants and 100 mcg/kg od
- Requires urgent referral for confirmation of diagnosis

6.2.3 lodine Deficiency Disorders (Endemic Goitre)

- More common in highland areas
- Much less likely since the introduction of iodised salt
- Only consult surgeons for treatment if large goiter causing obstructive problems or cosmetically unacceptable.

6. Endocrine disorders

Prophylaxis:

- Give Aqueous Iodine Oral Solution
 130mg/ml single dose
- Repeat every 2 years

7.0 Gastro-Intestinal Conditions

7.1 Amoebiasis

- Give health education on feacal disposal, hand-washing and food hygiene
- Consider in dysentery unresponsive to antibiotic treatment

7.1.1 Intestinal (Non-Invasive) Form

Treatment:

Adults:

 Give Metronidazole 800mg tds for 5 days preferably after food

Children:

Give Metronidazole 7.5 mg/kg tds for 5 days

7.1.2 Hepatic Form (Amoebic Liver Abscess)

Treatment:

- Give Metronidazole 800mg tds for 10 days
- Give orally (preferably after food) or IV (depending on the condition of the patient)
- If necessary, repeat treatment course after 2 weeks
- Consider aspiration in cases of large abscesses or superficial abscesses under ultrasound guidance.

Children:

- Give Metronidazole 10 mg/kg/dose every 8 hours for 10 days
- Give orally (preferably after food) or IV (depending on the condition of the patient).
- If necessary, repeat course of treatment after 2 weeks

7.2 Bacillary Dysentery

- If the only symptom is dehydration, give health education on hand-washing (the single most important preventive measure), correct faeces disposal, and food hygiene
- Ensure complete hygienic precautions by all in contact with the patient
- Isolate the patient if possible
- Investigate source of contamination and inform environmental health authorities
- Use antibiotics only if the patient is systemically unwell or septic or immunosuppressed

Treatment:

Adults:

- Give Ciprofloxacin 500 mg bd for 5 days
- Analgesic

Children > 3 months:

 Give Nalidixic Acid 50 mg/kg daily in 2-4 divided doses

7.3 Cholera

- Rehydration is of prime importance
- Ensure complete hygienic precautions by all in contact with the patient. The patient should be isolated if possible
- Investigate source of contamination, and inform environmental health authorities
- Trace close contacts and give antibiotics in the same doses as below
- Main treatment is by rehydration but antibiotics can shorten the diarrhoea episode and are therefore indicated.

Treatment:

Adults:

Give Doxycycline 300 mg stat

Alternatively in pregnancy and children <5 years:

• Give **Erythromycin** 250 mg qid for 3 days

Children >5 years:

Give Erythromycin 12.5 mg/kg qid for 3 days

7.4 Constipation

- Investigate and treat any identified cause
- Commonly related to inadequate dietary fiber intake and/or psychological factors
- Advise high residue diet, e.g. papaya seeds and increased fluid intake
- Reserve medication for severe cases only confirmed by examination.
- Do not use oral laxatives in children. Glycerine infant and pediatric suppositories can be used in the short term.
- If increased fiber and oral fluids are insufficient to cure constipation and a laxative is considered necessary use Liquid Paraffin 5-10 mls daily.
- Refer all infants with constipation for specialist assessment
- Constipation in the neonate is usually due to a significant underlying problem such as bowel atresia or Hirschsprung's disease.

If a neonate has not passed stools in the first 48 hours of life:

 Refer urgently for surgical and/or pediatric assessment especially if there is abdominal distension and/or vomiting

Note: All laxatives are contraindicated if intestinal obstruction is suspected

7. Gastro-Intestinal Conditions

Treatment:

Adults:

Give Bisacodyl 5-10mg at night

Alternatively

 Insert one Glycerol Suppository at night moisten with water before insertion.

If no response within 3-5 days:

• Refer for further management

Note: For hemorrhoids, anal fissure and other causes of persistent anal pain in adults:

Insert one Bismuth Subgallate Suppository rectally each night and morning after defecation

7.5 Diarrhoea

7.5.1 Acute Diarrhoea

- Replace fluid and electrolyte loss
- Maintain optimal hydration
- Establish and treat causal factors
- In adults with acute diarrhoea who are systemically unwell and/or have fever,
 Ciprofloxacin 500 mg bd may be considered if the patient is (suspected to be) HIV infected
- In children, if IV fluid is indicated but is impossible to administer, consider using the intra-osseus routes

Treatment:

- Give low osmolarity WHO ORS as soon as the patient's condition improves
- Measurement of BP and pulse may help in assessment of dehydration.

7.5.1.1 Use of drugs in children with diarrhoea

- Only use antibiotics for dysentery and suspected cholera cases with severe dehydration
- >6 month: Give **Zinc** 20mg per day for 10 days
- <6 month: Give **Zinc** 10mg per day for 10 days in addition to low osmolarity **ORS**
- Only use antiparasitics for:
 - Amoebiasis, after antibiotic treatment of bloody diarrhea for shigella has failed or trophozoites of E.histolytica containing red blood cells are seen in the faeces.
 - Giardiasis, when diarrhea has lasted at least 14 days and cysts or trophozoites of *Giardia* are seen in faeces or small bowel fluid

Note: Antidiarrhoeals and antiemetics should never be used in children with acute diarrhea and vomiting because they have no proven value and may be dangerous

7.5.1.2 Assessment of patients for dehydration

| | | | Α | В | С |
|----|-----------------------------|---------------|------------------------|--|---|
| 1. | Look at: | Condition | Well, alert | Restless, irritable | Lethargic, unconscious, floppy, very weak |
| | | Eyes | Normal | Sunken | Sunken and dry |
| | at. | Tears | Present | Absent | Absent |
| | | Tongue | Moist | Dry | Dry |
| | | Mouth | Drinks normally | Dry | Dry |
| | | Thirst | Not thirsty | Thirsty | Very thirsty |
| 2. | Feel: | Skin Pinch | Goes back quickly | Goes back slowly | Goes back very slowly |
| 3. | Decide: | | No sign of dehydration | If the patient has 2 or more signs including at least one sign*: | If the patient has 2 or more signs including at least one sign*: SEVERE |
| 4. | Treat (see belo w) | | Use Plan A | Weigh if possible Use Plan B | Weigh if possible Use Plan C |

Note: In severely malnourished children, skin turgor is not a reliable sign.

7.5.1.3 Treatment Plan A (to treat diarrhoea at home)

- Use this plan to teach the mother to continue to treat her child's current diarrhea at home and to give early treatment for any future diarrhea.
- Explain the 3 rules for treating diarrhea at home:

1. Give child more fluids than usual to prevent dehydration

- Suitable fluids include: Low osmolality ORS, plain water, food- based fluids (e.g. gruel, soup, rice water), breast milk
- Give ORS if the child has been on Treatment Plan B or C or cannot return to the health worker if the diarrhea gets worse
- Give ORS or water rather than a food-based fluid if the child is under 6 months old and not yet on solid foods
- Give as much of these fluids as the child will take
- Use the amount shown below
- for ORS as a guide
 - Continue giving these until the diarrhea stops

Special notes for severely malnourished children

- Dehydration tends to be overdiagnosed and its severity overestimated in severely malnourished children because it is difficult to assess dehydration in severely malnourished children using clinical signs alone.
- Do not use IV route for rehydration except in cases of shock.
- In shock give oxygen, keep the child warm, establish IV access and infuse
 15mls/kg over 1 hour of ½ Strength
 Darrows with 5 % Dextrose or Ringers
 Lactate with 5% Dextrose.
- The child needs to be observed closely and if the respiratory rate and pulse rate increases stop IV fluids.
- If the child is improving but still shocked repeat the same volume of fluids over another hour thereafter switch to oral or nasal gastric rehydration with **ReSoMal** alternatively with **F-75** for up to 10 hours.
- In severe dehydration give 5mls/kg of Resomal every 30 minutes for the
 - first 2 hours then give 5-10ml/kg/hour for the next 4-10 hours which should
 - be alternated with F-75 at the usual volumes.

Ingredients for home made ResoMal

| Ingredient | Amount | |
|---------------------|--------------------|--|
| Water | 2 litres | |
| WHO ORS | One 1 litre packet | |
| Sugar | 50 g | |
| Electrolyte/mineral | 40mls | |

- In mild diarrhea without dehydration, prevent dehydration developing by continuing milk feeds – ReSoMal is not necessary in these cases.
- Monitor carefully and frequently
- Watch for signs of heart failure due to over hydration which is common in children with kwashiorkor.

2. Give the child plenty of food to prevent malnutrition

Continue to breast-feed frequently

If the child is not breast-fed:

Give the usual milk feed

If the child is 6 months or older or already taking solid foods:

- also give cereals or another type of starchy food
- mix, if possible, with pulses, vegetables, and meat or fish
- add 1-2 teaspoonfuls of vegetable oil to each serving
- give fresh fruit juice or mashed banana to provide potassium
- give freshly prepared food
- cook and mash or grind food well to make it easier to digest
- encourage the child to eat
- offer food at least 6 times daily
- give the same food after the diarrhea stops
- give one extra meal daily for 2 weeks
 - 3. Take the child to the health worker if the child does not get better in 3 days or develops any if the following:
 - many watery stools
 - repeated vomiting
 - marked thirst
 - eating or drinking poorly

7. Gastro-Intestinal Conditions

- > fever
- blood in the stool
- If the child will be given ORS at home, show the mother how to mix ORS and how much to give after each loose stool (see
- table over)
- Give enough packets for 2 days treatment

| Age | Amount of ORS to | Amount of |
|---------|-------------------------|---------------|
| (years) | give after each | ORS to |
| | loose stool* | provide for |
| | | use at |
| | | home |
| Under 2 | 50-100 ml | 500 ml per |
| 2-10 | 100-200 ml | day |
| Over 10 | As much as wanted | 1 L per day |
| | | 2 L per day |

^{*} Describe and demonstrate the correct amount to be given using a locally available measure, e.g. cup or coke bottle

Show the mother how to give ORS:

For a child <2 years:

give a teaspoon every 1-2 minutes

For an older child:

give frequent sips from a cup

If the child vomits:

- wait 10 minutes, then give ORS more slowly If diarrhea continues after the ORS is used up:
 - tell the mother to give other fluids as described in rule 1 above or to return for more ORS

Explain how to prevent diarrhoea in child:

- give only breast milk for the first 4- 6
 months and continue to breastfeed for at
 least the first year
- introduce clean, nutritious weaning foods at 4-6 months
- give the child freshly prepared and wellcooked food and clean drinking water
- make sure all family members wash their hands with soap after using the toilet, and before eating or preparing food
- quickly dispose of the stool of young children in a latrine or by burying

7.5.1.4. Treatment Plan B (to treat dehydration)

1. Give ORS solution for the first 4 hours:

| Weight (kg) | Amount (mL) |
|-------------|-------------|
| Under 5 | 200-400 |
| 5- | 400-600 |
| 8 | 600-800 |
| 8- | 800-1,200 |
| 11 | 1,200- |
| 11- | 2,200 |
| 16 | 2,200- |
| 16- | 4,000 |
| 30 | |

- Encourage the mother to continue breastfeeding
- Give more **ORS** if the patient wants it
- For infants under 6 months who are not breast-fed, also give 100-200 ml of clean water during this period
- Observe the child carefully and help the mother give **ORS**
- solution:
- show her how much solution to give the child
- show her how to give it (see Plan A)
- check from time to time to see if there are any problems

If the child vomits:

- wait 10 minutes
- then continue with ORS, but more slowly

If the eyelids become puffy:

- stop ORS
- Give breast milk
- Give ORS again (as in Plan A) when the puffiness has gone
 - 2. After 4 hours, reassess the child using the assessment chart. Choose a suitable treatment plan to continue

If there are no signs of dehydration:

- use Plan A. When dehydration has been corrected the
- child usually passes urine and may also be tired and fall asleep

If signs showing some dehydration are still present:

- repeat Plan B, but start to offer food, milk and juice as
- described in Plan A

If signs of **severe dehydration** have appeared:

change to Plan C

If the mother must leave before completing Plan B:

- show her how much **ORS** to give to finish the 4 hour treatment at home
- give her enough ORS packets to complete rehydration and for 2 more days as shown in Plan A
- show her how to make the ORS solution
- explain to her the 3 rules in Plan A for treating the child at home

7.5.1.5. Treatment Plan C (to treat severe dehydration quickly)

Start IV fluids immediately

- If patient can drink, give ORS by mouth while the drip is set up
- Intra-ossueus route: if unable to get an IV line in quickly, consider using this useful method of rehydration fluid administration if familiar with the technique:
- If an intra-osseus needle is not available, use a 21G
- hypodermic needle or a large bore LP needle instead
- Ringer's Lactate 100 ml/kg divided doses (See Rate of Administration of IV Rehydration Fluid Table below)

Alternatively if not available

Normal saline

Note: Both of these IV solutions do not contain glucose and are thus not suitable for long-term IV fluid therapy in children – use Darrow's ½ strength + dextrose 5% instead

Rate of administration of IV rehydration fluid

| Age | First give 30 ml/kg | Then give 70ml/kg |
|------------|---------------------|-------------------------------------|
| <12 months | 1 hour * | 5 hours |
| >12 months | 30 minutes* | 2 ¹ / ₂ hours |

 Repeat once if radial pulse is still very weak or undetectable

Alternative method of IV rehydration fluid administration

- May be useful where the above schedule is difficult to follow on a busy ward
- give 10 ml/kg of the IV fluid as a bolus administered over 5 minutes using a syringe
- then give 100 ml/kg as an infusion over 6 hours

With either method:

reassess the patient every 1-2 hours

If hydration is not improving:

- give the IV fluid more rapidly
- also give ORS (approx. 5 ml/kg/hour) as soon as the patient can drink – usually after 3-4 hours (infants) or 1-2 hours (older patients)

 After 6 hours (infants) or 3 hours (older patients), evaluate the patient using the assessment chart, then choose the appropriate Plan A, B or C to continue treatment

If IV therapy cannot be given, but is available within 30 minutes and the child can take fluids orally:

 provide the mother with ORS solution and show her how to give it during the trip to the referral hospital

If IV therapy cannot be given and is not available, but nasogastric therapy is available:

- start rehydration by nasogastric tube with ORS solution
- give 20 ml/kg/hour for 6 hours (i.e. a total of 120 ml/kg)
- reassess the patient every 1-2 hours
- if the child has malnutrition, give half this amount

If there is repeated vomiting or increased abdominal distention:

• give the fluid more slowly

If hydration is not improving after 3 hours:

- refer for IV therapy
- reassess after 6 hours
- choose the appropriate treatment plan to continue treatment

If both IV and nasogastric therapy are not available, but the patient can drink:

- start rehydration by mouth with ORS solution
- give 20 ml/kg/hour for 6 hours (i.e. a total of 120 ml/kg)
- re-assess the patient every 1-2 hours

If there is repeated vomiting;

• give the fluid more slowly

If IV and nasogastric therapy are not available and the patient cannot drink:

- refer *urgently* for IV or nasogastric therapy
- If possible, observe the patient at least 6 hours after rehydration to be sure the mother can maintain hydration with ORS solution by mouth
- If the patient is under 2 years old and there is cholera in the area, treat for this after the patient is alert

7.5.2 Chronic Diarrhoea (Adults)

- Defined as LOOSE stools for 3 or more times daily or episodically for over 1 month
- Common presentation in HIV/AIDS patients.

Treatment:

Adults:

- Correct any dehydration and maintain hydration
- Consider Potassium Supplements
- Advice the patient to eat more potassiumrich foods if possible, e.g. bananas, oranges, tangerines, and other citrus fruits
- Give supplementary feeding when required/as tolerated
- Always do HIV test
- Investigate stool for presence of ova, cysts and parasites

If the condition persists

 Give Cotrimoxazole 960mg every 12 hours for 7 days

If the condition responds to treatment but recurs within 4 weeks:

- Re-treat in accordance with the initial response
 - Relapse may be due to short duration of initial treatment

7. Gastro-Intestinal Conditions

If improved after re-treatment:

Follow up as required

If still no response after adding Metronidazole or if there is no improvement after recurrence and retreatment with cotrimoxazole/metronidazole:

Give Albendazole 400mg twice daily for 2 weeks

If still not improved after albendazole treatment;

- Refer for microscopic examination of stool,
 See note (d) below
- Multiple stool examination may increase the diagnostic yield of parasites if present

If bacteria or parasite found;

Treat accordingly

If the condition responds to treatment but recurs within 4 weeks:

Re-evaluate

If bacteria or parasite is not found, use a constipating agent to control the diarrhoea for up to 5 days treatment

- Give Loperamide 4mg initially then 2mg after each loose stool
- Usual dose 6-8mg daily up to 16mg daily

Alternatively

• Codeine Phosphate 30mg every 6 hours

If still not improved within 1 week:

- Stop treatment
- Re-evaluate

If no treatable cause of the diarrhoea is found

Counsel the patient

Note:

- a. Do **not** use constipating agents in patients with bloody diarrhoea because of the risk of inducing toxic megacolon
- In persistent diarrhoea, perianal application of soft yellow paraffin may soothe anal mucosae
- c. Chronic diarrhoea caused by cryptosporidium infection and caused by HIV infection itself (HIV enteropathy) needs to be treated with antiretroviral therapy

7.5.3. Persistent diarrhoea (children)

- Defined as liquid stool 3 or more times daily for over 14 days
- A pathogen will only be identified in a small number of cases
- Consider TB and chronic infections e.g. UTI, as uncommon causes

7. Gastro-Intestinal Conditions

- Persistent diarrhoea is a common symptom of protein-energy malnutrition (PEM) and will resolve with treatment of this
- Correct any acute dehydration and maintain hydration

Maintain nutrition Including breast feeding, where appropriate Presence of fever and/or bloody stool makes bacterial infection more likely and malnutrition puts a child at increased risk of dying from persistent diarrhoea. Empiric (trial) antimicrobial treatment is therefore indicated in these conditions

If bloody stool and fever:

 Give Nalidixic Acid 50 mg/kg divided in 4 doses per day for 5 days

If malnourished give:

- Cotrimoxazole 24 mg/kg every 12 hours for 5 days and
- Supplemental feeding and supplements

If worsening, or not improving after 14 days:

- refer for microscopic stool examination
- consider giving Albendazole for worms (see Section 15.5 page 131)
- and/or give **Metronidazole** for giardiasis

7.6 Dyspepsia

 Meal related non-specific abdominal discomfort and pain

General measures:

- Advise patient to avoid hot spices, alcohol, tobacco, carbonated drinks
- Encourage regular meals

Treatment:

- Chew 2 Magnesium Trisilicate tablets every 6 hours or more frequently as required for 7days
 - > Take preferably before food
 - > Take the last dose at night

Alternatively:

 Give Ranitidine 300mg at night or 150mg every twelve hours for 4 weeks

or

- Give Cimetidine 400mg every twelve hours or 800mg at night OR
- Give Omeprazole 20mg at night for 2 weeks

7.7 Peptic Ulcer Disease/Gastritis

Commonly caused by Helicobacter pylori

Treatment (Triple Therapy):

- Give Omeprazole 40mg once daily for 2 weeks
- Give Metronidazole 400mg every 8 hours for 7 -10 days
- Give Amoxycillin 1g every twelve hours for 7 10days

Alternatively:

- Give Omeprazole 20 mg twice daily for 2 weeks
- Give Metronidazole 400mg every eight hours for 7-10 days
- Give Clarithromycin 500mg twice daily for 7-10 days

Red Flag:

- Refer for endoscopy and further management if ongoing pain and alarm symptoms
 - Weight loss, Haematemesis and signs and symptoms of gastric outlet obstruction

Note: Aspirin and other non-steroidal antiinflammatory drugs (NSAIDS) e.g. Indomethacin, Ibuprofen, are contraindicated in patients with a history of peptic ulcer

7.8 Vomiting

- Always look for a possible cause and treat accordingly
- Do not give symptomatic treatment without knowing the cause
- Always exclude mechanical obstruction
- Correct dehydration where necessary

Treatment:

 Give Metoclopramide 10 mg IM or slow IV (over 2 minutes) 3 times daily as required

Note: Patients less than 20 years require special caution. Observe dose requirements and use restrictions.

Children:

 All children with profuse vomiting must be admitted for hydration, observation and investigation. When a guardian comes back with a child who is still vomiting, admit or refer the child. Do not send them home.

8.0 Hepatic Disorders

8.1 Acute liver failure

- Withdraw the causative agent if possible e.g. drugs or alcohol
- Avoid hepatoxic drugs e.g. Paracetamol
- Ensure adequate intake of Glucose (Dextrose)
- Induce diarrhoea with Lactulose 5ml every 6 hours and/or Neomycin
 Sulphate 1g q6h (1 week)
- Give antibiotic prophylaxis for patients with liver failure: Ceftriaxone 2g od IV

Alternative: Give Ampicillin 1 g q8h

Prophylaxis of bleeding: Vitamin K 10 mg IM stat

If the patient is drowsy or comatose

- Give an IV infusion of **Dextrose** 50%
- Reduce protein intake
- Do not give sedatives or hypnotics
- Refer the patient for further diagnosis and management
- Give Thiamine 100 mg IV or IM
 3-7 days in case of known or suspected alcoholic cause

9.0 Infectious Diseases

9.1 HIV and related conditions

 People infected with HIV may develop HIV-related illnesses, the most common of which is TB.

In all cases of HIV-related illness, prompt diagnosis and proper management of the problem is crucial

For more details refer to the following MoH guidelines:

- MoH Management of HIV/AIDS Related Diseases (2008 2nd Edition)
- The Malawi AIDS case definition: Clinical and laboratory diagnosis of HIV related diseases.
- Treatment of AIDS Guidelines for the Use of Antiretroviral Therapy in Malawi
 3rd edition.
- PMTCT guidelines

9.1.1 Care and Support

General supportive care:

Proper nutrition is important.

Psychological support:

 Inform patients of any HIV support groups in their community.

Spiritual support:

• The spiritual needs of the patient and family should be properly addressed.

9.1.2 Immunisation of HIV (+) children

- Unless very ill, immunise with EPI vaccines (BCG, DPT, polio and measles) according to standard schedules
- Older children with clinical AIDS should not get BCG vaccine.
- Antibody response to vaccines may be less than normal but tends to be better in the early stages of HIV infection.

9.1.3 Breast-feeding by HIV (+) mothers

- Counsel HIV (+) women on the risks of future pregnancy
- Encourage Exclusive breast feeding up to 6months and may continue breastfeeding up to 2 years of age.
 - All lactating mothers must be fast tracked to start HAART if not yet on treatment. Refer to Option B+ on PMTCT guidelines.

9.1.4 End Organ Dysfunction

- These include:
 - Bone marrow (haematological abnormalities): see section on blood diseases
 - Central Nervous System, Gastrointestinal, Renal, Cardiac Systems
- For organ dysfunction directly caused by HIV, ART is indicated in combination with supportive treatment

9.1.5 Counselling

- Refer to the MoH Guide for Pre- and Posttest Counselling and AIDS Counselling information.
- If a child is too young, counsel the parents/guardians.
 - Counselling should be private, compassionate and confidential.
- Consider modes of transmission in discussions with parents or guardians.
- Offer HIV testing to parents of HIVinfected children, and advise them on:
 - The implications of HIV infection in themselves for further children.
 - The risk of transmitting infection sexually or as blood donors in the future.

9.1.5.1 Pre-test counselling

 Refer to the MOH Guide for Pre- and Post-test Counselling and AIDS Counselling information.

9.1.5.2 Post-test counselling

 Refer to the MOH Guide for Pre- and Post-test Counselling and AIDS Counselling information.

9.1.6 Health worker safety

- Refer to the MOH/National Prevention Services booklet *Recommended*
- Guidelines for Infection Control and Prevention

9.1.7 Post-Exposure Prophylaxis (PEP)

 PEP involves giving ARV's following possible exposure to HIV to prevent infection

- Situations where PEP should be considered:
 - Rape/sexual assault: women, men and children
 - Needle/sharps injury: usually health care workers
 - Multiple major trauma: eg. road traffic accident involving multiple passengers
 - Condom failure: especially if partner known to be infected

General Principal:

- The first dose should be given as soon as
 possible after exposure the sooner it is given,
 the more likely it is to prevent infection
- There is likely to be no benefit in giving PEP more than 72 hours after exposure
- Patients should receive a baseline rapid HIV test to confirm they are not already HIV positive
- However, if testing is not immediately
 available, the first dose can be given prior to
 testing to avoid delay
- If a patient is found to be HIV positive, PEP should not be initiated and they should be referred to an HIV clinic for ongoing care
- When the HIV status of the source is not known (usually the case in rape) they should be assumed to be HIV positive.

 When the source is known (for example a hospital inpatient) even if they are HIV negative, PEP may be given if there is a possibility that they may be in the window period.

PEP Regimens:

• Adults and children >35kg:

1st choice:

| Drug | Dose | Frequency | Duration |
|---|---------------|------------|----------|
| TDF/3TC | | | |
| Tenofovir 300mg/Lamivu dine 150mg | One tablet | Once daily | 30 days |

• Acceptable alternative if TDF/3TC not available:

| Drug | Dose | Frequency | Duration |
|---|---------------|----------------|----------|
| AZT/3TC | | | |
| Zidovudine 300mg/Lamivudine 150mg | One tablet | TWICE daily | 30 days |
| 2001118 | | | |

Children <12 years of age:

The paediatric formulation of AZT/3TC BD (Zidovudine 60mg/Lamivudine 30mg) is used twice daily, the dosing is dependent on weight:

| Weight band | Number of tablets BD |
|-------------|--------------------------------------|
| <6kg | 1 |
| 6 to <10kg | 1 ½ |
| 10 to <14kg | 2 |
| 14 to <20kg | 2 ½ |
| 20 to <25kg | 3 |
| 25 to <35kg | Use <i>adult</i> AZT/3TC 1 tablet BD |
| >35kg | Use TDF/3TC (see adults) |

For cases of rape:

 Always offer emergency contraception as soon as possible to girls and women of reproductive age, unless protected by a reliable form of contraception.

| Contraceptive drug | Immediately | After 12 hours |
|-----------------------------|-------------|-------------------|
| Postinor-2 | 2 tablets | - |
| Lo-Feminal or Microgynon | 4 tablets | 4 tablets |

 Always offer presumptive treatment for STI's, though this is less urgent than PEP and emergency contraception, and can be deferred (particularly the oral drugs) for a few days if necessary.

| Drug | Child <15 years | Adult |
|---------------|------------------|---------------|
| Benzathine | 50,000 IU/Kg IM | 2.4 mega unit |
| Penicillin | stat | IM stat |
| Gentamycin | 7.5mg/kg IM stat | 240mg IM |
| | | stat |
| Erythromycin | 12.5mg/kg QDS 7 | 500mg QDS 7 |
| | days | days |
| Metronidazole | 5mg/kg TDS 7 | 400mg TDS 7 |
| | days | days |

9.1.8 HIV infection in children

- Common presentation: diarrhoea, failure to thrive (FTT) and infections including tuberculosis, thrush (moniliasis), pneumonia, and meningitis;
- Treat these and malnutrition appropriately.
- Immunise HIV(+) children with DTP, hepatitis B, Hib, polio, and measles vaccines, unless they are very ill but, do not give BCG to older children with clinical AIDS.
- Encourage mothers to continue breast feeding.
- Whenever possible, treat HIV(+) children as out-patients to minimise the risk of crossinfection, as they may be immunocompromised.

Note: A reactive HIV ELISA test in a child under 12 months, even with no laboratory error, is not proof of HIV infection, as passively acquired maternal antibody may be the cause

9.1.9 Management of common clinical presentations in HIV (+) patients

- Treatment of these is often similar to the treatment of the same condition in HIV(-) patients, as covered elsewhere in these guidelines.
- Differences in treatment for HIV(+)
 patients are indicated in the
 appropriate treatment schedules.

9.1.9.1 Diarrhoea Disease

See Section 7.5

9.1.9.2 Respiratory Conditions (Adults)

- Common HIV associated respiratory diseases: bacterial pneumonia, pulmonary tuberculosis, Pneumocystis jirovecii pneumonia (PCP), lymphoid interstitial pneumonitis (LIP), pulmonary Kaposi's sarcoma.
- Cough without dyspnoea or tachycardia and associated with a runny nose is usually indicative of a viral URTI.

9.1.9.3 Pneumocystis Jirovecii Pneumonia (PCP)

Treatment:

- Give Cotrimoxazole po or IV 30 mg/kg q8h for 21 days
- Wherever possible give oxygen
- Give severely dyspoenic patients
 Prednisolone 40mg bd for 5 days then 40mg od for 5 days then 20mg od for the remaining 11 days
- Risk of recurrence after treatment is very high. For prophylaxis give
 Cotrimoxazole 960mg twice a day

9.1.9.4 Respiratory Conditions (children)

- Cough with or without difficulty in breathing.
- Pneumonia and pulmonary TB are common in HIV (+) children.
- Cough without dyspnoea or tachypnoea and associated with a runny nose is usually due to a viral URTI.

Treatment:

- If upper respiratory tract infection (URTI) with fever: exclude malaria (see Section 15.1)
 - ➤ If URTI without fever: Advise mother on general supportive home care (see Section 16.1)
- If Pneumonia suspected: assess severity and treat accordingly

- Presumed Pneumonia which fails to respond to treatment may be due other causes
 - Do a chest X-ray to rule out emphysema, effusion, TB, PCP, LIP, staphylococcal infection and other conditions common in HIV(+) patients
 - Manage according to X-ray findings
- If PCP found, treat as in Section 9.1.9.3 above
 - Give Prednisolone, 1-2mg/kg every 12 hours
- If LIP found:
 - Suspect LIP if chest xray shows a bilateral reticular –nodular interstial pattern. Distinguish from pulmonary TB.
 - The child may present with persistent cough, bilateral parotid swelling, persistent generalized lymphadenopathy, hepatomegaly and finger clubbing.
 - Antibiotics for bacterial Pneumonia
 - Give Prednisolone 1-2mg/kg daily for 2 weeks,
 - Decrease dose over 2-4 weeks depending on response.
 - Beware of reactivation of TB

9.1.9.5 Fever (adults)

- A body temperature of over 38⁰ C, continuously or intermittently, for more than 24 hours in any 72 hour period.
- Fever is common in HIV(+) patients Seriously ill patient:
 - Start treatment for presumed sepsis (See Section 9.7)
 - Maintain hydration
 - Refer urgently to hospital for further management

If patient not seriously ill:

- Maintain hydration
- Check bloodslide for malaria parasites, whether positive or negative:
 - give presumptive 1st line antimalarial treatment (see Section 15.1.1)
- Look for local causes of fever: otitis media, tonsillitis, skin infections, pneumonia, PTB and EPTB, urinary tract infections, joint infections and give appropriate treatment accordingly

- Consider giving antipyretic treatment *If not improved within 3 days:*
 - Refer for further investigations
 - Treat according to findings

If there are no suggestive laboratory or radiological findings:

- Consider empirical (trial) treatment for suspected sepsis
- (see Section 9.7)

If fever still persists but patient is clinically stable:

- Presume HIV related fever
- Give supportive care and assess for ART
- Seek a second opinion at the earliest opportunity

9.1.9.6 Fever, Persistent or Recurrent (children)

- Persistent fever: a body temperature of
 >38 degrees Celcius for more than 5 days
- Recurrent fever: a body temperature of >38 degrees celcius for more than 1 episode in a period of 5 days
- Look for: Meningitis, septicemia, Occult bacterial infections, TB, Fungal, viral or parasitic infections, Neoplasms
- Maintain hydration
- Maintain nutrition
- Give antipyretic treatment

Treatment:

Seriously ill child:

- Start treatment for presumed sepsis (see Section 9.7)
- Start 1st line antimalarial treatment (see Section 15.1.1)
- Refer the patient.

Child not seriously ill:

If the child has completed 1st line antimalaria treatment:

- Give Amoxycillin 15mg/kg every 8- 12 hours for 7 days.
- This is intended to treat non-serious bacterial infections, e.g. sinusitis, urinary tract.

If free of fever after 3 days:

- Complete treatment course for Amoxycillin
- Follow up as required.

If not improved:

- Give 2nd line antimalarial treatment (see Section 15.1.1)
- Follow up as required.

If the child has not completed 1st line antimalarial treatment

• Give 1st line antimalarial treatment (see Section 15.1.1)

If not free of fever after 3 days:

• Refer to the next level

9.1.9.7 Upper GIT Candidiasis

- Presumptive diagnosis:
 - White plagues in the mouth and lesions in the esophagus and stomach
 - Antifungal therapy is required treat according to Oropharyngeal conditions (see Section 14.1)

9.1.9.8 Mental disorders (adults and older children)

• Refer to Chapter 25 on Mental Disorders

9.1.9.9 Primary neurological disorders (adults)

- Refer for careful assessment and investigation
- Refer to Chapter 3 on Central Nervous System

9.1.9.9.1 CNS disorders

 Look for evidence of an opportunistic infection and treat the underlying cause accordingly

9.1.9.9.1.1 Protozoal, Viral, Fungal and Bacterial Infections

- Often affects the brain in HIV (+) patients
- Carry out a lumber puncture
- Refer to relevant treatment guidelines

9.1.9.9.1.2 Cerebral Toxoplasmosis

- Most frequently causes focal neurological signs in patients with advanced immune-suppression.
- Sulphadoxine/Pyrimethamine (SP) 2 tablets daily for at least 6 weeks, followed by lifelong cotrimoxazole 480mg twice daily.
- Start ART after some weeks on SP.
- May result in complete cure

9.1.9.9.1.3 Cryptococcal Meningitis

• Refer to Section 9.3.3

9.1.9.9.1.4 Tuberculosis

Refer to Section 9.5

9.1.9.9.1.5 Syphilis

Refer to Section 17.4

If there is severe spasticity and ataxia:

- Presume myelopathy
- Treat as neuro-syphilis with Benzylpenicillin 5
 MU IV 6 hourly for 2 weeks OR Doxycycline
 200 mg od for 3 weeks

 Give supportive and symptomatic treatment and counseling

9.1.9.9.1.6 AIDS Dementia

 Characterised by cognitive, behavioural and motor dysfunction, often overlooked in advanced HIV infection. ART is indicated and may give a favorable response.

9.1.9.9.1.7 Progressive Multifocal Leucoencephalopathy (PML)

- Caused by a polyomavirus (JC) virus, rapid evolution over weeks or months; altered mental state, visual defects, motor weakness, speech dysfunction, sensory deficits and cerebellar disorders.
- Prognosis is very poor, sometimes temporary relief from ART, no causative treatment available.

9.1.9.9.2 Peripheral Nervous System Conditions

If predominantly sensory: sensory peripheral neuropathy

- If on tuberculosis treatment, treat with Pyridoxine 25mg od for the duration of tuberculosis treatment
- Check if drugs that cause neuropathy are used (vincristine, stavudine, metronidazole) and consider modifying drug treatment
- Screen for diabetes mellitus and treat accordingly if present
- Consider other causes of neuropathy: alcohol abuse, renal disease, malignancies, vitamin B12 deficiency and treat if possible
- HIV associated peripheral neuropathy often improves with ART
- Give supportive and symptomatic treatment:
 - Give Amitryptilline 25-75mg nocte
 - Give painkillers if required (see Section 24.1)

9.1.9.10 Primary neurological disorders (children)

- Manage these in the same way as in HIV (-) children.
- Refer to hospital for careful assessment and investigation.
- Rule out treatable causes of acute episodes of neurological dysfunction such as TB, bacterial meningitis and cerebral malaria through careful history and examination in children.

9.1.9.11 Lymphadenopathy

Causes:

- Tuberculosis, bacterial (including syphilis), fungal or viral infections, malignancies (Kaposi's sarcoma, lymphoma), dermatological and other conditions.
- Persistent generalized lymphadenopathy (PGL), more than 3 separate lymph node groups affected, at least 2 nodes more than 1.5 cm in diameter at each site, duration of more than 1 month and no local or contiguous infection which might explain the lymphadenopathy is common and due to HIV infection alone; requires no treatment

General Measures:

 Ensure careful physical examination to identify any local or contiguous infection which might explain the lymphadenopathy.

If there is local or contiguous infection:

Treat as indicated

If TB is suspected:

Do fine needle aspiration for acid fast bacilli.
 Treat accordingly (see Section 9.5)

If there is a papulo-squamous rash and/or evidence of recent genital ulcer (adults only)

- Do a TPHA or RPR
- If positive, treat for syphilis see Section 17.4

If the patient has recent symptomatic lymphadenopathy of uncertain aetiology or if patient does not respond to empiric therapy:

 Refer for further assessment including lymph node biopsy

9.1.9.12 Failure to Thrive (FTT)

- This is seen by examining the child and checking the under-5 card
- When assessing FTT, it is vital to take a detailed dietary and social history to determine whether the child receives sufficient calories and a balanced diet
- Rule out TB and HIV
- Ascertain whether the child has any TB contacts
- Do chest x-ray and tuberculin PPD test, and also sputum test in an order child
- Tuberculin PPD testing is however usually unhelpful in severe FTT
- Identify any other associated problems and treat accordingly. These may include
 - Persistent Diarrhoea
 - Oral thrush
 - Respiratory conditions

- Assess the severity of FTT
 - Inability to feed and severe apathy are important indicators of severity
 - Severe malnutrition is often associated with extensive oedema and dermatitis

If able to feed and not severely malnourished:

- Give a trial of home feeding
- If possible, give exact recommendations on the schedule and content of the diet
- Assess the availability of food at home
- Encourage the mother to provide, whenever possible, a balanced nutritious diet including, for example soya, beans, groundnuts, bananas, other fruits, eggs, meat and fish
- Review after 2-4 weeks

If TB is not found

- Carry out other investigations as indicated from history and examination e.g.
 - Stool analysis and urea and electrolytes

If neither TB or other conditions are identified

Presume HIV-related FTT

If not able to feed and moderately or severely malnourished:

- Admit the patient
- Treat for malnutrition (see Section 23)

If the child is taking adequate oral feeds

- Continue these for 2-3 weeks
- Reassess the patient

If the child is unable to take adequate oral feeds, use NGT feeding until able to take oral feeds

Once the child is improving

- If aneamic, give Ferrous Sulphate and Folic Acid
- If helminth infestation is suspected, treat with **Albendazole**

If on re-assessment the child is not improving, refer to the higher level

Where severe FTT is considered to be HIV-related

- Improve nutritional and supportive care
- Prolonged NGT feeding may occasionally be appropriate
- Initiate HAART

9.1.9.13 Pain relief

• Refer to Section 24

9.1.10 Antiretroviral Treatment Regimens

9.1.10.1 First Line regimen

- Introduction of the first line regimen includes:
 - Staging and management of HIV related diseases
 - Group counseling on issues surrounding ART
 - Individual counseling and assessment of contraindications for ART
 - Baseline weight
 - Other blood tests may be done (full blood count, CD4 count, creatinine, liver enzymes) but are not mandatory

Note: ART eligibility for patients diagnosed with HIV

 All patients with HIV who are not already on ART should be assessed to determine if they require ART, and initiated or referred without delay.

Universal eligibility, regardless of stage or CD4 count:

- Pregnant/breastfeeding women
- Children <5 years of age

Adults and children >5 years:

- WHO Stage III or IV
- CD4 count <500

9.1.10.2 First line ART regimens

Adults and Children >35kg: regimen 5A

| Dose | Frequency |
|---------------|------------|
| | |
| One tablet | Once daily |
| | |
| | |
| | One |

- Children <35kg: regimen 2P
- Regimen 2P consists of AZT/3TC/NVP (Zidovudine 60mg/Lamivudine 30mg/Nevirapine 50mg).

Note: NVP requires to be initiated once daily for the first two weeks to minimise toxicity. This is achieved by giving AZT/3TC in the morning and AZT/3TC/NVP ir the evening for the first two weeks.

| Weight band | Number of tablets bd AZT/3TC/NVP |
|-------------|--------------------------------------|
| <6kg | 1 |
| 6 to <10kg | 1 ½ |
| 10 to <14kg | 2 |
| 14 to <20kg | 2 ½ |
| 20 to <25kg | 3 |
| 25 to <35kg | Use <i>adult</i> AZT/3TC 1 tablet bd |
| >35kg | Use TDF/3TC (see adults) |

Note:For detailed guidance on the management of adverse reactions to ART, diagnosing and managing treatment failure and switching to second line therapy, please refer to the *Malawi Integrated HIV Guidelines.*

9.1.11 Cotrimoxazole Prophylaxis

Adults

Give Cotrimoxazole 480mg bd to any HIV infected person in WHO clinical stage II, III or IV; any person with a CD4 count <500 cells/mm³ regardsless of symptoms; HIV + pregnant women after the first trimester

Children

- Give Cotrimoxazole 6-8mg/kg once daily to all HIV exposed children from 4-6 weeks till HIV infection has definitely been ruled out, and to all
 - > HIV infected children.
- If allergic to **Cotrimoxazole**, give **Dapsone**.

9.2 Leprosy

9.2.1 Multibacillary

- The Multi-Drug Treatment (MDT) regimen consists of:
- Monthly supervised doses of Rifampicin and Clofazimine taken on a fixed day at 4 week intervals
- Daily unsupervised doses of Clofazimine and Dapsone
- Continue treatment until 24 monthly supervised doses of Rifampicin and Clofazimine have been completed within a maximum of 3 years

9.2.1.1 MDT regimen for multibacillary leprosy

MDT regimen for multibacillary leprosy

| Medicine | Frequency | Age years/dose (mg) | | |
|------------|-----------|---------------------|-------------|--------------------|
| | | 0-5 yrs | 6-14 yrs | 15 yrs and over |
| Rifampicin | Monthly | 300 | 300 | 600 |
| Clofazimin | Monthly | 100 | 200 | 300 |
| Clofazimin | Daily | 25* | 50 | 50 |
| Dapson | Daily | 25 | 50 | 100 |

^{*25} mg capsules of clofazimine are not available. Give a 50 mg capsule every second day instead

9.2.2 Paucibacillary

- The MDT regimen consists of :
 - A monthly supervised dose of **Rifampicin** taken on a fixed day at 4-week intervals
 - > A daily unsupervised dose of **Dapsone**

Continue treatment until 6 monthly-supervised doses of **Rifampicin** have been completed within a maximum period 9 months

MDT regimen for paubacillary leprosy

| Medicine | Frequency | Age years/dose (mg) | | |
|------------|-----------|---------------------|------|------|
| | | 0-5 | 6-14 | 15 + |
| Rifampicin | Monthly | 300 | 300 | 600 |
| Dapsone | Daily | 25 | 50 | 100 |

9.2.3 Leprosy reactions

- Do not stop therapy for leprosy during treatment of reactions
- Refer urgently to Leprosy Control Assistant

9.2.3.1. Severe reversal reaction; paucibacillary patients

 This is a delayed hypersensitivity allergic reaction manifesting itself as exacerbated leprosy skin lesions and/or enlarged tender nerves with or without nerve deficit

Treatment

 Give Prednisolone daily with the dose being gradually reduced every 2 weeks until a total of 12 weeks are completed (see table)

Prednisolone regimen for paucibacillary severe reversal reactions

| Week | Prednisolone dose (mg daily) |
|-------|---------------------------------|
| 1-2 | 40 |
| 3-4 | 30 |
| 5-6 | 20 |
| 7-8 | 15 |
| 9-10 | 10 |
| 11-12 | 5 |

9.2.3.2 Severe reversal reaction: multibacillary patients

 This regimen is for multibacillary patients with a severe reversal reaction or recent nerve damage

Treatment

 Give Prednisolone daily being gradually reduced at intervals of 2 or 4 weeks over a period of 20 weeks (see table below)

Prednisolone regimen for multibacillary severe reversal reaction

| Week | Prednisolone dose (mg daily) |
|-------|---------------------------------|
| 1-2 | 40 |
| 3-6 | 30 |
| 7-10 | 20 |
| 11-14 | 15 |
| 15-18 | 10 |
| 19-20 | 5 |
| | |

9.2.3.3 Severe type 2 reaction

- This is an antigen-antibody complex reaction manifesting itself as fever, malaise and general body pain, and also frequently and typically with erythema nodosum leprosum (ENL)
- Rarely is iridocyclitis or orchiditis present

Treatment

- Standard short-course of daily Prednisolone with the dose being gradually reduced by 5mg every 2 days until a total of 12 days is completed (see table below)
- Do not stop therapy for leprosy during treatment of reaction
- Refer urgently to Leprosy Control assistant

Prednisolone regimen for severe type 2 reaction

| Day | Prednisolone dose (mg daily) |
|-------|---------------------------------|
| 1-2 | 30 |
| 3-4 | 25 |
| 5-6 | 20 |
| 7-8 | 15 |
| 9-10 | 10 |
| 11-12 | 5 |
| | |

9.2.3.4 Acute Dapsone Allergic Reaction

Symptoms/Signs:

- Itching, rash, exfoliative dermatitis or Stevens-Johnson syndrome
- Refer urgently to Leprosy Control Assistant
- Stop Dapsone
- Then observe

Treatment

Give Antihistamines, steroids or hospitalize depending on severity.

9.3 Meningitis

 Refer patient to hospital as soon as diagnosis is suspected

Note: A lumbar puncture is essential for diagnosis

 If possible, do a lumber puncture first and send the CSF in a sterile container along with the patient, but always start treatment before transfer

Treatment

Adults

- Give Benzyl Penicillin 5MU IV or IM stat
 - Plus (if available) Chloramphenicol 1g IV stat

Children

Give Benzyl Penicillin 100,000 units/kg IV or IM stat

- Plus **Chlorampenicol** 25 mg/kg IV stat *Neonates*
 - Give Benzyl Penicillin and Gentamycin IM or IV
 - When a lumber puncture cannot be done prior to referral, this should be done as soon as possible after admission

9.3.1 Bacterial meningitis

- In hospital, start an IV infusion for antibiotics using **Dextrose 5%** (not more than 50 mL/kg per day for an infant) and continue until oral medication can be tolerated
- Give antibiotics for at least 14 days, if there is a good response in meningococcal disease stop at 7 days

Treatment

Adults (empirical treatment pending test results)

• Give **Ceftriaxone** 2g IV q12h

Alternatively

- Give **Chloromphenical** 1g IV q6h
- Give Benzyl penicillin 5MU IV q6h

Note: use the alternative regime only when ceftriaxone is not available and evidence of culture and sensitivity of the invading organism to Chloramphenicol.

Children

Give Ceftriaxone 100mg/kg IM or IV q24h for 7 days

Alternatively

- **Chloramphenicol** 25mg /kg IV q8h plus
 - Benzylpenicillin 100,000 IU/kg q6h IV/IM

9.3.2 Meningitis in neonates

- Usually caused by gram(-) organisms and requires treatment for 21 days gram(+) infection). Otherwise give 14 days treatment
- Careful observation is essential
- While awaiting culture and sensitivity results or if they are not available:
- Give Benzylpenicillin 100,000 units/kg q6h, initially slow IV or IM plus Gentamycin 2.5 mg/kg IM or IV q8h or Gentamycin 5mg/kg IM q24h if 1 week old neonate.Or Gentamycin 7.5 mg/kg q24h if over a week old neonate.

Alternatively

• Give **Ampicillin** 50 mg/kg q6h initially IV then later IM as an alternative to **Benzylpenicillin**

If still febrile after 48 hours:

Add Cefotaxime

- Confirm diagnosis with Indian ink stain,
 Cryptococcal antigen and/or culture of the CSF.
- Refer for specialist management with
 Amphotericin B 0.7-1.0 mg/kg daily infused in
 Glucose 5% IV solution over 4-6 hours for 2
 weeks if tolerated, followed by Fluconazole 400
 mg od for 6 weeks and then 200 mg od lifelong.

9.3.3 Cryptococcal Meningitis

Alternatively if available

• Fluconazole 1200mg od po or IV q24h daily for 2 weeks, followed by Fluconazole 400 mg od for 6 weeks and then 200 mg od lifelong.

Children

 Give Fluconazole 12mg/kg for 10 weeks, then 10mg/kg ongoing. Therapeutic lumbar punctures may be indicated every 48 hours if any signs of increased intracranial pressure.

9.3.4 Meningococcal Meningitis (Prophylaxis)

 Recommended for selected groups living in very crowded conditions and for close household contacts

Adults

• Give Ciprofloxacin 500mg stat.

Alternatively

Give Doxycycline 300mg stat

Children

• Give **Doxycycline** 6 mg/kg

9.4 Tetanus

Immunization has significantly reduced the incidence of this

9.4.1 Adult Tetanus

- Good nursing care of the heavily sedated patient is essential
- Give active immunization against tetanus after recovery

General measures

- Nurse the patient in a quiet area
- Maintain adequate hydration and nutrition
- Prevent aspiration of fluid into the lungs
- Clean and debride necrotic wounds thoroughly
- Change from parenteral to oral medication as soon as possible
- Avoid provoking spasms
- Encourage active exercise after spasms have ceased

Treatment

Give Diazepam 20 mg IM or IV
 Chlorpromazine 50 mg IM or IV given alternately q3h

Alternatively

 Give Diazepam infusion 40 mg in one liter of IV dextrose or normal saline q6h to q8h

Note:

- Take care: respiratory depression may occur
- Dose sizes or frequencies of the above medicines can be increased if necessary to control spasms
- Give Anti-tetanus serum 20,000 units IV stat
- Give this after a test dose of 1,500 units s/c
- Benzyl penicillin 2 MU IV q6h every for 7 days
- Give Metronidazole 500 mg IV q8h or 400 mg tds for 7 days
- Give Tetanus toxoid vaccination: give the full course

9.4.2 Neonatal Tetanus

Symptoms:

- Poor feeding, constipation, stiffness and spasms
- Prevent if possible and aggressively treat respiratory complications as these are the main cause of death
- Start active immunization against tetanus once the child has recovered

General measures

- Nurse the baby in an intensive care area with close observation and attention to airway, temperature and spasms
- Maintain adequate hydration, initially with IV fluids
- Maintain nutrition with expressed breast milk via an NGT
- Have a mucous extractor or other suction available
- Avoid IM injections as much as possible by use of alternative routes (e.g. NGT, rectal administration) where indicated
- Change from IM injections to oral medication as soon as possible and keep handling to a minimum in order to avoid provoking spasms
- Thoroughly clean the umbilical area

Treatment

- Give Paraldehyde 0.2 ml/kg IM or 0.4 ml/kg rectally (see note below) followed by Phenobarbitone 15 mg/kg loading dose stat initially then10 mg IV plus 5 mg IM
- Continue with **Diazepam** 0.5 mg/kg by NGT or rectally (see note below)
- or slow IV and Phenobarbitione 5-10 mg/kg by NGT or IM
- Give these drugs alternatively every 3 hours
- Give Anti-tetanus serum 10,000 units IM or IV q6h for 5 days

Once spasms are controlled;

• Give **Phenobarbitone** 5-10 mg/kg od as maintenance dose

Note:

Paraldehyde: dissolves plastic so use a glass syringe. However if this is not available, use a plastic syringe but make sure the drug is given promptly and not left in the syringe before administration. The drug may also be given rectally using a syringe after removing the needle.

Diazepam: rectal administration (by syringe after removing the needle) is as reliable as IV and easier and safer to give.

9.4.3 Tetanus Prevention

- Promote Tetanus Toxoid Vaccination (TTV)
 in pregnant women and all women of child
 bearing age (see Section 12.3)
- Ensure adequate surgical toilet plus passive (ATS 1,500 units IM or S/C) and active (TTV) immunization after wounds, bites and burns

9.4.4 Tetanus Toxoid Vaccination (TTV)

9.4.4.2 Fully immunized but last booster >10 years ago:

Give one booster dose of 0.5 mL S/C or IM

Note: Fully immunized patients who have had a booster within the last 10 years do not need treatment with tetanus antitoxin (ATS) or tetanus toxoid vaccination (TTV)

9.5 Tuberculosis (TB)

- For detailed information on the control and management of TB refer to the *National Tuberculosis Control Programme Manual* (MOH, 2012)
- Suspect pulmonary TB if coughing for 3 weeks or more, usually with one or more of the following:
- Fever, Chest pain, Shortness of breath, Loss of weight and Hemoptysis

Symptoms/Signs:

- Enlarged lymph nodes (lymphadenopathy)
- Swelling of abdomen due to fluid (ascites)
- Tender swelling of the back (TB spine)
- Sometimes weakness of the legs (TB spine)
- Stiffness of the neck (meningitis)
- Confusion (meningitis)

- If TB is suspected, refer the patient to hospital
- Diagnosis and classification of pulmonary TB must be based on *sputum examination* (smear microscopy for acid-fast bacilli)

Note: Do not start TB treatment until a firm diagnosis has been made

- Effective treatment of TB and prevention of the development of medicine resistance depends on an appropriate combination of at least 2 medicines taken:
 - Regularly
 - > In the correct dose
 - > For the full recommended duration
 - Under direct observation (DOT)
 - HIV positive patients can be successfully treated for TB

9.5.1 Anti-Tuberculosis Medicines

- Malawi is now treating TB patients using fixed dose combinations (FDCs), single tablets and streptomycin injection as shown below:
- Combination Tablets

Adult Formulations

- RHZE contains: Rifampicin 150mg, Isoniazid
 75mg, Pyrazinamide 400mg, Ethambutol
 275mg
- RHE contains: Rifampicin 150mg, Isoniazid
 75mg, Ethambutol 275mg

• RH contains: Rifampicin 150mg, Isoniazid 75mg

Paediatric Formulations

- RHZ contains: Rifampicin 60mg, Isoniazid
 30mg, Pyrazinamide 150mg
- RH contains: Rifampicin 60mg, Isoniazid 30mg

b) Single tablets

- Z (pyrazinamide) contains:Pyrazinamide 400mg
- E (ethambutol) : Ethambutol 400mg,
 Ethambutol 100mg
- H100 (isoniazid) contains: Isoniazid
 100mg

c) Injections

• S (Streptomycin): Streptomycin 1g

9.5.2. TB Treatment Regimens

9.5.2.1 Treatment Regimen 1 (for new patients)

Initial intensive phase:

- Newly diagnosed TB patients are admitted for 2 weeks in hospital where they receive daily treatment under DOT.
- The remaining 6 weeks of the intensive phase is taken daily either in hospital or in the community according to the patient's DOT option.

 In central hospitals, patients are started on ambulatory treatment depending on the condition of the patient from the first day, but treatment is on daily basis just like the district hospitals.

Continuation phase:

- Patients take medicines under supervision.
- Medicines are collected from the nearest health facilities every fortnight.

Dosages of FDC formulations

| Adults | | | | | |
|-------------------------|---|---|--|--|--|
| Rody woight | Initial phase (2 months) | Continuation phase (4 months) | | | |
| Body weight (in kgs) | [RZE] [R150/H75/Z400/E275] Number of tablets* | [RZE] [R150/H75] Number of tablets* | | | |
| 30-37 | 2 | 2 | | | |
| 38-54 | 3 | 3 | | | |
| 55-74 | 4 | 4 | | | |
| 75 and over | 5 | 5 | | | |

| CHILDREN | | | | |
|-------------------------------|-----------------------------|---|------------------|--|
| Body weight in kg | Initial phase (2 months) | Conti- nuation phase (4 months) | | |
| | [RZE] | Е | [RZE] | |
| | (R60/H30/Z150) | 1 | (R60/H30) | |
| | Number of tablets | 0 | Number | |
| | or sachets* | 0 | of tablets or | |
| < | 1 | 1 | | |
| 8-9 | 1.5 | | | |
| 10-14 | 2 | | | |
| 15-19 | 3 | | | |
| 20-24 | 4 | | | |
| 25-29 | 5 | | | |
| Re-adjust dose as body weight | | | | |

9.5.2.2 Treatment Regimen 2

Indications for use: relapse, return after default, treatment failure and recurrent Tuberculosis.

- The regimen is 2SRHZE/1RHZE/5RHE
- Regimen consists of 2 months of SRHZE daily, 1 month of RHZE daily followed by 5 months of RHE daily, all under supervision.

Note:

- Sputum positive cases that have previously taken anti-tuberculosis medicines for 1 month or more must be suspected of discharging tubercle bacilli resistant to one or more anti-TB medicines.
- These patients must submit sputum specimens for medicine sensitivity testing before starting the re-treatment regimen.

Treatment regimen 2

| ADULTS | | | | | | | | | |
|-------------------------|--|---|---|----------|--|---|---|--|-----|
| Body weight in kg | Initial phase 2 months [SRHZE] | | Continuation phase 5 months [RHZE] [R150/H75/Z400/E275] | | | | | | |
| | Streptomycin daily injection for 2 months | | [RHZE] Number of tablets daily for 2 months | | [Z] Number tablets dail 5 month | y for | Nu | [RHE] 150/H75/E27 Imber of tab 5 months | 75] |
| 30-37 | 0.5g | | 2 | | 2 | | | 2 | |
| 38-54 | 0.75g | | 3 | | 3 | | | 3 | |
| 55-74 | 1g | | 4 | | 4 | | | 4 | |
| 75 and over | 1g | | 5 | | 5 | | 5 | | |
| CHILDREN | | <u> </u> | | | ' | | | | |
| | | Initial phase daily doses for months [SRHZ] and E [RHZ] and E | | aily dos | se for | Continuation daily dose 5 months [I [R60/H30/] E100 | for RH] and E | | |
| i I | Streptomycin daily | [RHZ] Number of tablets per day | r of Number of Number of tablets tablets | | [E] Numb tablet per da | S | [RH] Number of tablets per day | [E] Number of tablets per day | |
| < 7 | 15mg/kg | 1 | 1 | | 1 | 1 | | 1 | 1 |
| 8-9 | 15mg/kg | 1.5 | 1.5 | | 1.5 | 1. | 5 | 1.5 | 1.5 |
| 10-15 | 15mg/kg | 2 | 2 | | 2 | 2 | | 2 | 2 |
| 15-19 | 15mg/kg | 3 | 3 | | 3 | 3 | | 3 | 3 |
| 20-24 | 15mg/kg | 4 | 4 | | 4 | 4 | | 4 | 4 |
| 25-29 | 0.5mg | 5 | 5 | | 5 | 5 | i | 5 | 5 |

9.5.3. Tuberculosis Meningitis

- The regimen for adult and childhood cases of tuberculosis meningitis is different from above.
- The regimen is 2SRHZ/7RH and doses are as below

Dose regimen for tuberculosis meningitis

| Weight in kgs | Daily during weeks 1-8 | | | y during ks 9 – 32 |
|---------------|------------------------|------|------|-----------------------|
| | S | RH | Z | RH |
| Over 55 | 1 g | 4 | 4 | 4 |
| 40 – 55 | 0.75g | 3 | 3 | 3 |
| 25 – 39 | 0.5 | 2 | 2 | 2 |
| 20 – 24 | 15mg/kg | 1.5 | 1.5 | 1.5 |
| 15 – 19 | 15mg/kg | 1.5 | 1.5 | 1.5 |
| 9 – 14 | 15mg/kg | 1 | 1 | 1 |
| 5 – 8 | 15mg/kg | 0.5 | 0.5 | 0.5 |
| 0 – 4 | 15mg/kg | 0.25 | 0.25 | 0.25 |

 The regimen consists of 2 months of Streptomycin, Rifinah and Pyrazinamide given under supervision on a daily basis followed by 7 months of daily Rifinah.

9.5.4 Medicine Resistant Tuberculosis (MRT)

- Two types of medicine resistant TB:
- Primary resistance (resistance in newly diagnosed TB cases) and
- Secondary resistance (resistance in previously treated TB cases).
- Multi-Drug Resistant TB (MDR-TB) resistant to both rifampicin and isoniazid.
- Extensively-Drug Resistant TB (XDR-TB) also resistant to any fluoroquinolone, and
 at least one of the three injectable second
 line anti-TB medicines (Capreomycin,
 Kanamycin and Amikacin)

9.5.4.1 Treatment of Medicine-resistant TB

- MDR-TB treatment requires use of second line anti-TB medicines which have to be taken for 24 months.
- The patients are managed in their communities.

MDR-TB Treatment Regimen

| INTENSIVE PHASE: 6 MONTHS | | | | |
|---------------------------|---------------------|---------|--|--|
| Patient weight | Medicine | Dosage | | |
| | Kanamycin | | | |
| | (Km) | 750 mg | | |
| | Ethionamide | | | |
| | (Et) | 500 mg | | |
| | Pyrazinamide (Z) | 1000 mg | | |
| | | 600 mg | | |
| < 50 KGS | Ofloxacin (Of) | | | |
| | Cycloserine | | | |
| | (Cs) | 500 mg | | |
| | Kanamycin | 1000 mg | | |
| | Ethionamide | 750 mg | | |
| 50 - 65 kgs | Pyrazinamide | 1500 mg | | |
| | Ofloxacin | 600 mg | | |
| | Cycloserine | 750 mg | | |
| | Kanamycin | 1000 mg | | |
| | Ethionamide | 750 mg | | |
| >65 kgs | Pyrazinamide | 2000 mg | | |
| | Ofloxacin | 800 mg | | |
| | Cycloserine | 750 mg | | |

| CONTINUATION PHASE: 18 MONTHS | | | |
|-------------------------------|-------------|--------|--|
| Patient weight | Medicine | Dosage | |
| < 50 kg | Ethionamide | 500 mg | |
| | Ofloxacin | 600 mg | |
| | Cycloserine | 500 mg | |
| | | | |
| | Ethionamide | 750 mg | |
| 50 – 65 kg | Ofloxacin | 600 mg | |
| | Cycloserine | 750 mg | |
| >65 kg | Ethionamide | 750 mg | |
| | Ofloxacin | 800 mg | |
| | | | |

 For details see Guidelines for the Programmatic Management of Multi-medicine Resistant Tuberculosis in Malawi (MOH, 2012).

9.5.5 Management of Anti – TB Drug Reactions

The table below gives an outline of symptoms of the common drug reactions and how to manage them.

Minor side effects not requiring treatment to be stopped:

| Symptoms | Medicine | Management |
|----------------------------|---|---|
| Abdominal pain, nausea | Related to Rifampicin | Give oral medicines to the patient last thing at night |
| Burning of the feet | Related to Isoniazid peripheral neuropathy | Continue Isoniazid; give Pyridoxine 50 mg - 75 mg daily large doses of Pyridoxine may interfere with the action of Isoniazid (wherever possible, Pyridoxine 10 mg daily should be given routinely with isoniazid) |
| Joint pains | Related to Pyrazinamide | Continue Pyrazinamide ; use Aspirin or nonsteroidal anti-inflammatory medicine |
| Red urine | Related to Rifampicin | Reassure the patient |
| Women on Rifampicin | Rifampicin may reduce the effectiveness of the Oral contraceptive pill | Alternative contraception should be provided |

Major Side effects requiring treatment to be stopped:

| Symptoms | Medicine | Management |
|--|---|--|
| Deafness | Related to Streptomycin | Otoscopy to rule out other causes STOP Streptomycin if no other explanation. |
| Dizziness | If true vertigo and nystagmus, related to Streptomycin | STOP Streptomycin ; If just dizziness with no nystagmus, try dose reduction for one week, but if no better stop Streptomycin instead USE Ethambutol instead |
| Generalised reactions including shock, purpura | May be due to Rifampicin, Pyrazinamide and/ or Streptomycin | STOP all medication Use different combination of medicines |
| Jaundice | Related to drug- induced hepatitis. | STOP all antituberculosis medicines until jaundice and liver function tests revert to normal (see below) |
| Skin itching | Related to all anti- tuberculosis Medicines | STOP antituberculosis medicines |
| Visual impairment | Related to Ethambutol | Visual examination –seek clarification STOP Ethambutol |
| Vomiting ++/ confusion | Suspect drug- induced hepatitis | Urgent liver function tests (Ifts) If Ifts not available, STOP anti- tuberculosis medicines and observe. |

9.5.6 Complications of Tuberculosis and their Management

9.5.6.1 Pulmonary Tuberculosis

- Haemoptysis (coughing blood):
- If severe, refer to hospital for bed rest, sedatives and probably transfusion.
- Do a chest x-ray to rule out potentially treatable conditions such as aspergilloma or bronchiectasis.

Pleural effusion (collection of fluid between the lungs and the chest wall):

- Drain big pleural effusions to relieve symptoms of dyspnoea.
- Empyema (collection of pus alone) should be drained.

Spontaneous pneumothorax (collection of air between the lungs and the chest wall):

- This may cause sudden onset of shortness of breath.
- May require admission to hospital for drainage with an underwater seal.

Fibrosis (scarring) of the lungs:

- This may lead to cor-pulmonale (right-sided heart failure) in the long term.
- Symptomatic treatment may be required.

Bronchiectasis:

- Coughing due to residual lung damage sometimes with expectoration of large volumes of sputum and sometimes blood.
- Symptomatic treatment may be required provided that the sputum is negative for AAFB.

9.5.6.2 Extra-Pulmonary Tuberculosis

- Complications will depend upon the site of the disease:
- Tuberculosis of the spine: paraplegia (weakness of the lower limbs). Refer to hospital immediately.
- TB meningitis: cranial nerve damage. Give steroids.
- TB lymphadenitis: cold abscesses and suppurating fistulae. Do I &D.
- Pericardial effusion: heart failure. Give high doses of steroids.
- Cardiac tamponade (distress associated with shock). Refer to tertiary level.
- Pleural effusion: respiratory failure. Do therapeutic pleural tap.
- Tuberculoma: focal seizures or neurological signs. Order CT scan for diagnosis
- **TB Abdomen**: ascites. Do ascitic tap for diagnosis. Finding of lymph nodes on ultrasound scan of the abdomen can also be diagnostic.

9.5.7 Use of Anti-Tuberculosis Medicines in Special Situations

9.5.7.1 Pregnancy and Reproductive Health

- Streptomycin is potentially ototoxic and may cause deafness in babies.
- **Streptomycin** should **not** be given in pregnancy.
- Rifampicin stimulates formation of liver enzymes and therefore can reduce the effectiveness of the oral contraceptive pill.
- Advise patients on TB treatment to take alternative contraception while on Rifinah.

9.5.7.2 Renal impairment and renal failure

 Give normal dosage of Rifampicin, Isoniazid and Pyrazinamide to patients with renal failure.

Note:

Streptomycin and **Ethambutol** are given in reduced doses and less frequently in patients with renal failure.

9.5.7.3 Liver impairment and liver failure

- Isoniazid, Rifampicin and pyrazinamide are hepatotoxic.
- Patients with active liver disease who develop TB should not receive Pyrazinamide or Rifampicin.
- Give Streptomycin, Isoniazid and Ethambutol for intensive phase of treatment, and Isoniazid and Ethambutol for maintenance treatment.

Liver impairment with or without jaundice has a wider differential in patients on TB treatment

DIFFERENTIAL DIAGNOSIS

| Potential causes | Clinical clues and investigations |
|---------------------------------|--|
| TB IRIS in liver | 2-4 weeks after ART |
| | Fever, tender hepatomegaly |
| | Disproportionate rise in ALP/GGT |
| | ± jaundice |
| Viral hepatitis | Hep B,C and A |
| Alcohol or alternative remedies | History |
| | Signs of chronic liver disease |
| Sepsis associated cholestasis | Fever, clear source of infection, |
| | cholestasis without tender hepatomegaly |
| Acute cholangitis | Jaundice, right upper quadrant pain, |
| | obstructed biliary tree ± stones |
| HIV | CD4 < 100 |
| cholangiopathy | RUQ pain ±fever,diarrhoea,cholestasis ± jaundice |
| | USS Abdo:structuring and dilatation of |
| | bile ducts |
| Fatty liver | Firm hepatomegaly,± tenderness |
| | USS Abdo:hyperechoic texture |

It is often very difficult to manage patients with TB accompanied with severe liver impairement. The safest is to always stop all Hepatotoxic agents (Rifampicin, Isoniazid and Pyrazinamide) and refer patient to a tertially level

- If jaundice is acute and severe, treat initially with just Streptomycin and Ethambutol.
- Give Streptomycin, Isoniazid(find out from TB office whether this is in the protocals because Isoniazid is equally hepatotoxic, there is need for an alternative regime) and Ethambutol for intensive phase of treatment, and Isoniazid and Ethambutol for maintenance treatment.
- If jaundice is acute and severe, treat initially with just Streptomycin and Ethambutol.

9.5.7.4 Epilepsy

- Rifampicin reduces plasma levels of Phenobarbitone.
- Advise patients to increase the dose of Phenobarbitone.

9.5.7.5 TB/ART

- Rifampicin reduces plasma levels of Nevirapine by 30%.
- Low Nevirapine levels may increase the risk of the HIV becoming resistant to the medicine and thus compromise the effectiveness of the ART.

9.5.8 Corticosteroids and Tuberculosis

 Adjunctive therapy with corticosteroids, in conjunction with anti-TB medicines, may be appropriate in tuberculosis meningitis, pericardial and pleural disease.

- Prednisolone 40mg daily for 30 days in TB meningitis, patients with altered consciousness, neurological defects or spinal block, followed by a gradual reduction in dose in the succeeding weeks.
- Prednisolone 60mg for 4 weeks for pericardial effusion and constrictive pericarditis, followed by 30 mg for the next 2 weeks and tapering to zero over the next 2 weeks.
- Prednisolone 40mg daily for 1 2 weeks for large pleural effusion..

9.5.9 Management of Household Contacts of Smear-Positive TB Cases

9.5.9.1 Children aged 6 years and over

- Investigate for TB if symptoms are present
- Treat if TB is present.

Note: National TB Program is advocating for active case finding for all household members.

9.5.9.2 Children below 6 years:

- Screen all children using either clinical assessment, or tuberculin test or chest x-ray including those who are household contacts of smear-positive TB cases.
- Commence child on Isoniazid Preventive
 Therapy (IPT) (5mg / kg daily for 6 months) if there is no evidence of active TB.

suggestive of TB, refer for further assessment.

- If a child contact of any age has symptoms suggestive of TB, refer for further assessment.
- Register and treat the child for TB according to the National TB Program guidelines if diagnosed with TB.

9.5.9.3 Babies born to mothers with smearpositive Pulmonary TB

- Isoniazid (5 mg / kg daily for 6 months).
- Vaccinate with BCG at the end of 6 months.
- Continue breast-feeding.
- Should child develop symptoms while on Isoniazid Preventive Therapy, investigate for active TB.
- If TB is diagnosed, stop isoniazid and institute anti-TB treatment according to the guidelines.

9.6 Typhoid

- Prevent through clean water, improved sanitation and health education
- Diagnosis should be done by blood culture, Widal tests are inaccurate!

Treatment

 Give Ciprofloxacin 500mg oral or 400 mg IV q12h 14 days treatment the same for children

- Give Ceftriaxone 2g IV q24h for 14 days
- Switch to oral Ciprofloxacin when improving and patient able to tolerate

If severe

- IV treatment is preferred
- Continue for a total of 14 days, switch to oral
 Ciprofloxacin when improving and patient is able to tolerate oral medicines

Supportive measures:

- IV *fluids* may be needed
- Ensure meticulous hand washing and proper stool disposal
- Disinfect with chlorine
- Maintain good nutrition
- Give analgesic treatment for pain relief (see Section 24)
- Intestinal perforation and intestinal bleeding are complications
- Refer urgently for surgical attention if suspected

9.7 Sepsis

 Sepsis is a condition in which infection (mostly with bacteria) causes a systemic inflammatory response resulting in severe illness

- Try to find the cause and treat accordingly;
 where possible blood culture should be
 done before starting treatment
- Sepsis is common in HIV infected patients and is mainly caused by Pneumococcus and nontyphoidal Salmonella
- Always refer to hospital for systemic treatment, but in severely ill patients before referral give:

Adults

- Give **Chloramphenicol** 1g IV or IM stat *plus*
- **Gentamycin** 240 mg slow IV or IM stat *plus*
- Quinine 1200mg IV in 5% dextrose over 4 hours

Children

- Give Benzyl penicillin 50,000 units/kg IV or IM stat plus
- Give **Gentamycin** 7.5 mg/kg slow IV or IM stat plus
- Give Quinine 10 mg/kg IM stat
- The following hospital treatment regimens are empirical and should be amended accordingly based on the results of culture and sensitivity testing

Hospital treatment:

Adults:

Ceftriaxone 2g IV q24h for 10 days

Alternatively

- Ciprofloxacilin 400 mg IV every q12h or 500 mg orally bd *plus* Benzylpenicillin 2MU IV q6h
- Switch to oral Ciprofloxacin 500 mg bd plus
 Amoxycillin 500 mg tds, or oral Co-amoxiclav
 625 mg tds, when improved
- Antibiotics should be given for a minimum of 10 days

Note: use the alternative only with evidence culture and sensitivity

If intra-abdominal source suspected:

 Add Metronidazole 500 mg IV or 400 mg orally tds. If still febrile after 72 hours reassess the patient

Children

- Give Gentamicin 7.5mg/kg daily IV plus
- Benzyl penicillin 50,000 units/kg every 8 hours, initially slow IV later IM

Alternative to **Benzylpenicillin**

- Give Ampicillin 40 mg/kg IV q6h until oral medication can be tolerated then
- Give Amoxycillin 250 mg tds
 If still febrile after 72 hours:
 - Change to Ceftriaxone 100mg/kg once daily for Neonates
 - Give Benzyl penicillin or Ampicillin/Amoxicillin as for (older) children above plus
 - Give **Gentamycin** 2.5 mg/kg q8h.

If severe:

• Give **Ceftriaxone** 100mg/kg IV q24h

Causes: varicella zoster virus

More severe and extensive in HIV (+) patients

General Management

- Application of wet compresses or use of cool baths may help to control itching, (scratching may lead to disfigurement)
- In severe itching, give a sedative systemic antihistamine (e.g. oral **Promethazine** or **Chlorpherniramine**).
- Keep hands clean and nails clipped short to reduce problems caused by scratching

Treatment

Adults (symptomatic treatment):

- Apply Calamine + Sulphur 2% lotion nocte or 2 times daily
- Give Paracetamol 500mg every 4 hours
- Has more complications in adults, in particular those that are immunocompromised. Varicella pneumonitis is particularly serious. Therefore antiviral treatment is advised. Acyclovir 800 mg 5 times a day for 7-10 days.

10.0 ONCOLOGY

- Cancer is unregulated cell division with metastatic potential.
- In general there are two broad groups of cancers: solid and haematological malignancies.
- Solid malignancies include carcinomas and sarcomas, which can affect any part of the body.
- Haematological malignancies include leukaemias and lymphomas.
- The aetiology for cancer is multifactorial (extrinsic and intrinsic).
- Risk factors include radiation exposure, chemicals, viruses and genetic abnormalities such as mutation of the p53 gene; tobacco smoking being a single risk factor.
- General cancer prevention strategies need to target risk factors.

10.1 Cancer Diagnosis and Registration

Clinical presentation:

- Initial presentation is painles mass, whilst pain is a feature of advanced disease.
- Patients may present with non healing ulcer, a fixed mass and cachexia.
- Cachexia, recurrent fevers and night sweats may be a feature in advanced disease or leukemia.

Physical examination findings:

 An ulcer with irregular edges, easily bleeds, necrotic on any part of the body. For a mass, it could be fixed, not usually tender and limits the function of the organ involved.

Laboratory tests:

- Full blood count –elevated or decreased parameter in the blood, malignancy should be suspected. E.g: WBC >30; Hb >25mg/dl would suggest malignancy
- Tumour markers –elevated titers of betaHCG, CEA, AFP suggest particular cancer. Tumour markers may also be used to monitor response to cancer treatment. Use of these need to be treated cautiously.
- Histology is important confirmatory test for diagnosis.

Radiological tests:

- Mammography for breast cancer
- CXR for detection of lung cancer or boneforming tumours
- Scanning (USS/CT/MRI) investigating particular sites for cancer involvement.

Haematology, Cytology, Histology:

- Peripheral blood smear in haematological cancers
- Cytology for solid tumors
- Bone marrow aspirate and trepine
- Immunohistochemistry where available

Note:Currently cytology and histology services are done at Kamuzu Central Hospital in Lilongwe (catering for Central and Northern Region) and Queen Elizabeth Central Hospital (catering for the Southern Region).

Haematological services (peripheral blood films, bone marrow biopsies and interpretation etc..) are also available at Kamuzu Central Hospital and to an extent also at QECH.

Tests to undertake when specific tumours are suspected:

- Cervical biopsy for histology/or PAP smear, when cervical cancer suspected or in HIV positive females
- PSA, when prostate cancer is suspected, where possible prostrate biopsy
- HIV test, when HIV-related cancers (KS, cervical cancer, lymphoma) are suspected or diagnosed
- Barium swallow, endoscopic tissue biopsy for tissue analyses, when cancer of oesophagus suspected

- Breast uss, mammography, biopsy when breast abnormality or breast cancer strongly suspected
- Colonoscopy for suspected GIT malignancies
- Fecal occult blood for suspected GIT malignancies or screening for the same
- Sputum cytology for lung cancer

Cautions:

- Biochemistry tests do not give definitive diagnoses for cancers. They usually indicate the likelihood of having a particular type of cancer. Should be followed up with confirmatory tests such as cytological or histological diagnoses.
- Most radiological tests also do not give definitive diagnoses for cancers. They usually indicate the likelihood of having a particular type of cancer.
- Tumour markers are also produced by some normal cells in the body and levels may also be significantly elevated in non cancerous conditions. Results need to be interpreted with caution.
- Mammograms meant for screening purposes have high false-positive and false-negative results and are associated with overdiagnosis, overtreatment and radiation exposure.

10. Oncology

- Most diagnostic tests for cancer are not currently available at primary and secondary level healthcare in Malawi. With the exception of Kaposi's sarcoma, cancer diagnosis and treatment should be limited to central hospitals; suspected cancer patients should therefore be referred to central hospitals.
- Although CT is an important diagnostic tool, it has the potential to cause cancer, just like other sources of ionizing radiation.

Cancer Registration:

- Population based cancer registry, representing the Malawi National Cancer Registry
- Hospital based cancer registries at QECH, KCH, Zomba and Mzuzu Central Hospitals.
- Registers all cancer cases, which is helpful for cancer epidemiology and surveillance.

10.2 Treatment Options:

General Measures

- Treatment for cancers is multimodality.
- This includes surgery, radiation and chemotherapy, hormonaltherapy
- Most of the advanced cancers require all the treatment modalities.

- For better treatment outcome, multidisciplinary treatment planning conferences are encouraged.
- Targeted therapy exists but remains expensive

Treatment options depend on:

- Patient's preference
- Cost of treatment
- Stage of disease
- Grade and histological type of cancer
- Co-morbid conditions (hypertension, DM, COPD)
- Performance status (general condition of the patient)
- Age of patient
- Organ function (haematological, liver, renal, cardiorespiratory)

10.3 Most Common Cancer in Malawi

10.3.1 Kaposi's sarcoma

- Currently the number one cancer in men and second most common in females after cancer of the cervix.
- Presentation: mucocutaneous and visceral lesions.

- Mucocutaneous lesions usually manifest as red, purple or brown papules or plaques on the skin or mucous membranes.
- Visceral organs such as lungs or GIT and effusions in serous body cavities.

Diagnosis:

- Do a biopsy for histology.
- FNA is not helpful.
- Clinical diagnosis only without histology may be acceptable at a secondary care facility.
- Do HIV test, if this status is unknown.

Treatment:

- Depends on the extent of disease; early stage can resolve if the patient is on ART alone.
- All patients with HIV associated KS should receive ART.
- Local therapies: radiation and surgery are local therapies applied to KS lesions in general. Radiation therapy is reserved for disease that is limited but causing severe pain, bleeding and distress. Surgery is reserved for aggressive KS locally which is causing severe disfigurement and organ malfunction and overwhelming sepsis. In this case organ amputation is necessary, followed by systemic chemotherapy.

- Systemic therapy: several cytotoxic chemotherapy agents are effective in providing rapid improvement in majority of patients. Chemotherapy is indicated in patients with locally aggressive and disseminated KS.
- Vincristine 1.4mg/m²{max 2mg} once a week for six doses; if there is good effect continue with 2mg once every two weeks for six doses. If no effect refer to specialist for dual therapy. Dual therapy: BV (Bleomycin 15IU/m² IV and Vincristine 1.4mg/m² {max 2mg} every two weeks, maximum 20 cycles. Triple therapy: ABV (Adriamycin [Doxorubicin] 30mg/m², IV; Bleomycin 15IU/m² IV and Vincristine 1.4mg/m²). The cycle is repeated every 21 to 28 days for 8 cycles and in patients with good performance status.
- Second line treatment: Paclitaxel 120mg/m² every 2 weeks.
 - Second line treatment must be given by specialists.
- Monitoring of treatment for the first three cycles is important.
 - No treatment response, consult specialists.

Note: This is the only cancer that may be treated at secondary level healthcare (District Hospital).

 Where oncology facilities/capacity exists vincristine monotherapy should be discouraged

10.3.2 Head and neck cancers

- These involve the following malignancies:
 - Oral cavity cancers (tongue, floor of mouth), cancers of the oropharynx, nasopharynx, hypopharynx and larynx.

Presentation:

Depends on specific location of the cancer.
 Non healing ulcers in the mouth, which easily bleed, associated difficulties in mastication and progressive dysphagia. Persistent cough and hoarseness of voice in advanced cases.

Diagnosis:

 Done in conjunction with ENT surgeons and pathologists. Radiological imaging forms part of work up (e.g. CT Scan)

Treatment:

- Depends on the extent of disease.
 - Local therapies: Early stage disease

 surgery is the treatment of choice.

 For advanced cases, chemoradiation is more preferred to surgery for cosmesis and organ preservation as first line treatment.
 If recurrence, then surgery is treatment of choice.
 - Drugs of choice: Methotrexate,
 Adriamycin [Doxorubicin],
 Bleomycin, Vincristine, Cisplatin,
 Paclitaxel, 5Fluorouracil.
 - Follow up with CT scans with contrast of the head and neck to look for recurrences every six months.

10.3.3 Breast Cancer

 The commonest malignant tumors on this site is ductal Carcinoma. Sarcomas and lymphomas may also affect the breast.

Prevention and early detection

 It is advisable for women above age 20 years to do Self Breast Examination (SBE) monthly for potential masses.

- Women of reproductive age group should have clinical breast examination (CBE) by a nurse or clinician every 6 months.
- Mammography for all women of 50 years of age though recommended in countries like USA by United States Public Service Task Force (USPSTF) is not cost-effective for Malawi therefore is not feasible for population based program.

Diagnosis

Definitive diagnosis is through histology. This is more superior than cytology which should be understood as a preliminary diagnostic test.

Treatment

- Treatment for this condition is multimodality, involving:
- Surgery: mastectomy and axillary lymph node clearance of only level I and II nodes.
 Note: No role for lumpectomy or breast conserving surgery in state hospital due to lack of radiotherapy or associated delays with foreign referral for radiotherapy.
- Chemotherapy: This may be given as neoadjuvant (before surgery) or adjuvant (after surgery). The first option has advantage of down-staging to make difficult to operate tumors operable. There are many possible Chemotherapy protocols for instance AC, CAF, CMF, TAC, TC, PF. (Cisplatin/5FU)

- However, the first line will be AC in those with normal cardiac function and those otherwise or elderly CMF.
- Second-line Paclitaxel
- Third-line PF
- For Metastatic disease consideration of chemotherapy for younger patients or **Tamoxifen** in elderly patients who may not tolerate chemotherapy.
- Hormonal treatment: the commonest drug used is Tamoxifen 20 mg daily for 5 years. However recent evidence supports 10 years. The side-effects include hot-flushes, risk of cataract formation, thrombosis, and increased risk of endometrial carcinoma. Patient on this therapy have to be monitored by a qualified cancer expert. Other drugs in this category where Tamoxifen is contraindicated is Anastrazole 1 mg daily.
- Radiotherapy: this is recommended as adjuvant post-surgery to minimize recurrence and improves survival from breast cancer. Only patients that have been assessed to have no lung, liver or bone disease should be referred for this modality.

10.3.4 Oesophagus cancer

Prevention and early detection

- Risk factors are alcohol and smoking.
- Other factors include poor seed storage with Aflatoxin contamination.
- Patients presenting with unexplained dysphagia or odynophagia should promptly be investigated.

Treatment options

- Oesophagectomy
- Radio-chemotherapy (not locally available, consider in fit patients)

Note: All patients who are not fit for above curative treatment should be considered for:

- Endoscopic Stenting
- Feeding tube
- By-pass surgery
- Chemotherapy
- Radiotherapy (not locally available, so not feasible in patients with metastases or poor performance status).

Role of chemotherapy in cancer of the oesophagus

- Neo-adjuvant in good performance patients especially lower third tumours.
- Concurrent chemo-radiotherapy (currently not feasible in Malawi)
- Palliative to relieve dysphagia or other symptoms where patient doesn't prefer stenting.

10.3.5 Cervical cancer

Prevention and early detection

- Screening for any woman of reproductive age group above 25 years using VIA.
- The high HIV prevalence may not support age
 35 as onset for screening.
- Vaccination with HPV vaccine in girls between 9 years to 14 years. As this manual was being prepared cervarix by GSK and Gardasil by Merck were on the market.

Diagnosis

- Speculum and cervical punch biopsy for histological analysis.
- Speculum examination before any antibiotic course for women presenting with abnormal vaginal bleeding or foul smelling discharge.

Treatment

- Total radical hysterectomy with lymph node dissection
- Radiotherapy: EBRT and brachytherapy
- Radio-chemotherapy plus brachytherapy
- Palliative radiotherapy (not a feasible option on public referral system).
- Chemotherapy with Platinum, fluorouracil and Taxanes may be used.
- Palliative care is critical in providing pain control with morphine (see section on pain control).

Note: Assessment of this disease needs to be done at a tertiary institution with close discussions between Gynaecologists and Clinical Oncologists to jointly stage and decide on treatment.

Consider discussing with Oncology team for chemotherapy in the following patients:

- PV bleeding or excessive vaginal discharge
- Intractable pain on Morphine
- Symptomatic metastatic disease

10.3.6 Prostate cancer

- Lower prevalence rate of Prostate cancer in Malawi can be attributed to mis-diagnosis and lower life expectancy as to compared to the global prevalence rate of 7.9% of all cancers and 15% of male cancers.
- PSA screening is not recommended as there
 is no strong evidence in its favour. Where
 PSA is being ordered for diagnosis ensure
 there is no UTI or patient has not had recent
 para-rectal exam as these may give a false
 rise in PSA.

Treatment for early prostate cancer

- Prostatectomy
- Radiotherapy

Treatment for advanced or metastatic prostate cancer

- Orchidectomy
- Hormonal therapy: Bicalutamide 50 mg daily
- Chemotherapy may be used (Taxotere) in patients failing on hormonal treatment.

10.4 Haematological Malignancies

- Currently lymphoma is the commonest haematological malignancy in Malawi:
 - Aggressive NHLs such as HIV associated diffuse large cell lymphoma (DLCL)
 - Burkitts Leukaemia/lymphoma (BL/L).
- The incidence of other haematological conditions in Malawi such as leukemias is difficult to establish because of diagnostic challenges.

Diagnosis

- Peripheral blood film examination
 - All abnormal FBC results need peripheral blood film (PBF) examination.

- Patients with haematological malignancies may present with cytopenias (reduced blood cell counts) or cytoses (increased blood cell counts), lymphadenopathy, splenomegaly and/or hepatomegaly.
 - These may result in fevers, severe anemia and/or bleeding.
- Many other common conditions such as infections (e.g. TB) and solid malignancies will present with these features.
- Commonly patients with haematological malignancies are put on antibiotics for bacterial infections or on TB treatment which fail to resolve their clinical problems and, unless corrected in time, may delay appropriate treatment beyond "curative" stages.

• Bone marrow examination

Although we are only able to do morphological examination of bone marrow aspirates and biopsies in Malawi, bone marrow examination contributes significantly to the diagnosis of hematological conditions in our environment.

- In cases where leukemia patients present with cytopenia(s), bone marrow examination enables a diagnosis to be made.
- Lymphoma, although not a primary problem of the bone marrow, may involve the bone marrow.
- Other malignancies may also spread to the bone marrow.

Histopathology of other tissues

- The diagnosis of lymphomas also commonly requires biopsy of other tissues such as lymph nodes.
- Flow cytometry
- Flow cytometers are used for CD4 and CD8 enumeration in the care of HIV infected patients in the country.
- Used to diagnose leukemias from peripheral blood or bone marrow aspirates.

10.4.1 Lymphoma

- Refer patient to an oncology centre after diagnosis is made
- If available, refer patient with current FBC, U+Es and LFT results
- Basic staging investigations include CXR and abdominal ultrasound.

Chemotherapy

- Aggressive lymphomas have a reasonably good response to a combination of chemotherapeutic agents:
- Cyclophosphamide, Doxorubicin, Vincristine and Prednisolone given once every three weeks (CHOP 21).
 - Before starting patients on Doxorubicin, a cardiac ultrasound is required as Doxorubicin is cardiotoxic.
 - Intrathecal chemotherapy may be required depending on patient presentation

10.4.2 Acute Leukemias

- Acute leukemias are among the most aggressive hematological conditions.
- The chemotherapy available in Malawi is very limited in scope.
- PBF examination facilitates morphological diagnosis which may or may not be sufficient to reach a definitive diagnosis in terms of subtypes of acute leukemias. Treatment is depended on subtypes.

 Acute leukemias typically require four months of hospital admission in selfcontained patient rooms. For acute lymphoblastic leukemias, induction, intensification and maintenance treatment cycles last for a total of 2 years.

10.4.3 Chronic Leukemias

- Chronic lymphocytic leukemia (CLL) and chronic myeloid leukemia (CML) are manageable to some extent even in Malawi. Both can be diagnosed with reasonable certainty based on clinical presentation and PBF examination.
- Chlorambucil for CLL is not available in public hospitals but can be bought in private pharmacies. Cyclophosphamide can also be used for CLL.
- Hydroxyurea can be used for patients that are too poor to afford tyrosine kinase inhibitors. For patients with CML that can afford to have a Philadelphia chromosome test done but can not afford a tyrosine kinase inhibitor, a patient assistance program called Glivec® International Patient Assistance Program (GIPAP), makes imatinib available to such patients in Malawi.
- Potentially GeneXpert technology can provide relatively cheap means of performing molecular testing in Malawi for CML using Xpert BCR-ABL cartilages.

10.4.4 Multiple myeloma

- Multiple myeloma (MM) appears to be relatively rare in Malawi, perhaps due to younger population age structure than resource-rich countries.
- Diagnosis of MM relies on a combination of clinical presentation such as bone pain, morphological examination of bone marrow and basic investigations for the CRAB (calcium, renal disease, anemia and bone disease) criteria for MM by way of determination of calcium levels, renal function, full blood count and X-rays. Monoclonal gammopathy should also be demonstrated. Currently, there are no facilities for protein electrophoresis, immunoglobulin quantitation and immunofixation in Malawi.
- In terms of MM treatment, prednisone is commonly available in public pharmacies in Malawi and oral dexamethasone is occasionally available.
- Vincristine, doxorubicin (infusion) and dexamethasone (VAD regimen) can be given in Malawi.

11.0 Musculoskeletal Disorders

11.1 Arthritis (Non-Infective)

- Make specific diagnosis whenever possible and treat accordingly
- An acute mono-arthritis should always be considered to be infective until evidence to the contrary is obtained

11.1.1 Non-Specific Inflammatory Arthritis and Rheumatoid Arthritis

Signs and Symptoms

- Pain and swelling in joint
- Fever (especially in children)
- For rheumatoid arthritis key symptoms are:
 Chronic pain, joint swelling and deformity

Investigations

- FBC
- X-ray joint
- ESR

Treatment

Adults:

- 1. NSAIDS
 - Give Ibuprofen 1.2–1.8g daily in 3 divided doses after food

Alternatively:

- Give Aspirin 300-900 mg after food every 4 hours. Maximum of 4g daily in divided doses in acute conditions
- Review after 7 days
- If not responding: change to an alternative non-steroidal anti-inflammatory drug (NSAID):
 - Give Indomethacin 25 50mg tds after food for another 7 days or
 - Give Diclofenac Sodium 25-50mg tds after food or SR 75mg bd for another 7 days
 - Consider referral for specialist opinion
- 2. Intra-articular steroid injection with **Depo Medrol** or **Triamcinolone**
- Disease Modifying anti-rheumatic agents
 Methrotraxate (Needs specialist prescription)
 Children:
 - Always refer to hospital
 - Prior to referral, give Aspirin 20 mg/kg after food qid or Paracetamol 15mg/kg tds

11.2 Arthritis (Septic)

Symptoms and signs

- Fever
- Swollen, painful, warm joint
- Severe local tenderness
- Pain on joint motion
- Investigations

 Aspirate the joint for diagnostic and therapeutic purposes

Treatment

Adults:

- Give Flucloxacillin 1g IV every q6h for at least 14 days plus
- Give Ciprofloxacin 500mg bd
- A further 2-4 weeks of oral antibiotics (Flucloxacillin and Ciprofloxacin) may be required

Alternatively if penicillin allergic:

 Give Clindamycin 450mg qid or, if not available use Ceftriaxone 2g IV daily for 2 weeks followed by oral Erythromycin 500mg qid and Ciprofloxacin 500mg bd for a duration of 2-4 weeks

Children:

• Give Ceftriaxone 50mg/kg daily

Or alternatively when staphylococcal infection is suspected:

Give Flucloxacillin 25 mg/kg IV q6h for 14 days

Or if above drugs not available

 Give Chloramphenicol 12.5 mg/kg qid for at least 14 days, or 4 weeks if there is associated osteomyelitis, clinically evident by bone swelling or proven by X-rays after the initial 14 day course **Note:** Surgical drainage may be indicated TB septic arthritis is treated as for other forms of extra-pulmonary TB

Further management

 If pus present, always refer early for arthrotomy.

RED FLAG: Always refer early to hospital for systemic treatment and arthrotomy

11.3 Gout

11.3.1 Acute attack

Symptoms and signs

- Redness, tenderness and swelling around the joint
- Mimics septic arthritis

General measures

- Encourage Rest
- Ensure abundant fluid intake

Treatment

 Give Ibuprofen 800mg qid preferably after food in established cases until attack subsides

Alternatively:

- Give **Indomethacin** 50-75mg tds with food or
- Give Diclofenac Sodium 25-50mg tds preferably after food and preferably suppositories

- Give Colchicine 1.0mg followed by 0.5mg no more frequently than every 4 hours until pain is relieved or diarrhoea or vomiting starts.
 Maximum of 6mg per course; course should not to be repeated within 3 days
- Give **Prednisolone** 30 50mg od for 5 -7 days

11.3.2 Prevention of attacks

- Encourage physical exercise
- Encourage reduction in dietary protein (if intake is high)
- Avoid alcohol
- Only indicated in recurrent gout attacks.
- Allopurinol 100 mg daily after food
- Maintenance dose: Up to 200 -600mg od; dosage >300 mg to be given in divided doses; may be required life long
- Gradually increase over 1-3 weeks to 300 mg od, according to plasma or urinary uric acid concentration
- Do not start this treatment until an acute attack has completely subsided

11.4 Musculoskeletal Pain and Trauma

 Joint pain, lumbago, and chronic musculoskeletal disorders are common in adults

Adults (symptomatic treatment)

Give analgesics

If pain persistent or severe:

Refer to a Surgeon or a Physiotherapist

11.5 Osteomyelitis

11.5.1 Acute Osteomyelitis

Adults and Children:

- Refer to the hospital
- Admit to hospital for rest
- Splint the affected limb as required
- Give analgesics

If pain is severe:

- Give Pethidine 1mg/kg IM
- Repeat every 6 hours for a maximum 4 doses
- Drain pus surgically from the bone and send for culture and sensitivity testing
- Do not await culture results before starting antibiotic treatment

Children Over 2 years:

 Give Flucloxacillin 25 mg/kg up to a maximum of 500 mg qid, initially IV, then orally from 48 hours after fever has settled

Alternatively:

 Give Chloramphenicol 25 mg/kg up to a maximum of 500mg qid, initially IV, then orally from 48 hours after fever has settled

Children Under 2 years:

- Give Ceftriaxone 50mg/kg od or (Especially if staphylococcal infection is very likely):
- Give Flucloxacillin 25 mg/kg qid, initially IV, then orally from 48 hours after fever has settled

Alternatively, if the above not available:

 Give Chloramphenicol 25 mg/kg qid, initially IV, then orally from 48 hours after fever has settled

Note: Antibiotic treatment should be continued for 4 weeks under hospital supervision.

11.5.2 Chronic Osteomyelitis

Adults and Children:

Treatment

- Surgical treatment by sequestrectomy when an adequate involucrum has formed
- Antibiotic treatment for febrile flare-ups of infection as for acute osteomyelitis (See Section 11.5.1 above)
- Antibiotic cover for surgery as appropriate after culture and sensitivity testing
- Give Ibuprofen 1.2 1.8g od in 3 divided doses after food

Alternatively:

- Give Aspirin 10mg/kg tds, preferably after food, up to an adult maximum of 600 mg per dose
- For children, give Paracetamol instead of Aspirin

11.6 Rheumatic fever

Note: Always refer to hospital

11.6.1 Acute attack

Treatment

- Give Benzathine Penicillin 1.2MU IM single dose
 - Children <30 kg: Benzathine Penicillin 600,000 units

Alternatively, if compliance can be ensured:

- Phenoxymethylpenicillin 250 mg qid for 10 days
 - Children: Phenoxymethylpenicillin 12.5 mg/kg/dose

Alternatively, if penicillin allergy:

• Erythromycin 500mg tds for 10 days

11.6.2 Acute Carditis

• Strict bed rest until carditis has resolved

Adults and children:

Treatment

- Give Aspirin 25mg/kg, preferably after food, every 6 hours
- Reduce dose if tinnitus or other toxic symptoms develop
- Continue treatment with this until fever and joint inflammation are controlled

Then reduce dose gradually over a 2 week period

If symptoms recur:

Restart full dose

In severe carditis with heart failure and not responding to aspirin:

- Add **Prednisolone** 2 mg/kg od
- Reduce dose gradually after 3-4 weeks
- Treat heart failure

11.6.3 Chorea

Treatment

Adults and Children:

• Give Haloperidol 25 micrograms/kg tds

11.6.4 Prophylaxis of Rheumatic fever

11.6.4.1 Prevention of further attacks

• Continue treatment until at least age 25 Treatment

- Give Benzathine Penicillin 1.2 MU IM monthly
 - Children < 30 kg: 600,000 units/dose</p>

Alternative if compliance can be ensured:

• Give Phenoxymethylpenicillin 250mg bd

Alternative if penicillin allergy:

- Give Erythromycin 500 mg daily
 - Children <30 kg: 250mg</p>

11.6.4.2 Prophylaxis of Bacterial Endocarditis

 Needed to prevent bacterial endocarditis in those with previous rheumatic fever or any heart valve abnormalities of other cause.

11.6.4.3 Before Dental Extraction

Prophylaxis in Adults and Children > 30 kg:

 Give Amoxycillin 3g oral taken 1 hour before the dental procedure

Alternatively if penicillin allergy:

 Give Erythromycin 1.5g taken 1 hour before the procedure and 500 mg 6 hours later

Prophylaxis in Children > 30 kg:

 Give Amoxycillin 50 mg/kg taken 1 hour before dental procedure, and repeated 6 hours later

Alternatively if penicillin allergy:

 Give Clindamycin 600mg, orally, taken 1 hour before the procedure and 300mg 6 hours later

Prophylaxis in Children > 30kg:

- Give Clindamycin 600mg IM/IV 30min prior to the procedure followed by 300mg 6 hourly.
 (20mg/kg for children) OR
- Give Ceftriaxone 1g IV prior to the procedure or 50mg/kg

For high risk patients add:

Gentamicin 1.6mg/kg prior to the procedure OR Vancomycin 1g IV (20mg/kg/IV for children <10 years old) prior to the procedure.

11.6.4.4 Prophylaxis before other procedures

- a. For genito-urinary surgery or instrumentation:
 - Give Amoxycillin 1g IV or IM plus
 - Give Gentamycin 2mg/kg IV or IM 30 minutes before the procedure, then
 - Give **Amoxycillin** 500mg taken 6 hours later

Alternative if penicillin allergy:

- Give Erythromycin 500mg qid for 48 hours, instead of Amoxicillin
- b. For obstetric and gynecological procedures:
 - Not required except for those with prosthetic heart valves who should receive prophylaxis as for dental procedures

11.7 Cellulitis and Tropical Pyomyositis

- Treatment for cellulitis is both medical and surgical
- Medical treatment may prevent abscess formation at the start of infection, when the muscle is
- swollen, hot and painful
- Immobilize and give:

Treatment

Adult and children:

- Give Flucloxacillin 1g IV every 6 hours for at least 14 days plus,
- Give Ciprofloxacin 500mg orally twice daily

If penicillin allergic:

• Give **Ceftriaxone** 2g IV daily for 2 weeks **Note:** For Pyomyositis, perform surgical treatment (abscess drainage) when the swelling becomes fluctuant, alongside antibiotics

12.0 Sexual and Reproductive Health

12.1 Obstetric conditions

12.1.1 Antenatal Care

 Centred on ensuring, supporting and maintaining maternal and foetal well being throughout normal pregnancy and childbirth

Pillars of FANC:

- Early detection and treatment of problems and complications
- Prevention of complications and diseases
- Birth preparedness and complication readiness
- Health promotion

Frequency and timing of FANC visits:

- Appropriate scheduling depends on the gestational age of the pregnancy and also the woman's individual needs. For women whose pregnancies are progressing normally, four visits are sufficiently; one in first trimester, another one in second trimester and two in the third trimester
- Women with common discomforts, special needs, conditions that lie beyond the scope of basic care, or other problems may require additional visits

ANTENATAL CARE MATRIX

| Parameter | Weeks of Gestation | | | |
|--|-----------------------------------|---|----------------|-------------|
| | First visit or <16 weeks | | 28-32 weeks | 36 weeks |
| Registration | ü | | | |
| Comprehensive history-taking | ü | | | |
| History of complaints in current pregnancy | ü | ü | ü | ü |

Observations and clinical investigations

12.1.2 Hypertensive Disorders in Pregnancy

12.1.2.1 Pregnancy-Induced Hypertension (PIH)

At Health Centre

- Blood Pressure ≥ 140/90mmHg or an increase of Systolic BP of 20 and Diastolic BP of 15 from pre-pregnancy levels.
- Recheck Blood Pressure after resting for 10
 15 minutes

If still raised:

- Refer to the next higher level if it's severe PIH, Pre-eclampsia and eclampsia with accompaniment of a health worker.
- In case of eclampsia or eminent eclampsia; give a loading dose of 4g Magnesium
 Sulphate 20% solution in 500 ml of Normal
 Saline infused over 10 minutes plus 5 g of
 Magnesium Sulphate 50% solution in each buttock deep IM with 1 ml of 1% lignocaine
- Refer immediately

At the hospital

- Admit
- Re-assess the patient and exclude severe forms.

If it's Pregnancy Induced Hypertension (PIH)

 Give Methyldopa 500mg to 1g q8h until BP settles. If no improvement, add Nifedipine
 SR po to a maximum of 40 mg tds

12.1.2.2 Pre-eclampsia

Symptoms and signs:

 Raised BP ≥ 140/90mmHg, proteinuria ≥ + (dipstick), ± oedema, at the gestation ≥ 20 weeks

Treatment

If signs of pre-eclampsia

- Admit, evaluate and take full history
- Check for signs of imminent eclampsia, check weight, urine dip stick daily, and BP every 4 hours
- Evaluate wellbeing and gestation. Do ultrasound

If diastolic BP >90mm Hg and <110mm Hg then give

Give Methyldopa 500mg to 1g tds and review daily

If diastolic >110mm Hg

- Do not lower BP abruptly and avoid use of sublingual nifedipine
- Give **Hydralazine** 5mg IV slowly over 5 min.
- Repeat every 20 minutes until diastolic pressure is below 110 mm Hg.

If diastolic BP is still >110

 Give Hydralazine 40mg IV in 1 litre of Ringer's Lactate over 8 hours to maintain BP at below 110 mmHg.

Alternatively

Give Nifedipine 10mg. Recheck BP in 20 minutes. Repeat Nifedipine if diastolic ≥110mmHg, manage as severe pre-eclampsia.

- Watch carefully for eclampsia: If this develops see section for eclampsia
- Consider delivery regardless of gestational age if:
 - BP is difficult to control
 - Urine output is decreasing
 - Critical signs/symptoms persist (suggesting severe preeclampsia) eclampsia

12.1.2.3 Severe Pre-Eclampsia

Symptoms/Signs

 Diastolic BP >110, marked oedema, proteinuria ++/+++, headache, blurred vision, epigastric pain, oliguria, hyperreflexia

Management

- Admit in labour ward
- Put up an IV line normal saline
- Give 4g of 20% of Magnesium Sulphate (MgSO₄.7H₂O) solution IV over 5 minute period (20 mls)
- Administer 5g of 50% Magnesium
 Sulphate (MgSO₄.7H₂O) (20 mls) with 1ml of 1%
- Lignocaine IM deep in each buttock (total 10 g)
- Catheterize
- Monitor BP every 15 minutes until BP is lowered, then hourly

12. Sexual and Reproductive Health

- In the event of a convulsion after 15 minutes administer 2g of 50%
 Magnesium Sulphate solution IV over 5minutes (4mls)
- Monitor foetal heart every 30 minutes
- Refer to hospital labour ward and the midwife to escort the woman

Magnesium Sulphate Toxicity

Toxicity

- 1) Patella Reflexes are abscent
- 2) Respiratory rate < 16 per minute
- 3) Urine output < 30ml/hr since last admission of Magnesium Sulphate 4 hrs ago

Before Giving The Next Dose, Ensure That

- 1) Patella Reflexes are present
- 2) Respiratory Rate is ≥ 16/min
- 3) Urine output is ≥ 30 ml/hr since the last admission of Magnesium Sulphate 4 hrs ago

In Case of Toxicity

- 1) Don't give the next dose of Magnesium Sulphate
- 2) Assist ventilation if respiratory arrest (mask & bag; if severe respiratory depression, endotracheal intubation & mechanical ventilation)
- 3) Calcium gluconate 1g IV slowly until toxic effexts are reversed

Magnesium Sulphate & Nifedipine

If both are administered simultaneously, BP may drop precipitously because both are Calcium antagonists. The risk is there though rare.

NB: The **Magnesium Sulphate** used in Pre-eclampsia and Eclampsia has the formula

MgSO₄.7H₂O

12.1.2.3.1 Foetal Well-being or Maturity

Check

- Full blood count, urea & creatinine in serum
- Liver function tests
- Electrolytes: sodium, potassium, and chloride
 - Refer to central hospital if lab results are derranged

If >34 weeks gestation:

- Stabilize patient
- Deliver within 24 hours either by induction, if cervix is favourable, or by caesarean section

If <34 weeks gestation:

- Inform clinician
- Assess fetal well-being using Ultrasound scan or Cardiotocograph
- Give **Dexamethasone** 6mg every twelve hours
 IM for 4 doses

Alternatively

 Betamethasone 12mg IM once daily for a total of 2 doses

12.1.2.4 Eclampsia

Signs and symptoms

- Convulsions
- Diastolic BP ≥ 90 mm Hg after 20 weeks gestation
- Proteinuria ≥ 2+

Note: Convulsions can occur prior to labour, intrapartum or postpartum.

If a pregnant woman convulses start treatment for eclampsia, but rule out other organic causes, including Epilepsy, Meningitis, Cerebral malaria, Encephilitis, Hypoglycemia

General management

- Place the woman on her side to reduce risk of aspiration
- Secure airway, aspirate secretions or vomitus
- Give adequate oxygen supply by nasal prongs or face mask
- Protect the woman from injury
- Put up an IV line normal saline
- Control convulsions with magnesium sulphate (see dose below)
- Give 4g of 20% of Magnesium Sulphate IV over 5 minute period (20 mls)
- Administer 5g of 50% Magnesium Sulphate
 (20mls) with 1ml of 2% Lignocaine IM deep in each buttock (total 10 g)
- Catheterize
- Monitor BP every 15 minutes until BP is lowered, then hourly
- In the event of a convulsion after 15 minutes administer 2g of 50% Magnesium Sulphate IV over 5 minutes (4mls)
- Monitor foetal heart half hourly

Treatment

- Closely monitor for signs of magnesium sulphate toxicity such as presence of patella reflexes, the respiratory rate (not less than 16), and urinary output should not be less than 25mls an hour.
- Continue magnesium sulphate for 24 hours post- delivery or 24 hours after the last convulsion whichever was the last
- Maintenance dose: Magnesium sulphate
 5g of 50% solution every 4 hours deep IM till
 24 hours post-delivery or 24 hours after the
 last convulsion which ever was the last.
 Addition of 1.0ml of 2% Lidocaine minimizes
 discomfort

Note: Once Magnesium Sulphate is administered a decision must be made to deliver the pregnant woman within 12 hours

If magnesium sulphate is not available give:

- Loading dose of **Diazepam** 10mg IV slowly over 2 minutes
- Maintenance dose of **Diazepam** 40mg in 500mls of normal saline or **Ringer's Lactate**
- Do not give more than 100mg in 24 hours
- If not already at hospital, refer the patient to hospital

RED FLAG: In case of Magnesium Sulphate toxicity administer Calcium Gluconate 1gm as IV stat dose and stop Magnesium Sulphate

Mode of delivery

- Carry out an obstetric assessment to decide on appropriate mode
- Only allow assisted vaginal delivery if labour is progressing quickly
- Consider caesarean section if unlikely to deliver in 6-12 hours regardless of gestational age
- Give Oxytocin 10 IU (1mL amp) by IV push in the 3rd stage
- Do not use ergometrine

Monitoring

 Continue careful observation (and treatment if necessary) for at least 48 hours after delivery

12.1.3 Prelabour/Premature Rupture of Membranes

 Rupture of the membranes before labour (prelabour is >37 weeks gestation while PPROM is before 37weeks)

Signs and Symptoms

- Watery vaginal discharge
- Management If gestation less than 34 weeks
 - No digital vaginal examination should be done
 - If the diagnosis is in doubt, perform sterile speculum examination as well rule out cord prolapse
 - Check vital signs 4 times hourly, assess fetal heart rate

- Provide a pad and observe for colour amount and smell of liquor daily
- If at a Health Centre,
 - Give Erythromycin 250mg tds for 7 days
 - Give Metronidazole 400mg tds for 7 days
 - Give corticosterioids if <34
 weeks. Refer immediately
- If at the Hospital
 - If still draining, deliver at 34 weeks if there are no other contraindications
 - If signs of intra-uterine infection (temperature 37.5 °C or more, purulent or offensive liquor), or fetal distress. Inform the most senior person available, and plan urgent delivery regardless of gestational age
- If gestation 34 weeks or greater
 - Do sterile speculum exam to confirm draining and rule out cord prolapse
 - If membranes have been ruptured for more than 12 hours, give antibiotics:
 - Give Ampicillin 2g IV every 6 hours
 - If no signs of infection evident; give Erythromycin 500mgs qid

Alternatively

- Give Benzylpenicillin 2 MU IV every 6 hours until delivery
- Stop 48 hours after delivery unless there are signs of sepsis
 - ➤ If labour does not begin spontaneously within 24 hours, assess the cervix and induce labour, if no contraindication to vaginal delivery. If unsuccessful deliver by caesarean section
 - If signs of intra-uterine infection develop, inform the most senior person available, to decide on mode of delivery

12.1.4 Chorioamnionitis

Intra-uterine infection

Signs and Symptoms

- Foul-smelling vaginal discharge after 28 weeks of pregnancy,
- Fever/chills, abdominal pain, fetal tachycardia

Treatment

- Give Benzylpenicillin 2.5 MU IV stat and/or
- Give Ampicillin 2gms IV stat
- Refer to hospital

At hospital

- Give Metronidazole 500mg IV q8h, and
- Ampicilline 1g qid/Benzylpenicillin 2 MU IV q6h and
- Give Gentamycin 240 mg IM single dose daily

- Continue for 48 hrs after the fever subsides, but not less than 5 days.
- Deliver urgently. Induce or accelerate labour with Oxytocin if the cervix is favourable and no contraindications; do ceasarean section if necessary.
- If mother has amnionitis or if membranes were ruptured for more than 12 hours before delivery, start newborn on:
- Give Benzylpenicillin 50,000 IU/kg/dose IM q12h
- Give Gentamycin 5 mg/kg IM od for 5 days if birth weight >1500 g).

12.1.5 Anteparturm Haemorrhage

Vaginal bleeding from 28 weeks gestation to before delivery

 Consider: abruptio placenta, placenta previa, Ruptured uterus, cervicitis

Management

- Explain to the patient
- Avoid digital vaginal examination
 - Check vital signs
 - Insert large bore canulae, with Ringer's Lactate or normal saline
 - ➤ If at a Health Centre refer
- At the hospital
 - Urgent FBC, group and cross match
 - Start plasma expanders or transfusion if haemodynamically compromised. FFP if DIC suspected
 - Ultrasound examination (if feasible)

Plan for urgent delivery and/or laparotomy regardless of Gestational age

12.1.6 Medical Conditions in Pregnancy

12.1.6.1 Malaria in Pregnancy

- Similar presentation as in non-pregnant woman, however prone to complications:
 - Hypoglycaemia and preterm delivery

12.1.6.2 Uncomplicated Malaria

- In 1st trimester of pregnancy give oral **Quinine** 10mg/kg body weight, administered q8h for 7 days.
- In 2nd and 3rd trimester give **Lumefantrine**Artemether (LA) bd for 3 days.

Note: Pregnant women are susceptible to hypoglycemia when taking quinine.

12.1.6.3 Severe malaria in pregnancy

Treatment

- Same as adults
- Avoid usage of Artesunate in 1st trimester.
- Refer to Severe Malaria section

12.1.6.4 HIV/AIDs in Pregnancy

- Ascertain HIV status in all pregnant women
- Offer HTC if unknown status
- Manage as per current PMTCT program guidelines

12.1.6.5 Anaemia in Pregnancy

 Anaemia in pregnancy is defined as HB of less than 11g/dl, (severe anemia is HB less than 7g/dl at any gestational age).

Signs and symptoms

 Fatigue, headache, weakness, heart palpitations and dizziness

Physical Examination

Pallor of the skin and mucous membrane.

At Health Centre

• If HB is < 8 g/dl or any complications, refer

Prevention

- Provide all antenatal women with FeFol 325mg po bd
- Advise on diet rich in green leafy vegetable, liver, fish, eggs
- To prevent hook worm give Albendazole (400 mg stat)
- To prevent malaria give 2 doses SP (three tablets each dose) 4 weeks apart, starting after quickening (16 weeks gestation)
- Advise to keep adequate interval between pregnancy > 2 years minimum
- All breastfeeding mothers should take iron supplements

At District Hospital

Management

Prevention (as above in Health Centre)

Treatment

- FBC and treat according to the result
 - If HB is < 7g/dl especially if symptomatic, then blood transfusion
 - Transfuse rapidly if anemia due to blood loss.
 - Transfuse slowly and with diuretics if chronic anaemia. (To reduce risk of congestive cardiac failure due to sudden circulatory overload.)
 - Treat with ferrous sulphate/ folic po bd and recheck HB in 2 to 4 weeks.
- Urine and stool examination and treat according to results
- Treat malaria or schistosomiasis if indicated
- If Haemoglobinopathy (e.g. sickle cell anemia) is suspected, then refer

12.1.6.6 Diabetes in Pregnancy

 It is a group of metabolic disease characterized by hypersensitivity from defects in insulin secretion, action or both. Can be gestational or preexisting.

Signs and symptoms

History: Polyuria, polyphaghia, polydipsia.
 Suspect/screen in macrosomia, unexplained IUD, FHx of DM, Maternal obesity, excessive weight gain

At Health centre

Refer to the next level

At District hospital

Management

 Goal is to maintain FBS at 6 – 8MMOI/I for gestational

Investigations

- Routine Investigations (FANC), others RBS, Hb A₁C
- Ophthalmic examination

Note: (It is advisable to manage only uncomplicated diabetes mellitus)

12.1.6.7 Prepregnancy Care

- Pregnancy counselling: diet, ideal weight and sugar levels
- Switch to insulin if unstable on oral drugs

12.1.6.8 Antenatally

- Continue pre- pregnancy regimen if blood sugar is controlled.
- Consider in patient admission for DM5 education and glucose control.
- Antenatal care every 2 weeks until 30 wks. gestation, then weekly until delivery
- Screen all pregnant women at 24 30 weeks
- Patient log book to self-record daily insulin dosages and daily blood glucose levels before each meal (3 meals) and 2200 hours (before going to bed)Check FBS

- If <6mmol/l (108 mg/dl), patient is managed by diet alone.
- ➤ If > 6 mmol/l Insulin must be started.
 - If first trimester, total daily dose = weight x 0.7 units
 - If second trimester, total daily dose = weight x 0.8 units
 - If third trimester, total daily dose =
 weight x 0.9 1.0 units
 - Given as 2/3 of total daily dose in the morning at breakfast: 1/3
 Soluble Insulin and 1/3 as Long Acting Insulin
 - Given as 1/3 of total daily dose in the evening at dinner (17 hrs): ½
 as Long Acting Insulin
 - For example, for weight of 72 kgs in third trimester, give 16 units
 Soluble Insulin and 32 units Long
 Acting Insulin at breakfast and 12 units
 Long Acting Insulin at supper.
- Ultrasound every 4 weeks

12.1.6.9 Intrapartum Management

- Hospitalization at 34 36 weeks
- Elective delivery at 38 -39 weeks
 - No specific treatment if labor progresses normally and quickly
 - For induction, or prolonged labor: add 1/3 of her daily insulin as soluble to 1 L of Dextrose Normal Saline (DNS) and treat 40dpm

- For caesarean: skip AM Insulin, start DNS
- Place Oxytocin in separate bag of Normal
 Saline (NS) fluid using separate IV access
- At 39 weeks gestation for women with well controlled blood sugar and no vascular disease.
- At earlier gestation for class D and higher, polyhydromnoios, macrosomia, poor blood glucose control, Chronic Hypertension on medication or IUGR and IUD
- Caesarian delivery for EFW > 4500 on UD.

12.1.6.10 Postnatal period

- Breastfeeding infant early and notify pediatric clinician of maternal diabetes
- Use insulin sliding scale for 5 days post vaginal delivery and then resume pre pregnancy regimen
- Treat with DNS at 3L daily post C/S until tolerating PO and then use insulin sliding scale
- Advise mothers to start diabetic diet as soon as possible

Note: COMPLICATED DIABETES (Refer to central hospital)

12.1.7 Dysfunctional Labour Syndrome

 Labour not progressing according to expectation; has passed alert line

Management

- Explain to the mother
- Ensure adequate pain relief and hydration

- Check vital signs
- Analyze partogram
- Consult senior or refer as soon as possible
- Consider 3 P's:
 - Passenger: assess size of baby, presentation, position, fetal distress
 - Power: if contractions are inadequate, then augment
 - Passage: assess pelvis adequacy and consider C/S
 - ✓ If obstructed labour (caput ++, moulding>++, Cervix poorly applied, Bandls ring, maternal and fetal distress)
 - Secure IV access, Group and Xmatch, secure blood
 - Start IV antibiotics: GiveAmpicillin 1g IV Stat
 - Monitor vital signs
 - Catheterize
 - Prepare for C/S stat.
 - ✓ If Ruptured uterus suspected: Do laparotomy

12.1.8 Caesarian Section

 This operation is the process whereby the child is removed from the uterus by direct incision through the abdominal wall and the uterus following indications that arise from the mother, the fetus, or both.

At Healthe Centre

• Refer to the District Hospital

At District Hospital

- Elective indications
 - Prophylaxis: Give Ceftriaxone IV 2g stat OR Ampicillin 2g stat (30 min- 1hr before operation)
- Emergency Indications
 - ➤ All cases should be done within 2hrs

Management

- Do FBC
- Antibiotherapy:
 - 1st line, give Ampicillin 1g q6h for 7 days + Gentamycin 160mg q24h for 5 days (if no contraindications)
 - 2nd line, give Metronidazole IV 500mg q8h for 5 days or Metronidazole tablets 400mg tds for 7 days
 - > 3rd Line, **Ceftriaxone** 1g q12h for 7-10 days
- Sitz bath twice daily if there is episiotomy or tears- Care of incision wound
- Correct anaemia if it is there
- High protein food in patients with malnutrition
- Proper information, education and communication (IEC) concerning hygiene for the baby and mother

12.1.9 Intrauterine Death (IUD)

 Death of the fetus > 24 weeks gestation or > 500gms weight

Clinical signs

 Patient complains of absence of fetal movement. If IUD is few days old breast size may diminish and colostrum secretion may start in some cases.

At Health Centre

Refer to the District Hospital

At District Hospital

- Diagnosis: Decreased or absent fetal movement
- Exam: No fetal heart heard on fetoscope or doptones, fundal height may be less than expected.
- Investigations: US with no fetal cardiac (Verify by 2 health care providers); check FBC, RBS, grouping VDRL

Management

- Induction of labour (refer to Reproductive Health protocols)
 - If augmentation of labour, then manage similar to live birth
 - Ensure privacy to the extent possible
 - Provide bereavement counselling
 - Placental evaluation and perinatal autopsy recommended
 - ARM is avoided in early labour as it can lead to ascending infection.
 - Prophylaxis: Give Ceftriaxone IV 2g stat.
 - Antibiotherapy: if signs of infection or macerated (Monotherapy or triple therapy)

- ✓ 1st line: give Ampicillin 1g q6h for 7 day
 + Gentamycin 160mg q24h for 5 days
 (if no contraindications)
- ✓ 2nd line: give Metronidazole IV 500mg q8h for 5 days or Metronidazole tablets 400mg tds for 7 days
- √ 3rd line: give Ceftriaxone 2g q24h for 7-10 days
- Triple therapy: give Ampicillin IV 1g q6h 7 days + Gentamycin IV 160mg q24h for 5 days + Metronidazole IV 500mg q8h for 7 -10 days
- Caesarian section: performed if spontaneous delivery failed, if there is contraindication of SVD or contraindication of induction.
- Craniotomy should be performed at central hospital level only with Craniotomy set.
- Decapitation of the baby should not be performed.

12.1.10 Neonatal resuscitation

 Refer to WHO neonatal resuscitation flow chart second edition

12.1.11 Cord care

- Wait for 1 to 3 minutes or for cessation of the cord pulsation, then clamp
- Tie the cord with two sterile ties; one 2 cms from the baby's body and the next 3cms from the first tie
- Apply 7.1% Chlorhexidine Digluconate to the cord stump soon after cutting to prevent infection

- Continue monitoring the stump for bleeding, especially in the first 24 hours
- Bleeding later on from the cord might indicate haemorrhagic disease of the newborn (due to Vitamin K deficiency) or an infection. Refer urgently.

12.1.12 Postpartum Haemorrhage (PPH)

 Blood loss from the genital tract of more than 500ml after delivery of a baby.

Causes

 Uterine atony, retained products of conception, genital tract trauma, coagulation problems, ruptured uterus, endomyometritis

12.1.12.1 Primary PPH

- Abnormal vaginal bleeding within 24 hours of delivery
- Resuscitate
- Set up an IV line and empty bladder
- Replace blood loss with IV fluids / blood
- Identify and treat the cause
- If uterine atony:
 - Rub up a contraction
 - Give 10 units Oxytocin IV stat then 40 units infusion
 - ➤ Give Misoprostol 1000mcg rectally
 - Refer to hospital with nurse accompaniment

- If retained placenta:
 - Attempt manual removal or evacuation in theatre
 - Inspect the vagina thoroughly for perineal injuries and repair
 - ✓ If above interventions fail:
 - Take patient to theatre for exploratory laparotomy; B-lynch suture or hysterectomy

12.1.12.2 Secondary PPH

Abnormal bleeding 24 hours or more after delivery

Causes

Retained products, often with infection

Treatment

- Set up an IV line and resuscitate
- Replace blood loss with IV fluids/blood
- Empty bladder
- Rub up a contraction
- Give Oxytocin 10 units IM
- Give Amoxycillin 500 mg tds plus
 Metronidazole 400 mg tds
- Alternatively if penicillin sensitive: Give
 Erythromycin 500 mg qid
- Refer immediately for evacuation of the uterus

Supportive measures

If patient is septic start antibiotics; Ampicillin
1g qid/ Benzyl penicillin 2 MU IV q6h,
Gentamycin 240mg od, Metronidazole 500mg
tid

Give IV fluids to sustain a high degree of perfusion

12.1.13 Post-Natal Care

 For care of a woman from immediately after birth to 6 weeks, refer to Reproductive Health protocols

12.2 GYNAECOLOGY

12.2.1 Abnormal Vaginal Bleeding

 Bleeding which deviates from normal menstrual pattern: interval, duration amount.

12.2.1.1 Acute Vaginal Bleeding

- Resuscitate
- Exclude pregnancy
- To reduce/stop bleeding
 - Give Ibuprofen 400mg q8h for 5 days
 - Give Traxenamic Acid 1g q8h hourly po for up to 5 days
 - Combined oral contraceptive 2 tabs od x10days od then 2-6 cycles unless contraindicated
 - Give Ferous Sulphate 20mg od. Refer if persistent
- Young adolescents, usually physiological, pathology rare
 - Exclude pregnancy and its complications and manage conservatively
 - If bleeding is heavy manage as above

- Women of child bearing age:
 - Strongly suspect complications of pregnancy including ectopic pregnancy. Other causes: use of hormonal contraceptives, DUB, ICD, fibroids, choriocarcinoma, cervical cancer
 - Speculum examination mandatory to rule out local causes

12.2.2 Dysfunctional Uterine Bleeding

Usually no organic cause found

Management

- Give Cyclic Progesterone 5-10mg bdx 14 days on second part of the cycle; alternatively, oral COC 1 od 3-6 cycles
- If no improvement refer to specialist

Post-menopausal women

- Important causes are endometrial cancer, cervical cancers, cervical polyps, atrophic vaginitis
- Always need investigation therefore refer early

At the hospital

- Speculum examination mandatory
- USS for masses and endometrial thickness. If >4mm, do diagnostic D&C

12.2.3 Abortion and its complications

Loss of pregnancy before viability 28 weeks or

- Fetal weight 500g
- Post abortal Sepsis

Complications

Post abortal haemorrhage

- Resuscitate and stabilize the patient
- Carry out vaginal examination
- Remove products of conception and/or foreign bodies
- Give Oxytocin 10 units IM
- Alternatively, give Misoprostol 600mcg po or 400mcg sublingually
- Perform (or if not possible refer)
 evacuation or manual vacuum aspiration
 (MVA) if gestation < 12 weeks
- If septic abortion suspected stat doses of antibiotics (refer to section on septic abortion)

Key features

Persistent fevers for >24hrs, foul smelling lochia, persistent bleeding

Management

- Maintain hydration: set up an IV line and give IV fluids
- Give Paracetamol 1 g stat
- Give Oxytocin 10 units IM to contract uterus
- Refer to hospital for evacuation of the uterus

At hospital

- ➤ Give **Misoprostol** 600mcg PR stat
- Give antibiotic treatment for 7 days as follows:

For sepsis

➢ If patient is able to take po drugs; give Doxycycline 100mg bd x 7 days and Metronidazole 400mg tds x 7 days

If patient is septic use IV drugs as follows:

- Give Metronidazole 500 mg tds
- ➤ Give **Gentamycin** 320mg IM stat
- Give Benzyl Penicillin 2 MU IV q6h
- Consider uterine careful evacuation in some cases

If not improving:

- Reassess and consider the appropriate intervention
- Change of antibiotics

• Comprehensive post arbortal management plan

- Counseling for contraception, HTC, future fertility plans/pregnancy care
- Cervical cancer screening
- STI/HIV prevention and treatment

12.2.4 Ectopic Pregnancy

- Often comes ruptured
- Cardinal signs and symptoms include: bleeding following amenorrhoea, PT positive, abdominal distension/tenderness, signs of haemorrhagic shock (low BP, high PR)
- Diagnosis is primarily clinical, Couldoscentis may support diagnosis
- Refer immediately to a hospital

At Hospital

- FBC, group and cross match
- Laparotomy stat. Do not wait for resuscitation
- Cover for chlamydia with **Doxycycline** 100mg bd x7 days

12.2.5 Vaginal Candidiasis (Moniliasis)

 Common infection occurs more frequently in patients taking antibiotics, pregnant women, HIV/AIDS patients and patients with diabetes.

Treatment

Adults:

 Give Clotrimazole 500mg intravaginally as single dose or Clotrimazole 200mg intravaginally once daily for 3 days

Alternatively

- Give Miconazole 200mg intravaginally once daily for 3 days or
- Give Fluconazole 150mg orally as single dose (contra-indicated in pregnancy)

OR

 Give Nystatin Pessary 100,000 IU vaginally every 12 hours for 7 days

To avoid re-infecting her partner, the male partner should:

- Apply the same vaginal cream as the patient
- Recurrent vulvalvaginal candidiasis should be referred to hospital

12.2.6 Dysmenorrhoea

 Rule out organic causes like PID, endometriosis, fibroids

Management

- Non pharmacological management (e.g. hot water bottle)
- Give **Mefenamic Acid** 500mg every 8 hours during menses for not more than 7 days

Alternatively

• Give **Ibuprofen** 400mg tds (see section on analgesics)

If no response:

Give cyclical courses of Low Estrogen
 Combined Oral Contraceptive tablets once daily for 3 – 6 months

If there is still no improvement

Refer to hospital

12.2.7 Mastitis

General Measures

- Apply hot compression and a constrictive bandage or bra to support the breast and relieve pain
- Maintain lactation in the infected breast if there are no nipple fissures to prevent stasis
- In severe cases, avoid engorgement by reducing milk production (e.g. Bromocriptine 2.5mg bid x 7-14 days)

Treatment

- Give Flucloxacillin 500 mg every 6 hours for 7 days
- Doses should be taken at least 30 minutes before meals

Alternatively

- Give **Amoxycillin** 500mg tds for 5-7 days
- If penecilline sensitive give Erythromycin
 500mg tds for 5 7 days
- Give Aspirin 600 mg tds after food

12.2.8 Breast abscess

- If breast abscess forms, drain surgically
- Change dressing every day
- Give antibiotic treatment as above

12.2.9 Obstetrical Fistula

 Obstructed labour is the most common cause of urogenital fistula in Malawi. Other aetiologies include surgery, radiation therapy and trauma or instrumental vaginal delivery.

Signs and symptoms

Draining urine or stools through the vagina

Prevention

- Encourage food rich in nutrition to girls before they get pregnant.
- Discourage early pregnancies.
- Encourage all pregnant women to attend antenatal care.
- Encourage pelvimetry during antenatal to all pregnant women
- Screen the high parturient (woman in labour).
- Encourage use and interpretation of partograph to all parturients.
- Encourage all parturients to deliver with a skilled birth attendant
- Ceasarian section must be done by skilled personnel

At Health Centre

 Refer all patients with draining urine and/or stools to the District Hospital

At District Hospital

- Reassess the patient to confirm the diagnosis with physical examination and dye test
- Repair minor and uncomplicated fistulas (a small hole from the bladder to the vagina).
 These must be repaired within 2 to 72 hours post delivery
- Refer all complicated fistulae to the Central Hospital, such as: VVF of more than 2cm, rectal vaginal fistulae, ureteral vaginal and urethral vaginal fistulae

12.3 Contraceptives

Types

 Combined hormonal contraceptives, progestogen only contraceptive, contraceptive devices, emergency contraceptive, barrier methods.

12.3.1 Combined Oral Contraceptive (COC)

- Most effective preparations for general use.
- Contain oestrogen and progestogen. Lowest effective dose preparations preferable.

Indications

- Contraception
- Menstrual disturbances

12.3.2 Oral Progestogen Only Contraceptive Pill

- Have a higher failure rate than COCs.
- Suitable alternative when Eostrogens are contraindicated; hypertensives, migraine, valvular heart disease and diabetes mellitus.
- Menstrual irregularities

 (oligomenorrhoea, menorrhagia)
 are more common but tend to
 resolve on long term treatment.

Dose

 1 tablet daily at the same time each day starting on day 1 of the menstrual cycle

12.3.3 Parenteral Progestogens

- Suitable for long term family planning option
- Injectable preparations (e.g. Depo-Provera®)
- Implant preparations (e.g Jadelle® and Norplant®, Implanon®)

12.3.4 Intrauterine Devices

- Offers long-term hormonal and nonhormonal contraception
- Not appropriate for those at high risk for STI's/PID
- Intrauterine infection (endometritis/PID) common in risky clients

Indications:

- Contraception for post-delivery/arbortal
- Emergency contraception after 120 hours
 - Example: Copper T 380®

12.3.5 Emergency Contraception

- Effective if taken within 120 hrs of unprotected intercourse.
- Give COC 4 tabs or Levonorgestrel 0.75mg every 12 hours for 24 hrs.
- If vomiting occurs within 3hrs of taking hormonal tablet, give replacement dose and anti-emetics can be considered.
- Explain the following:
 - > The next period may be early or late
 - A barrier method needs to be used until the next period.
 - Patient should return promptly if lower abdominal pains develop to rule out ectopic pregnancy or of any problems.
- An IUCD can be inserted up to 5 days of unprotected sexual intercourse.

12.4 Sexual Assault

- Treat as emergency with empathy, make sure your client feels safe
- Collect comprehensive general and gynae history, including the circumstances of the incident
- Examine the patient thoroughly (refer to One Stop Centre Guidelines)

- Fill the forms and inform the police and any relevant authorities
- Do tests: HTC, VDRL, pregnancy test
- Prophylactic as follows:
 - Emergency depending on age and time since incident (refer above)
 - PEP as per guideline; if HIV positive refer to HIV care clinic
 - > STI prophylaxis:
 - ✓ Give **Gentamicin** 240mg or 6 mg/kg single dose IM stat
 - ✓ Give **Metronidazole** 2g stat or 5mg/kg tds for 7 days
 - ✓ Give Benzathine Penicillin 2.4 MU IM (adult) stat or if < 25 kg: 600,000 IU stat, > 25-35 kg 1,200,000 IU stat
 - ✓ Give Doxycycline 100mg bdx 7 days if breast feeding, pregnant.
 - ✓ If < 8 yrs, give:</p>
 Erythromycin 12.5 mg/kg
 qid for 7 days
 - ✓ If > 8 yrs, give:Erythromycin 25 mg/kg qid for 7 days
- Treat other injuries as appropriate
- Offer counselling and protective custody where relevant

12.5 Neonatal Problems

12.5.1 Routine care of the newborn at delivery

- Most babies require only simple supportive care at and after delivery.
 - > Dry the baby with a clean towel.
 - Observe baby while drying.
 - Give the baby to the mother as soon as possible, place on chest/abdomen.
 - Cover the baby to prevent heat loss.
 - Encourage initiation of breastfeeding within the first hour.
 - Skin-to-skin contact and early breastfeeding are the best ways to keep a baby warm and prevent hypoglycaemia.

12.5.2 Neonatal Resuscitation

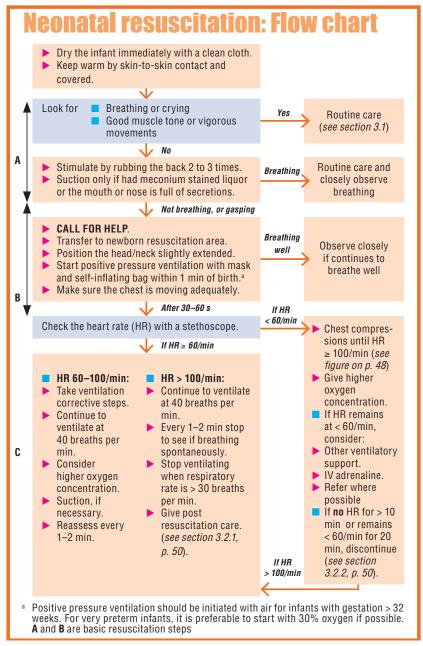
- For some babies the need for resuscitation may be anticipated
 - Those born to mothers with chronic illness
 - Where the mother had a previous fetal or neonatal death
 - A mother with pre-eclampsia, in multiple pregnancies
 - In preterm delivery
 - In abnormal presentation of the fetus

- With a prolapsed cord, or
- Where there is prolonged labour or rupture of membranes, or meconium-stained liquor.

Therefore,

- Be prepared for resuscitation at every delivery,
- Follow the assessment steps

Cessation of resuscitation



If after 20 minutes of resuscitation the baby is:

- Not breathing and pulse is absent: cease efforts.
- Explain to the mother that the baby has died, and give it to her to hold if she wishes.

12.5.3 Routine care for all newborn babies soon after delivery

- Keep dry in a warm room away from drafts, well covered
- Keep the baby with the mother, rooming in
- Initiate breastfeeding within the first hour
- Let the baby breastfeed on demand if able to suck
- Give Vitamin K (phytomenadione), according to national guidelines
- 1 ampoule (1 mg/0.5ml or 1 mg/ml) IM once
- (Do NOT use 10 mg/ml ampoule)
- Keep umbilical cord clean and dry
- Apply antiseptic ointment or antibiotic eye drops/ointment (e.g. Tetracycline eye ointment) to both eyes once, according to national guidelines
- Give oral polio, hepatitis B and BCG vaccines, depending on national
- Guidelines

12.5.4 Prevention of Neonatal Infections

- Many early neonatal infections can be prevented by:
 - Good basic hygiene and cleanliness during delivery of the baby
 - Special attention to cord care: use of spirit no longer recommended
 - > Eye care
- Many late neonatal infections are acquired in hospitals. These can be prevented by:
 - Exclusive breastfeeding
 - Strict procedures for hand washing for all staff and for families before and after handling babies
 - Not using water for humidification in incubators (where Pseudomonas will easily colonize) or by avoiding incubators (using kangaroo mother care instead).

12.5.5 Danger signs in newborns and young infants

- Neonates and young infants often present with non-specific symptoms and signs which indicate severe illness.
- These signs might be present at or after delivery, or in a newborn presenting to hospital, or develop during hospital admission.

 Initial management of the neonate presenting with these signs is aimed at stabilizing the child and preventing deterioration.

Danger Signs

- Unable to breastfeed
- Convulsions
- Drowsy or unconscious
- Respiratory rate less than 20/min or apnoea (cessation of breathing for >15 secs)
- Respiratory rate greater than 60/min
- Perinatal Asphyxia
 - Grunting
 - Severe chest indrawing
 - Central cyanosis

Management

- Give oxygen by nasal prongs or nasal catheter if the young infant is cyanosed or in severe respiratory distress.
- Start CPAP if there is no improvement
- Give bag and mask ventilation, with oxygen (or room air if oxygen is not available) if respiratory rate too slow (<20).
- Give Ampicillin (or Benzylpenicillin) and Gentamicin.
- If drowsy, unconscious or convulsing, check blood glucose.
- If glucose <2.2 mmol/l (<40 mg/100 ml), give Glucose IV.
- If you cannot check blood glucose quickly, assume hypoglycaemia and give Glucose IV.

- If you cannot insert an IV drip, give expressed breast milk or glucose through a nasogastric tube.
- Give phenobarbital if convulsing.
- Admit, or refer urgently if treatment is not available at your hospital
- Give vitamin K (if not given before).
- Monitor the baby frequently.

12.5.6 Care of Low Birth Weight (<2.5kg) and Premature Babies

 Low birth weight and premature babies are at risk of a number of problems – it is important to try and anticipate these and treat if necessary.

12.5.7 Hypoglyccaemia and Feeding

- Small babies should be fed every 3 hours.
- The daily feed volume should be calculated as below, and divided by 8 to give the 3 hourly feed volume, or divided by 12 to give 2 hourly feeds

| Day | 1 | 2 | 3 | 4 | 5 onwards |
|----------------|-----------------|----------------|----------------------|----------------------|------------------|
| Amount /day | 60 ml/ kg | 80- 90ml/kg | 110- 120ml /kg | 140- 150ml /kg | 150- 200ml/kg |

- If the baby shows any signs of hypoglycaemia (lethargy, floppiness, convulsions, apnoea, abnormal neurological behavior) please check the blood glucose level.
- If it is not possible to check then go ahead and treat:

Treatment

- If Blood Glucose < 2.2mmol/l
 - ➤ 1ml/kg 50% **Dextrose** via NGT or
 - > 5ml/kg 10% Dextrose IV
 - Recheck after 30 minutes to ensure the blood glucose level has improved

12.5.8 Hypothermia

- Prevent hypothermia by keeping in hot cot/under heater and ensuring baby is well-wrapped and wearing a hat.
- When the baby is stable, it should be nursed in 'Kangaroo' position.

12.5.9 Infection

- Premature delivery may be due to maternal infection.
- Have a very low threshold for starting antibiotics in premature babies.
 Remember that signs of sepsis in a newborn can be very non-specific.

12.5.10 Respiratory Distress Syndrome

- Very premature babies lack surfactant and therefore may have marked respiratory distress.
- Put the baby on O2, cover with antibiotics and ensure they are kept warm. For babies
 <1.5kg, start CPAP with any signs of distress.

12.5.11 Apnoea of Prematurity

- Very premature babies have immature respiratory centres in the brain and sometimes 'forget' to breath.
- Use Aminophyllin as a respiratory stimulant for babies <34/40 gestation:
 6mg/kg stat, then 2-3mg/kg twice a day.

12.5.12 Neonatal Jaundice

 For all jaundiced babies, consider whether infection may be the cause and investigate (blood culture, lumbar puncture) and treat appropriately.

Investigation

- Any baby who appears visibly jaundiced should have a bilirubin level.
- If the bilirubin level is greater than the value shown in the chart below, the baby should be started on phototherapy.
- If no birilubin measurement available, start phototherapy if jaundice extends to extremities.

Phototherapy treatment thresholds for jaundiced babies

| | Healthy Term | Preterm, LBW, |
|-----------------|--|---------------|
| | Baby | Sick |
| Day Of Life | mg/dl | mg/dl |
| Day 1 | Treat any visible jaundice with phototherapy | |
| Day 2 | 15 | 13 |
| Day 3 | 18 | 16 |
| Day 4 and after | 20 | 17 |

Once Under Phototherapy

- Ensure the baby is feeding well top up with EBM via cupt or NGT if necessary.
- Babies under phototherapy should have their bilirubin level checked on a daily basis.
- Phototherapy should be stopped when the value is more than 3mg/dl below the threshold shown above.

Other Problems:

- Patent Ductus Arteriosus
 - Suspect in the premature baby with full pulses and a blowing systolic murmur.

Treatment

- Give Ibuprofen 10mg/kg Day 1, 5mg/kg Day 2 and Day 3.
- Necrotising enterocolitis
 - Suspect if abdominal distension, tenderness, bile stained vomitus, bloody stool, 'septic appearance'.

Management

- Order abdominal Xray,
- Inform surgeons
- Keep NPO
- Begin treatment with Ceftriaxone and Metronidazole.

13.0 Ophthalmic Conditions

13.1 Conjunctivitis

Presentation

- Foreign body sensation, burning, tearing, redness, photophobia and discharge. Usually bilateral though one eye may become affected 1-2 days later. On waking the eyelids are frequently stuck together and difficult to open.
- Purulent discharge, and lid oedema in infective causes

13.1.1 Allergic Conjunctivitis

- Avoid triggers when identified
 - > Mild
 - ✓ Artificial tears e.g. Hypromellose and Viscotears tds
 - ✓ Cool compresses over the eyelids
 - Moderate
 - ✓ Mast cell stabilizers (Sodium
 Cromoglycate 2% qid, Nedocromil
 Sodium 2% BD) and Antihistamines
 (Emedastine bd or qid)
 - ✓ Combined antihistamines and mast cell stabilizers (**Ketotifen** bd) can also be used

Severe

- ✓ Treat as moderate disease and a short course of low potent steroid such as **Fluorometholone** 0.1% tds, otherwise **Dexamethasone** 0.1% gutt tds may be used
 - Consider using Acyclovir
 200mg 5x/day or 400mg bid in patients on prolonged immunosuppresants.

13.1.2 Bacterial Conjunctivitis

Treatment

- Eyelids to be cleaned of discharge before using topical antibiotics
- Eye drops include Chloramphenicol 0.3%,
 Ciprofloxacin 0.3%, Ofloxacin, and,
 Moxifloxacin 0.3% maybe used.
- Eye ointments such as Chloramphenicol and Tetracycline provide higher concentrations for longer periods than drops but inappropriate for day use because of blurred vision.

13.1.3 Ophthalmia Neonatorum

Key Diagnostic features

 This is conjunctivitis developing within the first month after birth as the result of infection transmitted from mother to infant during delivery.

Prophylaxis

- Give Povidone-iodine 2.5% (a cheap and effective agent against all of the common pathogens that cause Ophthalmia neonatorum).
- Give Erythromycin 0.5% ointment or Tetracycline 1% ointment

- Give Erythromycin elixir 50mg/kg/day for 14 days with Erythromycin ointment or Tetracycline ointment qid (to treat Chlamydial infection).
- If Pneumonitis is suspected, treat for 3 weeks, and also treat the mother and her sexual partner.
- For Gonococcal infection,
 - Give Ceftriaxone 25-50mg/kg IV or IM for 7 days or Cefotaxime 100mg/kg IV or IM for 7 days
- Other bacterial infections
 - Give Chloramphenicol or Neomycin eye ointment qid
 - Systemic antibiotics may be considered in severe cases.
- For suspected Herpes simplex virus infection
 - Give Acyclovir 45-60mg/kg in 3 divided doses for 14 days and topical Acyclovir 5 times daily.

13.2 Keratitis

13.2.1 Bacterial Keratitis

Key diagnostic features

- Risk factors include contact lens wear, trauma, ocular surface disease (such as herpetic keratitis, bullous keratopathy, and dry eyes), chronic blepharitis, trichiasis, and exposure keratopathy, severe allergic eye disease and corneal anaesthesia).
- Other factors include topical or systemic immunosuppression such as diabetes, vitamin A deficiency and measles.

Presentation

- Painful red eyes, with a purulent discharge, with circumlimbal injection and with corneal opacities ± oedema ± anterior uveitis
- May complicate with limbal and scleral extensions, corneal perforations and endophthalmitis

- Topical antibiotics
 - Initially at hourly intervals day and night for up to 48 hours. Later reduced to 2-hourly during waking hours for a further 48 hours then qid for 7 days or until the epithelium has healed.

Consider using eye ointments eg
 Chloramphenicol eye ointment 2%
 or Tetracycline eye ointment 2%

Oral antibiotics

Give Ciprofloxacin 750mg bd 7-10 days or Augmentin 625mg bd 7-10 days (in threatened or actual corneal perforation or a peripheral ulcer in which there is scleral extension, for isolates for which there are potential systemic complications e.g N. Meningitides).

• Subconjunctival antibiotics

- Only indicated if there is poor compliance with topical treatment.
- Cycloplegics (Atropine 1% gutt bd or Cyclopentolate I% gutt tds).
- Topical steroids therapy to reduce corneal scarring ONLY after the ulcer has been sterilized and fungal infection has been excluded.

RED FLAGS:

- Consider using a bandage contact lens in impending or actual corneal perforation
- Counsel patient and guardians on visual prognosis

13.2.2 Viral Keratitis

Treatment of epithelial keratitis

- Give **Acyclovir** 3% ointment five times daily.
- Debridement may be used for dendritic but not geographic ulcers.
- The majority of dendritic ulcers will eventually heal spontaneously without treatment.

Treatment of Disciform Keratitis

- Initially with topical steroids (Prednisolone
 0.5% eye drops) with Acyclovir 5% eye
 ointment both qid for 4 weeks, then tapering
 of both can be done
- A weak steroid such Fluorometholone 0.1% on alternate days maybe used subsequently for several months

Treatment of Stromal Necrotic Keratitis

- Lowest effective topical steroid therapy to control inflammation
- Topical Acyclovir 5% eye ointment five times a day
- Oral Acyclovir (400mg bd for a year) to reduce the rate of recurrent epithelial and stromal keratitis. The benefit is greatest in patients with frequent debilitating bilateral disease or if involving an only eye

13.2.3 Fungal Keratitis

Treatment

- Epithelium debridement over the lesion enhances penetration of antifungal agents.
- Local treatment initially hourly for 48 hours and then reducing as signs permit.
- Use topical Natamycin 5% gutt or Econazole 1% gutt Amphotericin B 0.15% for either filamentous or candida infections.
- Subconjunctival Fluconazole may be used in severe cases with hypopyon.
- Systemic anti-fungals may be required for severe keratitis or endophthalmitis such as daily Fluconazole 100mg for a week
- Cycloplegics (Atropine 1% gutt bd or Cyclopentolate I% gutt tds.
- A broad spectrum antibiotic should also be used as bacterial co-infection is common

13.3 Immune-Mediated Uveitis

- Treatment of immune-mediated uveitis involves predominantly the use of antiinflammatory and immunosuppressive agents.
- Mydriatics
 - > Tropicamide (0.5% and 1%), Cyclopentolate (0.5% and 1%),

Phenylephrine (2.5% and 10%), and Atropine 0.5% to 1%

Steroids

- Topical steroids useful only for anterior uveitis because therapeutic levels are not reached behind the lens.
- Periocular steroids
 - ✓ Therapeutic concentrations maybe achieved behind the lens e.g. Triamcinolone Acetonide (Kenalog) and depot steroids such as Methylprednisolone Acetate (Depomedrone).
- Intraocular steroids
 - ✓ Intravitreal injection of

 Triamcinolone Acetonide (4mg in 0.1ml) or Slow-release steroid implant (Flucinolone Acetonide) via pars plana.

 Useful in patients with posterior uveitis.
- Systemic steroids
 - ✓ Give oral Prednisolone 1mg/kg
 - ✓ Intravenous injection of Methylprednisolone 1g/day for 3 days.

Antimetabolites

- Give Azathioprine 1mg/kg/day od or in divided doses. Double dose after 1-2 weeks. For Sightthreatening uveitis and as a steroid-sparing therapy in patients with intolerable side effects from systemic steroids
- ➢ Give Methotrexate 10-15mg/week (children can be given 30mg/week) as a steroid-sparing agent in patients with uveitis associated with Sarcoidosis. Folic Acid 2.5-5.0mg/day is co-administered to reduce bone marrow toxicity. Patients must refrain from alcohol.
- Give Mycophenolate Mofetil 1g bd which may be increased to 4g daily. A good alternative to azathioprine in unresponsive or intolerant patients.
 Contraindicated in children.
 Monitoring involves a weekly full blood count for 4 weeks and then monthly.

13.4 Cytomegalovirus (CMV) Retinitis

Ascertain HIV status and CD4 count. Consider drug failure or drug noncompliance

Systemic treatment

- Either of the following may be used in the treatment of CMV Retinitits
 - Give Ganciclovir 5mg/kg IV BD for 2-3 weeks, then 5mg/kg od during the induction phase. Oral Ganciclovir 300-450mg daily for prophylaxis and maintenance maybe given when retinitis is stable until CD4 count is more than 100-150 cells/μl. Ganciclovir is marrow toxic and hence the need for regular FBC checks.
 - Give Foscarnet 60mg/kg tds for up to 3 weeks for induction and 90-120mg od for maintenance. Foscarnet is nephrotoxic and causes electrolyte imbalances and seizures.
 - Give Valganciclovir 900mg bd in the induction phase and 900mg od in the maintenance phase.

Intravitreal treatment

- Intravitreal injections
 - ➤ **Ganciclovir 4mg** in 0.05ml
- Give Ganciclovir slow-release device (Vitrasert) effective for about 60 days.

13.5 Endophthalmitis

Endogenous Bacterial Endophthalmitis

- The choice of agents is dependent of on culture and sensitivity results
- Empirically treat endophthalmitis with systemic infection with
 Ceftazidime 1g bd and
 Vancomycin 1g bd for 2-3 weeks
- ➤ Isolated endophthalmitis is treated with oral **Ciprofloxacin** 750mg bd for 7-14 days and intravitreal antibiotics as in postoperative endophthalmitis below.

Postoperative Bacterial Endophthalmitis

Intravitreal antibiotics are the key to management.

> Empirical Intravitreal Injections

- ✓ Give Vancomycin (2mg in 0.1ml) and Ceftazidime (2mg in 0.1ml) OR
- ✓ Give Vancomycin (2mg in 0.1ml and Amikacin (0.5mg in 0.1ml)

Periocular Antibiotic Injections

- ✓ Give Vancomycin 25 mg in 0.5 ml and Ceftazidime 100 mg in 0.5 ml OR
- ✓ Give Vancomycin 25 mg in 0.5 ml and Amikacin 25 mg in 0.5 ml

OR

- ✓ Give Vancomycin 25 mg in 0.5 ml and Gentamycin 20 mg in 0.5 mL
- > Oral Antibiotics (Oral antibiotics are of uncertain benefit).
 - ✓ Give Fluoroquinolones (eg
 Ciprofloxacin 750mg b.d for 10 days) have a better penetration of the eye

Steroids

- ✓ Topical Dexamethasone 0.1% qid OR
- ✓ Oral steroids. High dose Prednisolone can be given if fungal infection has been excluded.
- ✓ Periocular steroids eg
 Dexamethasone 6 mg in 0.25 ml OR Triamcinolone 1mg should be considered if systemic steroids are contraindicated

13.6 Glaucoma

 This is an optic neuropathy with a characteristic visual fields defects

13.6.1 Primary Open Angle Glaucoma

Medical Treatment

- Postaglandin analogues the first line treatment, avoid in uveitis
 - > Travopost 0.004% (Travatan) od
 - > Latanoprost 0.005% (Xalatan) od
 - **Bimatroprost** 0.03% Lumigan
- Beta adrenergic antagonists, contraindicated in Asthma
 - > Selective β 1 antagonists
 - ✓ **Betaxolol** 0.5% bd
 - Non Selective β antagonists
 - ✓ Timolol 0.5% bd
 - Alpha 2 adrenergic agonists
 - ✓ Brimonidine 0.2% tds
 - Carbonic anhydrase inhibitors
 - ✓ Systemic
 - Acetazolamide 125 –250mg qid
 - ✓ Topical
 - Brinzolamide 1% (Azopt)bd
 - Dorzolamide 2%(Trusopt) tds
- Parasympathomimetics (Cholinergics)
 - Pilocarpine 2%, 4% qid gel or drops

Surgical Treatment

- Trabeculectomy when maximal medical treatment is suboptimal in IOP control
- Consider transcleral cyclophotocoagulation in eye with poor visual potential

13.7 Chemical Injury

- Treat and ask questions later
- Prognostics features of chemical corneal injuries depend on
 - ➤ The pH alkalis are more damaging than acids
 - Duration of contact
 - Corneal involvement
 - Limbal involvement
 - Associated non chemical injury such as thermal injury and blunt trauma
 - Conjunctival involvement

- Immediate irrigation with water or 2 litres of normal saline until pH is normalized
- Double evert the upper eye lids and remove any retained particulate matter
- Repeat pH after 20 minutes, if abnormal repeat the irrigation
- Daily pH tests, any derangements may indicate the presence of retained chemical particulates and warrants further irrigations and forniceal inspection

Acute management:

- Use preservative free drugs where possible
 - Preservative free topical antibiotics eg Chloramphenicol 0.5% QID
 - Topical cycloplegia eg Atropine 1% ocular BD
 - Topical lubricants eg Carmellose 1 hourly
 - Oral analgesia

13.7.1 Severe Chemical Injuries

- Admit
- Give
 - Topical steroids eg Prednisolone 1% 2-3 hourly for < 10 days –
 - ➤ Topical ascorbic acid **Sodium**Ascorbate 10% 2 hourly for < 10 days
 - Give Oral Ascorbic Acid 1g bd
 - Systemic Tetracyclines 100mg od for 3 months
 - ➤ Give **Acetazolamide** 250mg qid ± **Timolol** 0.5% bd
 - ➤ Patients need to be couselled for long term follow up to manage complications

13.8 Orbital Cellulitis

Key diagnostic features

- This is an ophthalmic emergency.
- Risk factors include sinus disease, traumatic orbital septal perforation and post orbital surgery
- Symptoms; fever, periocular pain, inflamed eye lids, proptosis, restricted and painful extraocular movements, poor visual acuity, RAPD, reduced color vision
- Investigations
 - FBC, BC, CSF analysis if meningeal or cerebral signs develop
 - CT scan of the orbits, paranasal sinuses and brain

- Hospital admission
- ENT assessment is mandatory.
- Antimicrobial therapy
 - ➤ Ceftazidime 1g IV q8h and Metronidazole 500mg IV q8h. Therapy should be continued until the patient is apyrexial for 4 days.
- Optic nerve function monitoring
- Visual acuity, color vision, pupillary reactions, and light brightness appreciation.
- Surgical intervention should be considered in orbital or subperiosteal abscesses, unresponsiveness to antibiotics and in decreasing vision.

14.0 Oral and Maxillofacial Conditions

14.1 Candidiasis/Oroesophageal

Give Nystatin oral suspension / pessaries
 100,000 units every 6 hours for 10 14 days

Note: Pessary should be sucked and taken after food

- Review after 14 days
- Paint Gentian Violet aqueous solution 0.5 % on the lesions tds for 7 days
- Give Clotrimazole Troches 10 mg tds for 4 weeks (children)

Alternatively

- Give Chlorhexidine 0.2% mouth rinses tds (should <u>not</u> be used together with Nystatin)
- If not resolved after 7 days:
- Continue with above treatment and add
 Ketoconazole 200-400 mg bd for 10-14 days
- Children: 1-4 years: Give Ketaconazole 50 mg
 bd for 10 14 days
- 5-12 years: Give **Ketaconazole** 100 mg bd for 10 -14 days

Note:

 Ketoconazole interacts with the following ARVs: Nevirapine, protease inhibitors and Didanosine

Alternatively

Adults

Give Fluconazole 50-100 mg qid for 14 days

Children

Give **Fluconazole** 6 mg/kg on day 1, then 3 mg/kg qid for 13 days

Prophylaxis:

Adults

Give Fluconazole 100 mg daily for long term

Children

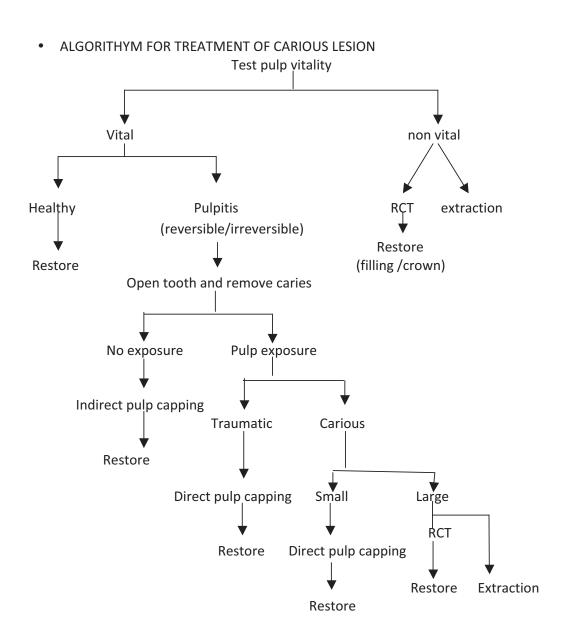
Give Fluconazole 3-6 mg/kg daily for long term

14.2 Caries, Toothache

- Depends upon the extent of caries and clinical judgment:
 - If minimal and confined to the enamel
 - ✓ Apply topical Fluoride
 - If in enamel and dentine but not involving the pulp
 - ✓ Filling

14. Oral Maxillofacial Conditions

- ➤ If involving the pulp and there is periapical infection and/ or pulp inflammation
 - ✓ Root Canal Therapy
- > If severe
 - ✓ Tooth extraction



Note:

- Antibiotics are not indicated unless there is infection
- Reinforce oral hygiene practices including use of fluoridated toothpaste to all patients

14.3 Dental Abscess

Treatment

- Consider incision and drainage
- Give Amoxycillin 250-500 mg tds for 7 days and
- Give Metronidazole 200-400 mg (Children:7.5 mg/kg/dose) tds for 7 days or
- Give Benzyl Penicillin 1-2 MU (Children: 25,000 units/kg/dose) IM or IV q6h for 7 days and
- Give Metronidazole 250-500 mg IV q8h for 7 days
- Give **Aspirin** 300 600 mg tds
- If abscess persists after 2 weeks, do culture and sensitivity
- Remove source of infection :
 - Extraction of infected tooth
 - Root canal therapy /Apicectomy /Apicectomy

Alternatively

- Give Erythromycin 250-500 mg qid for 7 days (if allergic to penicillins) and
- Give Paracetamol 1g tds for adults
 Children give Paracetamol 250mg tds

14. Oral Maxillofacial Conditions

 Give Brufen 400mg tds for adults (if Paracetamol is ineffective)
 Children give Brufen 200mg tds (if Paracetamol is ineffective)

Note: For moderate to severe pain, refer to pain management ladder

14.4 Gingivitis

Treatment

- Reinforce oral hygiene practices; i.e, brushing at least twice a day to remove plaque; in the morning after breakfast and in the evening before going to bed
- Conventional therapy; scaling and cleaning to remove all tooth surface adherents
- Gargle with antibacterial mouthwash twice a day until symptoms resolve
- Antibiotics are <u>not</u> indicated unless one has Acute Necrotizing Ulcerative Gingivitis (ANUG) or Linear Gingival Erythema (LGE)

14.5 Periodontal abscess

Treatment

- Give antibiotics as for dental abscess
- Reinforce oral hygiene practices
- Antibacterial mouthwash
- Scaling and/or root planing
- Topical fluoride application

14.6 Periodontitis

Treatment

- Root planing +/- antibacterial irrigant (Tetracycline)
- Reinforce oral hygiene practices
- Antibiotics are <u>not</u> indicated unless the following exists:
 - Necrotizing ulcerative Periodontitis (NUP)
 - Exudate discharging from the periodontal pockets
 - Patient is non-responsive to conventional therapy
 - Juvenile Periodontitis
 - Aggressive Periodontitis

14.7 Odontogenic and Maxillofacial infections

Signs and Symptoms

 Patients who present rapid progressive swelling, difficulty in breathing, elevated tongue, drooling, difficulty in swallowing (dysphagia), facial space involvement, elevated temperature, severe jaw trismus (< 10 mm), toxic appearance, compromised host defenses

Treatment

- Ensure airway patency
- Culture and sensitivity
- Incision and drainage

- Removal of the cause
- Extraction of the offending tooth or
- Treat the tooth endodontically with root canal therapy
- Sequestrectomy (removal of necrotic bone)
- Keep patient hydrated (2-3L Fluids/24 hours for maintenance)
- Give analgesics for pain relief
- Encourage high-calorie food intake
- Prescribe appropriate antibiotics for rapidly progressive swelling as follows:
 - Give Amoxycillin 250-500mg tds for 7days or Benzylpenicillin 0.5-2.0 MU IM or IV q6h for 7days
 - Give Metronidazole 200-400mg tds or 7days or Metronidazole 250-500mg IV q8h for 7 days
 - Give Tetracycline 250mg every six hours or 500 mg twelve hourly for 7 days
 - Give Erythromycin 250-500 mg qid for 7 days
 - ➤ Give Clindamycin 150-300 mg

Note: Antibiotics are indicated for diffuse swelling, compromised host defenses, involvement of fascial spaces, severe pericoronitis, osteomyelitis

14.7.1 Salivary Gland Disease

 Retention cysts, thyroglossal duct cysts, brachial cleft cysts, enlargements etc.

Treatment

- Surgical procedure (if indicated)
- Give Ciprofloxacin 250 500 mg bd for at least 5 days plus
- Give Metronidazole 200 400 mg tds for at least 5 days if there is infection

14.7.2 Trigeminal and Glossopharyngeal Neuralgia

Signs and Symptoms

Unilateral pain with trigger zone

Treatment

- Give Carbamezepine 600-1,600mg
 nocte for a month then review
- Give **Phenytoin** 300-500mg tds per day
 - Can be administered by IV for severe TN pain
- Give Gabapentin 300mg tds
- Give Baclophen 5mg bd or tds a day which can be increased
 - Usual effective dose is 50-60mg per day
 - Can be used alone or in combination withCarbamezepine

Note: If trigeminal neuralgia persists after 2 months refer to neurosurgery

14.8 Local Anaesthesia Toxicity in Dental Surgery

14.8.1 Mild Local Anaesthetic toxicity

Signs and Symptoms

 Talkativeness, anxiety, slurred speech, confusion

Treatment

- Stop administration of local anaesthetic
- Monitor vital signs
- Observe for 1-hour

14.8.2 Moderate Local Anaesthetic toxicity

Signs and Symptoms

 Stuttering speech, nystagmus, tremors, headache, dizziness, blurred vision, drowsiness

Treatment

- Place in supine position
- Monitor vital signs
- Administer oxygen
- Observe for 1-hour

14.8.3 Severe Local Anaesthetic toxicity:

Signs and Symptoms

Seizure, cardiac dysrythmia or cardiac arrest

Treatment

- Place in supine position
- If seizures, protect from nearby objects,
- Suction oral cavity if vomiting occurs

14. Oral and Maxillofacial Conditions

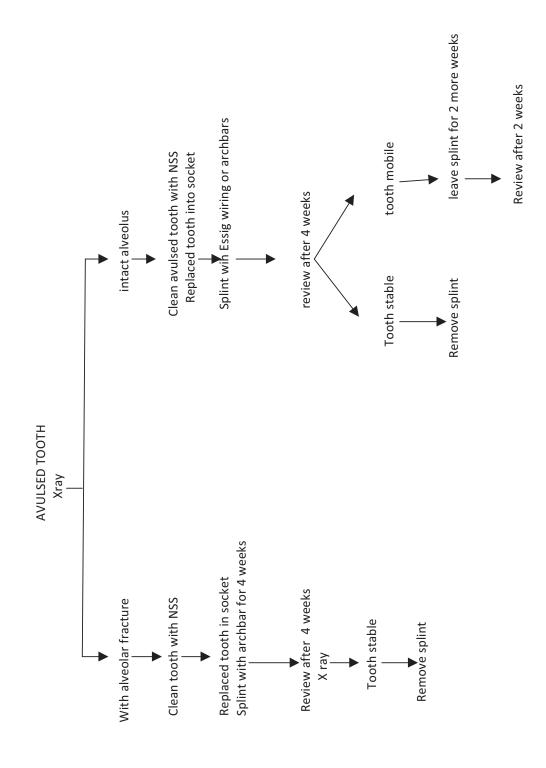
- Administer oxygen
- Give **Diazepam** 5-10 mg IV slowly as stat dose
- Transport to emergency care facility / intensive care unit
- Monitor vital signs
- Summon for medical assistance

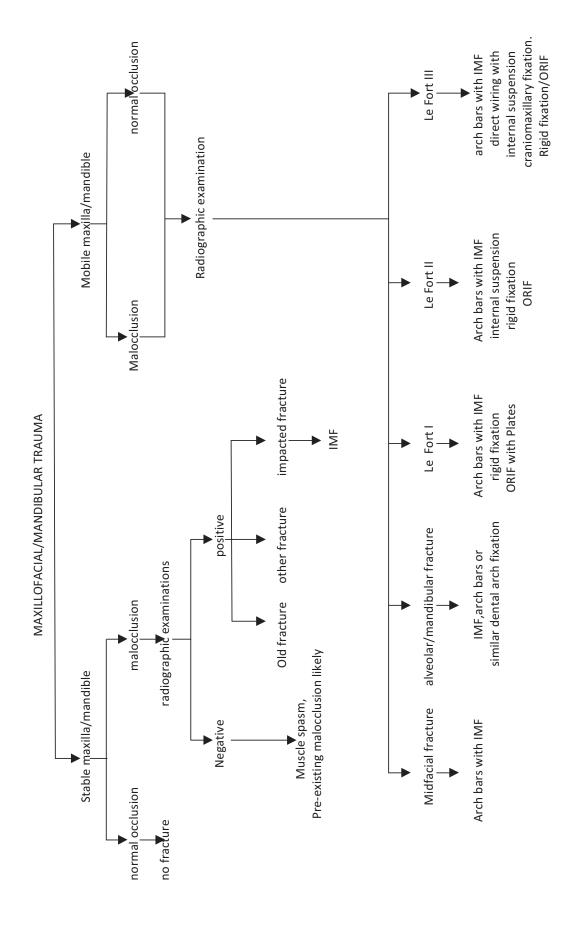
14.9 Mouth Ulcers (Sores)

- If related to HIV infection, please refer to section on Management of the HIV-Related Diseases,
- All ulcers in the mouth regardless of the HIVstatus, lasting more than three weeks, should be referred for further management

14.10 Oral Trauma

- Clean the area with NSS
- Check for fractures and break in the skin or mucosatreat accordingly
- Sutures and simple fractures can be treated under local anesthesia
- Complex and complicated fractures maybe treated under LA and premedicated with pethidine 100mg IM for 18 years and above
- Fractures not suited under local anesthesia must be treated under general anesthesia

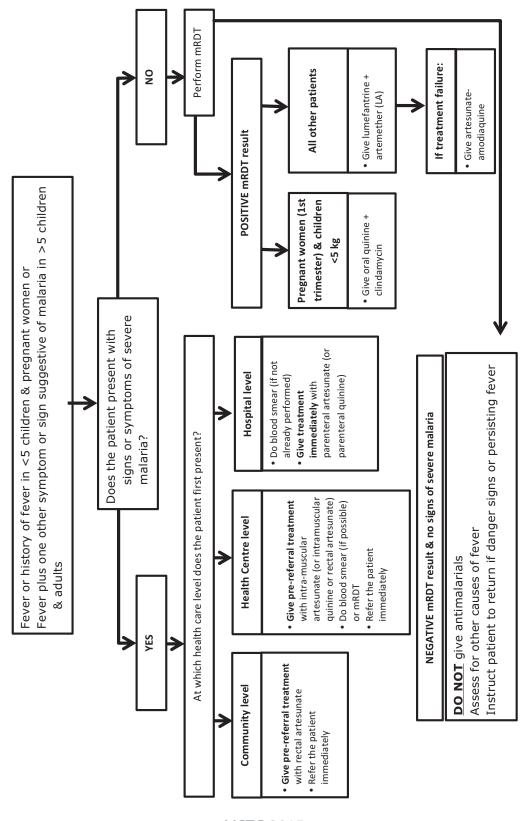




15.0 Parasitic diseases

15.1 Malaria

Flow Chart for the Diagnosis and Treatment of Malaria



15.1.1 Malaria, non-severe, uncomplicated

 Refer to the MOH National Malaria Control Programme Revised Guidelines for the Treatment of Malaria in Malawi, July 2013, for full details of malaria management

Signs and Symptoms

 Fever or recent history of fever in pregnant women or children under the age of five years and fever or history of fever plus one other symptom or sign suggestive of malaria in over five children and adults

Diagnosis

- All suspected uncomplicated malaria cases at all levels of the health care delivery system should be tested using malaria rapid diagnostic tests (mRDTs) or by microscopic examination of blood film wherever possible
- Testing (film or mRDT)
- Repeat diagnostic test (film or mRDT) if:
 - the first test was positive and there is persistent fever or worsening condition despite suitable antimalarial treatment

the first test was negative and antimalarial treatment was not given, but the patient's fever persists or condition deteriorates

Treatment

 In the event that both mRDT and microscopy are not available, treat on the basis of presumptive diagnosis

First-Line Treatment

- Give Lumefantrine 120mg/Artemether 20mg (LA)
- Lumefantrine-artemther comes in two oral formulations:
 - Non-dispersible LA [LA(ND)] for older children weighing 25kgs or more including adults and
 - Dispersible LA [LA(D)] for children weighing from 5kg to less than 25kg

Note: If LA(D) is not available, LA(ND) can be used to treat children weighing from 5kg to less than 25kg

Dosage Schedule for dispersible (D) and nondispersible (ND) Lumefantrine-Artemether (LA-120mg/20mg tablets)

| Body weight in Kg (age in | No. of tablets at approximate timing of dosing | | | | | |
|------------------------------|--|----------------|-------|----|-------|----|
| years) | Day 1 | | Day 2 | | Day 3 | |
| | Start dose | After 8 hrs | AM | PM | AM | PM |
| LA(D) 5-14.9 kg (<3) | 1 | 1 | 1 | 1 | 1 | 1 |
| LA(D) 15-24.9 kg (≥ 3-8) | 2 | 2 | 2 | 2 | 2 | 2 |
| LA(ND) 25-34.9 kg (≥9-14) | 3 | 3 | 3 | 3 | 3 | 3 |
| LA(ND) ≥ 35 kg (>14) | 4 | 4 | 4 | 4 | 4 | 4 |

- First dose should be given as DOT. If vomiting occurs within 30 minutes, repeat the dose
- Dose is given according to body weight
- If possible each dose should be taken with milk, which improves the absorption of lumefantrine component of the combination
- If fever persists beyond 72 hours, do malaria microscopy, and if the result is positive, give second line treatmentIn patients with contractions or intolerance to LA give second line treatment

Second-Line Treatment

 Give Artesunate 4 mg/kg/day and Amodiaquine 10 mg/kg/day od for 3 days

Dosage schedule for fixed combination dose of artesunate-amodiaquine

| Body Weight (kg) | Age | Daily dose for 3 days artesunate-amodiaquine | Preparation strength per tablet |
|------------------------|------------------------------|--|---------------------------------|
| 5.0 – | 2 – 11 | 1 tablet | 25 mg/67.5 |
| 8.9 | months | | mg |
| 9.0 – | 1 – 5 | 1 tablet | 50 |
| 17.9 | years | | mg/135 mg |
| 18.0 – | 6 – 13 | 1 tablet | 100 mg/270 |
| 35.9 | years | | mg |
| ≥ 36 | 14 years old and above | 2 tablets | 100 mg/270 mg |

Note:

- Treatment failure to first line treatment (LA) should be suspected if symptoms persist or the patient clinically deteriorates three to 14 days after initiation of LA drug therapy
- Side effects: Transient rise in transaminases and transient reduction in white blood cell count. The dosing schedule is indicated in Table 1.4 below
- Pregnant women in the first trimester with a confirmed treatment failure to quinine plus clindamycin should be treated with LA

If pregnant and in first trimester and children weighing < 5kg

- Give oral Quinine plus Clindamycin
- The table below shows the recommended dosages for oral Quinine and Clindamycin:

Dosage schedule for oral Quinine plus Clindamycin

| Population | Recommended dosage |
|-------------------------------------|--|
| Pregnant women in the 1st trimester | Quinine tablets 600 mg tds for 7 days plus Clindamycin tablets 300 mg tds for 7 days |
| Children < 5 kg | Quinine (10 mg salt/kg body weight) tds for 7 days plus Clindamycin (20 mg base/kg body weight) tds for 7 days |

Second Line Treatment in pregnancy during first trimester

 If confirmed treatment failure to quinine plus clindamycin, mother should be treated with LA

15.1.2 Severe Malaria

- Most severe malaria occurs in children under 5 years of age
- Severe malaria is a medical emergency and as such treatment should begin immediately, whether the patient presents at the community, health centre, or hospital level
- Suspect severe malaria if a patient has one or more of the following conditions (mostly seen in combination):

Clinical manifestations and some laboratory findings

| Cillical maintestations and some laboratory midnigs | | | |
|---|--------------------|------|--------------------------|
| Clinica | l manifestations | Some | laboratory findings |
| • | Impaired level of | • | Severe anaemia: |
| | consciousness | | (Hb<5 g/dl) (i.e. Hb <5 |
| | (cerebral malaria) | | g/dl or Hct < 15 %) |
| • | Respiratory | • | Hypoglycaemia: (<2.2 |
| | distress (acidotic | | mmol/l or <40 mg/dl) |
| | breathing) | • | Hyperlactataemia |
| • | Repetitive | | (lactic acidosis) (blood |
| | convulsions | | lactate >4 mmol/l) |
| • | Circulatory | • | Electrolyte imbalance |
| | collapse | | (hyponatraemia) |
| • | Pulmonary | • | Acute kidney injury |
| | oedema | | (serum creatinine |
| • | Prostration | | >265 µmol/l) |
| • | Excessive or | • | Haemoglobinuria |
| | persistent | | |
| | vomiting | | |
| | | | |

- Extreme pallor
- Shock (weak pulse, cold extremities)
- Jaundice (yellowish coloration of eyes)
- Little or no urine output (think about acute kidney injury)
- Very dark coloured urine
- Spontaneous bleeding (mouth, nose, skin, eyes)

Hypovolaemia

- Although most children with malaria have a (history of) fever, this may be variable in patients who have progressed to severe malaria
- Examine children with suspected severe malaria for other conditions (e.g. pneumonia, meningitis) as a possible cause of their symptoms and, if found, manage appropriately

Note:

Patients with hyperparasitaemia: 4+ (40,000 – 400,000/µl or ring stage ≥ 5% of RBCs) who do not have any of these indicators of severe (disease) malaria should be admitted for observation. Treat with first-line antimalarial (LA).

 If severe malaria is diagnosed in an outpatient, refer the child for hospitalization (see below)

15.1.2.1 Pre-referral treatment at Community level

 Refer any patient with severe malaria to the nearest hospital

Treatment

- Give Rectal Artesunate at 10 mg/kg body weight in a single dose, followed as soon as possible by definitive therapy for severe malaria at a hospital
- In the event that Artesunate Suppository is expelled from the rectum within 30 minutes of insertion, a second suppository should be inserted
- If referral is not possible within 12 hours, a second dose of Rectal Artesunate should be administered at 12 hours after the initial dose, then once in every 24 hours until patient is transferred to a hospital

Initial (pre-referral) Dosage of Artesunate Suppositories for patients aged ≥16 yrs

| Weight (kg) | Artesunate dose | Regimen (single dose) |
|-------------|-----------------|---|
| <40 | 10 mg/kg bw | Use appropriate no. of 50 mg rectal suppositories |
| 40 –59 | 400 mg | Two suppositories of 200 mg each |
| 60 – 80 | 800 mg | Four suppositories of 200 mg each |
| >80 | 1200 mg | Six suppositories of 200 mg each |

Note:

- For children, hold the buttocks together for 10 minutes to ensure retention of the rectal dose
- Treatment with Rectal Artesunate is suboptimal and every effort should be made to refer the patient as soon as possible
- The table below shows the recommended pre-referral doses of Artesunate
 Suppositories for children aged <16 years
 - As in adult patients, if referral is not possible within 12 hours, a second dose of **Rectal Artesunate** should be administered at 12 hours after the initial dose

➤ Thereafter the dose may be repeated every 24 hours. Refer to the malaria treatment guidelines for RA insertion procedure

Initial (pre-referral) Dosage of Artesunate Suppositories for Children Aged 2months –15 Years (and weighing at least 5 kg)

| Weight (kg) | Age | Artesunate dose (mg) | Regimen (single dose) |
|-------------|---------|----------------------|-----------------------|
| 5 – 8.9 | 2 – 12 | 50 | One 50 mg |
| | months | | suppository |
| 9 – 19 | 13 – 42 | 100 | Two 50 mg |
| | months | | suppositories |
| 20 – 29 | 43 – 60 | 200 | One 200 mg |
| | months | | suppository |
| 30 – 39 | | 300 | One 200 mg |
| | 6 – 13 | | suppository + |
| | years | | Two 50 mg |
| | | | suppositories |
| >40 | >14 | 400 | Two 200 mg |
| | years | | suppositories |

15.1.2.1 Pre-referral treatment at Health Centre Level

Treatment

- Give Artesunate 2.4 mg/kg (0.12 ml/kg) IM
 - Artesunate should be given by intramuscular injection into the upperouter quarter of anterior thigh and should not be injected into the buttocks
 - To administer IM **Artesunate**, weigh the patient and determine the number of vials needed for treatment as per the table below:

Number of Required Vials of Parenteral Artesunate by Body Weight

| Weight | 60 mg vials required |
|----------------|----------------------|
| 5 kg – 25 kg | 1 |
| 26 kg – 50 kg | 2 |
| 51 kg – 75 kg | 3 |
| 76 kg – 100 kg | 4 |

- Each 60 mg vial of injectable Artesunate must be reconstituted with 1 ml of Sodium
 Bicarbonate
- Dilute the Artesunate-Bicarbonate mixture with 2 ml of 5% Dextrose Solution or Normal Saline (0.9% Sodium Chloride) to produce a 20 mg/ml solution

- Withdraw the appropriate volume in a syringe ([2.4 mg x body weight in kg]/20 mg/ml) for intramuscular injection, rounding to the next whole number in milliliters
- Administration of pre-referral IM Artesunate should be followed as soon as possible by definitive therapy for malaria at a hospital
- If referral is not possible within 12 hours, a second dose of IM **Artesunate** should be administered at 12 hours after the initial dose
- If referral is still not possible after 24 hours, a third dose of IM Artesunate should be given

Alternatively

- If IM Artesunate is unavailable or contraindicated, treat with high dose Quinine IM, administered in the thigh not the buttock
- Give IM **Quinine** 10 mg (0.2 ml) per kg body weight

 If the volume to be injected exceeds 3 ml, give half into each thigh. An example of body weights and dosing (ml) for IM quinine is given in the table below.

Dosage of Parenteral Quinine per body weight

| Body weight | Quinine (ml) | Number of injection sites |
|----------------|-----------------|---------------------------|
| Under 5 kg | 1.0 ml | 1 |
| 5.1 – 7.5 kg | 1.5 ml | 1 |
| 7.6 – 10.0 kg | 2.0 ml | 1 |
| 10.1 – 12.5 kg | 2.5 ml | 1 |
| 12.6 – 15.0 kg | 3.0 ml | 1 |
| 15.1 – 17.5 kg | 3.5 ml | 2 |
| 17.6 – 20.0 kg | 4.0 ml | 2 |
| 20.1 – 22.5 kg | 4.5 ml | 2 |
| 22.6 – 25.0 kg | 5.0 ml | 2 |
| 25.1 – 27.5 kg | 5.5 ml | 2 |
| 27.6 – 30.0 kg | 6.0 ml | 2 |

- Administration of pre-referral IM Quinine should be followed as soon as possible by definitive therapy for malaria at a hospital
- If referral is not possible within 12 hours, a second dose of IM quinine should be administered 12 hours after the initial dose
- If referral is still not possible after 24 hours, a third dose of IM **Quinine** should be given
 - ➤ Give 0.4ml/kg of this solution as the first (loading) dose this is 20mg/kg

- Subsequent (12-hourly) doses should each be 0.2ml/kg (10mg/kg)
- ➤ The dose of **Quinine** for an adult at any one time should not exceed 1,200mg

Injectable artesunate or quinine should be for patients unable to take oral drugs.

- Where there is no scale, weight of the child can be estimated as follows:
 - For children of 3months to 12months old
 - ✓ Weight (Kg) = Age (months) + 9/2
 - For children of 1 year to 6years old
 - ✓ Weight (Kg) = [Age (in years) x 2] + 8

If IM Artesunate and IM Quinine are unavailable

• Give Rectal Artesunate

15.1.2.2 Additional management and supportive measures

- Reduce fever:
 - Tepid sponging with lukewarm (not cold) water
 - Give an antipyretic (paracetamol 10 mg/kg; 6 to 8-hourly) as required until fever is reduced. See dose tables in Section 10.1

Start resuscitation, particularly maintenance of a patent airway.

Establish IV line.

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Take 8 immediate measures:

- Ensure adequate fluid intake:
 - ➢ If the patient can drink: give ORS 20 mL/kg plus one added teaspoon of glucose powder or sugar
 - If the patient cannot drink: use nasogastric tube to provide this fluid
 - Repeat after 2-4 hours

15.1.2.3 Management of Severe Malaria Patient in Hospital Setting

 Consider the following (8-8-8) points in the management of patients with severe malaria

Make a thick blood smear for immediate malaria parasite count, (if microscopy is not available, an mRDT may be useful to indicate whether malaria infection is present or not)

Classify the degree of dehydration, assess patient's fluid requirements and correct accordingly.

Control fever if the axillary temperature is 38.5°C or above: Tepid sponge, fanning and oral or rectal paracetamol (15 mg/kg every 4 to 6 hours).

Control convulsions: maintain airway, treat with rectal diazepam (0.5 mg/kg) or slow IV diazepam (0.3 mg/kg, maximum 10 mg in an adult), or paraldehyde 0.1 ml/kg IM.

Remember to correct any hypoglycaemia or hyperpyrexia in a convulsing patient.

Detect and treat hypoglycaemia: hypoglycaemia can be induced by high parasitaemia, fasting and quinine therapy. Hypoglycaemia can recur, especially in pregnant women and children. If blood glucose ≤3 mmol/l or ≤54 mg/dl; give 1 ml/kg of 50% dextrose IV, diluted with an equal volume of 0.9% saline or 5% dextrose, give slowly over 3-5 minutes, and check blood glucose after 30 minutes and as required after treatment. Follow with 10% dextrose infusion at 5 ml/kg/hr. If there is no test for blood glucose, treat as if the patient is hypoglycaemic.

Start intravenous or intra-muscular artesunate. Dosage schedule is provided from section 2.2.2.2 below. If intravenous or intra-muscular artesunate is unavailable, use intravenous or intra-muscular quinine.

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Look for and Deal with the Following 8 Complications:

- 1. **Shock**: If cold peripheries, delayed capillary refill, or Systolic BP <50 mmHg in children 1 5years or <80 mmHg in >5 years, suspect Gram-negative septicaemia. In such cases take blood samples for culture. Give parenteral broad-spectrum antimicrobials. Correct fluid disturbance, and then continue with maintenance fluid as follows:
 - for children weighing <10
 kg, give 4 ml/kg/hr.;
 - for children weighing 10 –
 20 kg, give 40 ml/hr. plus additional 2 ml per kg for each kg of weight in excess of 10 kg;
 - for children weighing >20 kg, give 60 ml/hr., plus additional 1 ml per kg for each kg of weight in excess of 20 kg. Give oxygen if possible.

- 2. Altered consciousness and/or convulsions: Check for hypoglycaemia, hyperpyrexia and 'subtle' seizures. In a comatose patient, convulsions (seizures) may be 'subtle' i.e. minor movements [flicker of eyelid, mouth or finger] or unusual repetitive movements [a rhythmical cry, unusual breathing, or 'pedalling' of legs]. If seizure suspected, treat as in item 6 in Box1.
- 3. **Severe anaemia:** Consider the need for blood transfusion: Assess the degree of pallor (no pallor, some pallor or severe pallor look especially at palms of hands, also mucous membranes). Assess signs that increase the danger of severe anaemia respiratory distress, altered consciousness, shock and hyperparasitaemia.

Note: The decision to transfuse with blood should not only be based on low laboratory values, but on a full assessment of the patient**. As a guide, all patients with PCV<12% or Hb<4 g/dl should be transfused, whatever the clinical state; those with any of the above danger signs may be transfused even if PCV is 13-18% or Hb 4-6g/dl.

- 4. Transfuse packed red cells in most cases; in shock or severe acidosis, use whole blood. The volume transfused should be 20 ml/kg.
- 5. Metabolic acidosis (deep, fast breathing): exclude or treat hypoglycaemia, hypovolaemia and gram negative septicaemia. Give isotonic saline 20 ml/kg of body weight rapidly or screened whole blood 10 ml/kg if PCV <18% or Hb<6 g/dl. Consider lactic acidosis and enquire whether the patient has been taking ART (lactic acidosis is a side effect of stavudine).
- Spontaneous bleeding or coagulopathy: 6. If patients have underlying malnutrition, concomitant hepatic obstruction and bile salt excretion defects or prolonged fasting for more than 3 days, transfuse screened fresh whole blood, give Vitamin K 10 mg IV slowly once a day for 3 days. For Children give 2 – 3 mg/day slow IV. Vitamin K injections should not be given to "all" severe malaria patients with spontaneous bleeding, the risks and benefit of Vitamin K administration should be considered. Serious adverse events of Vitamin K injection include hypotension, difficulties in breathing, bradycardia or anaphylaxis.

- a) Acute pulmonary oedema in adults: prevent by avoiding excessive rehydration. Treatment: prop patient up; give oxygen. Stop IV fluids if pulmonary oedema is due to over-hydration, give a diuretic (furosemide IV 40 mg for adult and 0.5 1 mg/kg/dose for children).
- b) Acute respiratory distress syndrome: supportive treatment +/- ventilation
- 7. Acute kidney injury in adults: detect this by monitoring fluid balance. Identify and correct any dehydration or hypovolaemia. Maintain strict fluid balance. Consider peritoneal dialysis if oliguria persists beyond a few days.
- 8. **Common infections** and other conditions that present like severe malaria: Perform urinalysis, lumbar puncture (unless contraindicated), blood culture if possible, and chest x-ray.

Box 3: Monitor the Following 8 Observations:

Where possible use Critical Care Pathways (CCPs).

- 1. Level of consciousness (using coma score)
- 2. Vital signs every 4 hours (temperature, pulse, respiration, blood pressure)
- 3. Fluid balance (urine volumes, intake volumes IV and oral puffy eyes, chest crepitation, elevated jugular venous pressure)
- 4. Increasing anaemia (pallor, heart failure with increasing liver size)
- 5. Occurrence of convulsions see item 2 in previous Box
- 6. Blood glucose every 4 hours while unconscious and also if convulsions occur
- 7. [Hb]/Packed Cell Volume at least daily, or more often if anaemia is suspected
- 8. Ability to suck, drink, eat, sit and walk measures of overall strength.

15.1.2.4 Management of Severe Malaria in Paediatric In-Patients

Treatment

- Give Artesunate 2.4 mg/kg body weight IV (for both adults and children) on admission (at 0 hour)
 - Repeat at 12 hours and 24 hours, after initiating the first dose then once daily for not more than six days
 - Switch to LA once the patient can take oral treatment after at least 24 hours of Parenteral Artesunate
 - There should be an interval of at least 8 hours between the last dose of Artesunate and the first dose of LA

Alternatively

Give Artesunate 2.4 mg/kg body weight IM into the upper-outer quarter of anterior thigh if intravenous bolus is not feasible
 Note: Artesunate solution should be freshly prepared prior to administration and should never be stored

In case **Artesunate** is not available or is contraindicated, then

- Give Parenteral Quinine
- Refer to section 15.1.2.1 to determine the number of vials needed for treatment

- For children, Quinine IV is administered as follows:
 - Initial (loading) dose 20 mg (Quinine Salt)/kg body weight: inject this dose into 10 ml/kg of 5% Dextrose or half strength Darrow's and infuse over 3-4 hours
 - If patient has already received **Quinine** for this illness, the first dose IV infusion should be 10 mg/kg diluted as above and given over 3-4 hours with no loading dose
 - Subsequent doses of 10 mg/kg should be given every 12 hours
 - ➤ The infusion should run for 3 4 hours. Continue the 5% **Dextrose** or half strength Darrow's IV fluid (10 ml/kg given over 3 – 4 hours) between doses of quinine
 - Switch to LA once the patient can take oral treatment after at least 24 hours of Parenteral Quinine

Note: LA should only be taken 12 hours after last dose of quinine to avoid cardiotoxicity

15.1.2.5 Management of Severe Malaria in Adult In-Patients

Treatment

If the patient can be weighed, intravenous
 Quinine is administered in the same manner as for children

- If the patient cannot be weighed, IV Quinine should be given as follows:
 - First dose 900 mg in one litre of 5%
 Dextrose or ½-strength Darrow's Fluids
 given over 3 4 hours
 - Subsequent doses 600 mg in one litre 5%
 Dextrose or ½-strength Darrow's Fluids
 q12h given over 3 4 hours
 - ➤ Continue the same IV fluids or Ringer's

 Lactate (10 ml/kg given over 3 4 hours)

 between doses of Quinine (Give a

 maximum of about 3 litres per 24 hours to
 avoid fluid overload)
 - Stop intravenous Quinine as soon as the patient can take food and fluids orally and at least 24 hours of Parenteral Quinine has been administered

Note: What if 60+ kg?

 Give the appropriate dose of LA beginning 12 hours of the last dose of quinine for 3 days. For pregnant women in the first trimester give oral quinine plus clindamycin for a total of 7 days

Complication that may arise in adults

- Apart from cerebral malaria and anaemia, in adults other complications may develop such as:
 - Acute renal failure
 - Respiratory distress syndrome (presenting as severe breathlessness)

- Disseminated intravascular coagulation (DIC) – presenting as prolonged or spontaneous bleeding
- Jaundice from severe haemolysis or liver cell damage
- Management must be appropriate to each complication that develops
- Fluid and antimalarial drugs are given as for children

15.1.4 Treatment of Severe Malaria in Pregnancy

- Special attention must be paid to anaemia, hypoglycaemia and pulmonary oedema
- See below for further information on the management of complications

Box 4: Management of complications: (see The 8-8-8 schedule above)

Manage complication as for any adult. Of special importance in pregnancy are:

- Pulmonary oedema: careful fluid management, diuretics if necessary, oxygen if possible, nurse patient in semiupright position.
- Hypoglycaemia: consider this complication if there is altered consciousness or seizure.
 - Treat as in Item 2 in Box 2.
- Anaemia: be prepared for blood transfusion, especially if the patient is close to parturition. Otherwise, indications for blood transfusion are the same as in others – (see Box 2).
- Acute kidney injury: a particular danger if there has been eclampsia or shock.
 Identification and management as above.
- Shock: consider concealed haemorrhage, continuing blood loss, and septicaemia.
 Pay special attention to fluid needs.
 Culture blood if possible. Administer broad spectrum antibiotics in addition to quinine.

Treatment

- Parenteral artesunate is the recommended treatment for severe malaria in the second and third trimesters of pregnancy.
- Refer to 15.1.2.5
- Parenteral Quinine is the recommended treatment for severe malaria in the first trimester of pregnancy
 Note: Random blood glucose should be measured before and after quinine administration
- Shift to oral Quinine plus Clindamycin (during 1st trimester) and LA (in 2nd and 3rd trimester) as soon as the patient is able to take oral medication and at least 24 hours of parenteral therapy has been administered (refer to malaria treatment guidelines for details)

15.1.5 Malaria: Selective Chemoprophylaxis

• The appropriate regimen for an individual depends on the circumstances.

15.1.5.1 Risk Groups

- The following high risk groups should be given antimalarial cheprophylaxis:
 - Patients with immunosuppression caused by illness (e.g. Leukaemia, but not HIV infection or malnutrition) or splenectomy

- > Tropical splenomegaly syndrome
- Under 5s with recurrent febrile convulsions
- Individuals with sickle cell disease
- Non-immune visitors (i.e. visitors from non-malarial countries)
- Pregnant women

15.1.5.2 Antimalarial Prophylaxis Regimens

- Give **Mefloquine** (Lariam) 250 mg weekly
 - Contraindicated in pilots, people with history of cardiac disease, neurological disease or depression, and in those taking beta-blocking drugs
- Give Atovaquone-proguanil ('Malarone') –
 one tablet daily
- Take for only one week after exposure end
 Give Chloroquine 300 mg 2 tablets weekly
 - Should be combined with daily proguanil (see below)
 - Chloroquine causes itching in 40% of black people. Contraindicated in persons with psoriasis or epilepsy
 - Risk of retinal damage if taken every week for more than 6 years - advise a change

- Give **Proguanil** (Paludrine®) 200 mg daily
 - Should combine with an additional drug such as weekly Chloroquine

15.1.5.3 Intermittent Presumptive Treatmnet of Malaria in Pregnancy (IPTp)

- Intermittent Presumptive Treatment of malaria in pregnancy (IPTp) is one of the major malaria preventive strategies in Malawi
- Pregnant women should receive at least three doses of Sulfadoxine-Pyrimethamine (SP)
 525mg after the first trimester
- Administer three tablets of SP with each scheduled antenatal care visit after quickening
- The doses should be administered at least four weeks apart and given as directly observed therapy (DOT)
- The last dose of SP can be delivered safely up until the time of delivery
- **Sulfadoxine-Pyrimethamine** can be given either on an empty stomach or with food

Note: HIV positive women receiving Cotrimoxazole Prophylaxis should not receive SP

15.2 Onchocerciasis (River blindness)

 Occurs mainly in highland areas such as Thyolo

Treatment

Adults and children > 5 kg:

Give Ivermectin 150 mcg (0.15 mg)/kg single dose

Children<5 years:

- ➤ Give **Albendazole** 200mg
- Repeat annually or on return of symptoms
- No food for at least 2 hours before or after dosage
- Mothers should not breast-feed during treatment

Ivermectin dose table

| Weight | Height | Age | N | Total |
|--------|---------|---------|-----|-------|
| (kg) | (cm) | (years) | 0 | (mg) |
| > 64 | > 158 | > 40 | 4 | 12 |
| 45-64 | 141-158 | 25-39 | 3 | 9 |
| 26-44 | 120-140 | 14-24 | 2 | 6 |
| 15-25 | 90-119 | 5-13 | 1 | 3 |
| <15 | <90 | <5 | n/r | n/r |

^{*}ivermectin 3 mg tablets n/r = not recommended

- **Ivermectin** is not recommended for:
 - Pregnant women (or for those who think they might be) – it is often possible to delay treatment for a few month
 - Breast-feeding mothers with babies < 1 week old</p>
 - Children < 5 years old (i.e. < 15 kg or 90 cm height)</p>
 - Seriously ill patients

15.2 Schistosomiasis (Bilharzia)

Prevention

- Health education
- Vector control
- Improved sanitation and water supply
- Modification of the environment, e.g. clearing of vegetation in certain areas
- Avoidance of re-infestation

15.3.1 Schistosomiasis Haematobium

Treatment

- Give Praziquantel 40 mg/kg as a single dose
- Children below 4 years of age Praziquantel
 20mg/kg as stat dose

15.3.2 Schistosomiasis Mansonii

- Consider this diagnosis in cases of:
 - Unexplained chronic abdominal complaints with hepatosplenomegaly
 - Ascites with splenomegaly
 - Chronic bloody diarrhoea with no fever
 - Paraparesis/paraplegia
- Refer patients with above clinical features
- Treat after laboratory confirmation

Treatment

- Give Praziquantel 40 mg/kg as a single dose.
- Children below 4 years of age give Praziquantel
 20mg/kg as stat dose

15.4 Trypanosomiasis (Sleeping Sickness)

- Suspect in any patient presenting with fever from areas near:
 - Wildlife Reserves: Vwaza, Nkhotakota, Majete, Mwabvi
 - National Parks: Kasungu, Liwonde, Lengwe
 - Phirilongwe (Mangochi), Machinga, Mwanza
 - Lower Shire borders with Mozambique or from any other areas where Tsetse fly is found
- Suspect in children from these areas who remain sick after presumptive malaria treatment
- Increased suspicion in any sick patient from these areas with a history of:
 - Headache, Vomiting, Weakness, Changes in mood, Convulsions, Drowsiness, Mental slowness
- Travel history is very important
- Suspect also in any patient from these areas where the cause of illness is not otherwise apparent

- Trypanosomiasis can be acute in children (resembling malaria) and can be more chronic in adults
- Early stage trypanosomiasis may cause myocarditis
- Examination may reveal anaemia, lymph gland enlargement and spleen enlargement
- Nearly all cases have a hard and painful subcutaneous nodule (chancre) which is evidence of an infected bite

Procedure at Health Centres

- Refer the patient immediately to the nearest hospital
- Request close family members of the patient to undergo examination at the hospital as they may also be infected

Hospital management

- Request for a thick blood smear
- If negative more tests will be needed to confirm this diagnosis
- If diagnosis is confirmed by blood smear or other blood test:
 - Start Suramin
 - > as follows: Day 1: 5mg/kg
 - > Day 2: 20mg/kg
 - > Day 3: Do a lumbar puncture
- If LP is normal (stage 1 trypanosomiasis): give
 Suramin 20mg/kg on day 3,
 10, 17, 24, 31

- If LP is abnormal (stage 2 or CNS trypanosomiasis): stop suramin, start
 Melarsprolol (Mel-B) as follows:
 - > Day 3: 1.2mg/kg
 - > Day 4: 2.4mg/kg
 - > Day 5: 3.6mg/kg
 - > Day 6: 3.6mg/kg
- Repeat this 4-day Melarsprolol cycle after one and two weeks

Notes on treatment regimen

- If any medicine reaction occurs (e.g. skin rash, exfoliative dermatitis, reactive encephalitis) stop treatment and inform the clinical officer or medical officer immediately
- Do a lumbar puncture (LP) on day 3.
 Subsequent treatment depends on whether this is found to be normal or abnormal
- Freshly reconstitute the Suramin (Sur) 1 g vial of powder with 10 mL water for injection to make a 10% solution (100 mg/mL)
- Add the required dose of 20 mg/kg (0.2 mL of injection/kg) up to a maximum of 1 g (the whole vial) in adults of 50 kg or over to 200 mL of dextrose 5% and infuse over 2 hours. Alternatively give the dose as a slow IV injection.

- Melarsoprol (Mel B) dose is 3.6 mg/kg (=0.1ml/kg). Give this as a slow IV push. Take great care to avoid extravasation as the medicine is highly irritant. In adults of 50 kg or over the dose is the maximum permissible 180 mg (i.e. one 5 mL ampoule)
- Prednisolone may be added to Melarsprolol with a dose of 40mg once daily. The dose in children is 1 mg/kg once daily
- Control any seizures with Diazepam 5-10 mg slow IV with or without the addition of Phenytoin 150-300 mg as a single daily dose taken with water
- Anti-trypanosomal treatment may cause abortion in pregnancy, but this must be regarded as an unavoidable risk
- Follow-up: review the patient for repeat blood film and LP at 3, 6, 12 and 24 months posttreatment

15.5 Worm Infestations

15.5.1 Ascaris, Enterobius, Ancylostoma, Trichuris Infestation

- Ascaris limbricoides = roundworm,
 Enterobius vermicularis = pinworm,
 Ancylostoma duodenale = hookworm,
 Trichuris trichiura = whipworm
- Hookworm can contribute considerably to anaemia especially in children

Treatment

Adults and Children > 2 years:

Albendazole 400 mg single dose

Children below 2 years

Albendazole 200mg

NOTES:

- Albendazole (and Mebendazole) are contraindicated in pregnancy
- Heavy trichuris infections generally require treatment for 3 consecutive days
- In enterrobiasis, all family members must be treated concurrently

15.5.2 Strongyloides Stercoralis Infestation

 A hyper-infection syndrome in immunosuppressed persons can occur and may be lethal and therefore requires specialist treatment.

Treatment

 Give Ivermectin 200mcg/kg daily for 2 days Alternatively

Adults

Give Albendazole 400 mg daily for 5 consecutive days

Children < 2 years

- Give Albendazole 200 mg daily for 3 consecutive days
- Check stool 3 weeks after treatment, at least 3 specimens
- Repeat above treatment if eggs or larvae still found

15.5.3 Taenia Saginata/Solium (Tapeworm) Infestation

Treatment

Adults

- Give **Praziquantel** 10 mg/kg stat
- Children < 2 years
 - Give **Praziquantel** 10 mg/kg stat

16.0 Respiratory Conditions

16.1 Acute Respiratory Infections (ARI) in Children

- Most ARI are mild, self-limiting viral infections
- The Malawi ARI Control Programme emphasizes standard case management as its main strategy. This includes:
 - > Early diagnosis
 - Appropriate drug use
 - > Timely referral
 - Advice on suitable home care
- Refer to ARI Control Programme Guidelines,
 MOHP 1998 for more information
- Refer to the WHO's Management of the Child with Cough or Difficult Breathing for a summary of patient assessment, classification of illness and treatment instructions

Note ARI Case Management

- Refer all cases for severe disease/pneumonia to hospital for admission after initial IM doses of recommended antibiotics
- Treat all pneumonia cases as out-patients with Cotrimoxazole or Amoxycilin
- Do not use cough mixtures they have no role to play in ARI management

16.1.1 Home care of children with ARI

16.1.1.1 Home care of child with ARI (2 months – 5 years)

Advise guardian to:

- Watch out for these danger signs (which may indicate pneumonia) and return quickly to the health facility if any occur:
 - Breathing becomes difficult
 - Breathing becomes fast
 - Child cannot drink
 - Child becomes more ill
- Feed the child
 - Continue feeding the child during illness
 - Increase feeding after illness
 - Clear blocked nose if interfering with feeding
- Increase fluids
 - If > 6 months old, offer the child extra fluids to drink

- Increase breast-feeding
- Soothe throat and relieve cough
 - Give sips of water or other (preferably warm) fluids
- Treat fever
 - Give Paracetamol in the recommended dose every 6 hours until the high fever stops
 - Increase fluids (see above)
 - Do not overdress or overwrap the child, i.e. keep the child lightly dressed
- Complete prescribed treatment
 - Complete this even if the child becomes better
- Return for follow-up assessment after 2 days if child is being treated for pneumonia.

16.1.1.2 Home care of child with ARI (young infant)

Advise guardian to:

- Watch out for these danger signs and return quickly to the health facility if any occur
 - Breathing becomes difficult
 - Breathing becomes fast
 - Young infant not able to feed properly
 - > Young infant becomes more ill
- Keep the young infant warm
- Breast-feed often
- Clear blocked nose if interfering with feeding

16.1.2 Common cold (nasopharyngitis)

- Coughs are commonly associated with colds
- Antibiotics should not be given
- Often causes fever in young children which may last up to 72 hours
- In infants, nasal discharge may interfere with breast-feeding and cause difficult in breathing
- Rule out pneumonia, otitis media, and streptococcal pharyngitis
 - Advise mother on how to correctly provide suitable home care (see section 16.1)

16.1.3 Sinusitis

- Most sinusitis is viral and self-limiting, requiring no antibiotics
- Pain in sinusitis is NOT an indicator of severity
- Steam inhalation may help drainage of blocked sinus
- Purulent nasal discharge may be caused by a foreign body in the nose
- Bacterial sinusitis is usually caused by S. pneumoniae or H. influenzae. It is characterized by:
 - Persistent purulent nasal discharge >7 days plus

- Sinus tenderness and/or
- Facial or periobital swelling and/or
- Persistent fever
- Extract tooth under antibiotic cover
 Benzathine Penicillin 1.2g IM stat or
 Amoxycillin 3g orally stat 1hour prior to procedure and Metronidazole 400mg
- Only if there are definite signs of bacterial sinusitis give Amoxycillin 500mg tds for 7 days

Alternatively if penicillin hypersensitivity:

- Give **Erythromycin** 500 mg qid for 7 days *If there is pain or fever give*:
 - Analgesic/antipyretic treatment as required

16.1.4 Pharyngitis, Tonsillitis and its Complications

- Most sore throats are due to viral infections such as adenovirus and CMV, and should not be treated with antibiotics
- For pain or fever give analgesic treatment as required
- Be sure to rule out streptococcal pharyngitis to prevent acute rheumatic fever and other non-suppurative (endocarditis) and suppurative complications (retropharyngeal and peritonsillar abscesses).

 Signs suggestive of bacterial pharingitis are abrupt onset of pain, fever, tender, enlarged cervical lymph nodes, white or greyish pharyngeal exudates, absence of lower respiratory tract signs and symptoms absence of signs suggesting viral nasopharyngitis (e.g. rhinorrhoea, conjunctivitis, cough)

Note: Do not use cotrimoxazole as it is not effective

Treatment

Adults:

Give Benzathine Penicillin 1.2 MU single dose
 IM

Alternatively (if assured of compliance)

- Give **Amoxycillin** 500mg tds for 7 days *Alternatively in penicillin hypersensitive patients:*
 - Give Erythromycin 500mg qid for 7 days

Children > 30 kg:

Give Benzathine Penicillin 1.2 MU single dose
 IM

Alternatively (if assured of compliance)

- Give **Amoxycillin 250mg** tds for 7 days *Alternatively in penicillin hypersensitive patients:*
- Give **Erythromycin** 500 mg bd *Children < 30 kg*
 - Give Benzathine Penicillin 600,000 IU single dose IM

Alternatively (if assured of compliance)

Give Amoxycillin 125mg-250mg tds for 7 days

Alternatively in penicillin hypersensitive patients:

• Give **Erythromycin** 7.5-12.5 mg/kg orally qid **Note:** If there is pain or fever give analgesic treatment as required

16.1.5 Peritonsillar and Retropharyngeal Abscesses

Peritonsillar abscess (quinsy) and retropharyngeal abscess

Signs and Symptoms

- Fever
- May cause difficulty in swallowing
- Tenderness at the angle of the jaw
- Features of upper airway obstruction such as difficulties in breathing, stridor
- Patient is unable to drink at all

Treatment

Peritonsillar Abscess

Adults:

- Give Amoxycillin 500mg tds and
- Give Metronidazole 400mg tds

Alternatively

- ➤ Give **Benzylpenicillin** 2 MU IV q6h
- Switch when possible (usually after 48-72 hours) to oral Amoxycillin 500mg tds
- Continue for a total of 14 days antibiotic treatment

Alternatively if penicillin hypersensitivity:

Give Erythromycin 500mg qid

Children:

- Give Benzylpenicillin 25,000 units/kg/dose
- Switch when possible (usually after 48-72 hours) to oral Amoxycillin 15mg/kg tds.
- Continue for a total of 14 days antibiotic treatment

Alternatively if penicillin hypersensitivity:

Give Erythromycin 12.5 mg/kg/dose

Further treatment

- Give analgesic/antipyretic for pain and fever
- Refer to hospital for the following
 - ✓ If pus is present and does not drain spontaneously then carry out incision and drainage
 - ✓ If quinsy is present carry out needle aspiration for analgesic and therapeutic effect

• Retropharyngeal Abscess

Signs and symptoms

- See above
- Surgical drainage is usually necessary, therefore should be referred.

Treatment

Adults:

Give Co-amoxiclav 625mg tds (or Co-amoxiclav 375mg plus Amoxycillin 250mg) for 14 days

Alternatively

- For a total of 14 days antibiotic
- Analgesic for pain and fever

Children:

- Chloramphenicol 25 mg/kg tds, initially IM or IV. later orally for a total of 14 days
- Analgesic/antipyretic for pain and fever

16.1.6 Bronchitis (Acute)

 Cough productive of purulent sputum, not improving after 3 days, without signs of pneumonia

Treatment

- Give **Amoxycillin** 500mg tds for 5 days *or*
- Give Doxycycline 200mg on first day followed by 100 mg od for a further 5 days

16.1.7 Cervical Adenitis

Adenitis may be due to bacterial infection,
 TB, KS and/or HIV infection among other causes

Treatment

- Give **Amoxycillin** 500mg tds for 10 days *Alternatively* if penicillin hypersensitivity:
 - Give **Erythromycin** 500mg qid or
 - Give Doxycycline 200mg on first day followed by 100 mg od for a further 9 days
 - For pain or fever give analgesic

Note: If no improvement, do a fine needle aspiration for AFBs

16.2 Lower Respiratory Tract Infections

16.2.1 Asthma (Recurrent Wheezing)

- Exclude stridor and upper airway obstruction (inspiratory difficulty rather than expiratory wheeze of asthma) before diagnosing and treating for asthma.
- Wheezing can also be a sign of heart failure
- Prevention of attacks is important
- Drugs delivered by metered dose inhaler (MDI) should always be given via a spacer device, especially during acute attacks. This will maximize effective drug delivery and minimize side effects.
- You can make an effective spacer device using a plastic bottle. Cut a small hole in the bottom of a plastic bottle that you can fit the inhaler mouthpiece to with an airtight seal. Shake the inhaler before each puff
- Prime the spacer device with 2 puffs before use

- If a spacer is unavailable, check patient inhaler technique to ensure good delivery of the drug into the lungs, not the throat.
- Antibiotics are not routinely indicated in asthma. They are indicated if there is good evidence of a precipitating respiratory infection (e.g. fever, bronchial breathing)

16.2.2 Severe Asthma

- Consider other causes of acute severe breathlessness with careful examination: acute left ventricular failure, pneumothorax, pulmonary embolus, upper airway obstruction, massive pleural effusion, severe pneumonia.
- Severe or life threatening asthma in adults is suggested by any of the following:
 - Silent chest
 - Central cyanosis
 - ➤ Tachypnoea RR >30, exhaustion, inability to complete sentences
 - Persistent tachycardia >110bpm, bradycardia, hypotension, pulsus paradoxus
 - Use of accessory muscles
 - Confusion, agitation or coma
 - > Peak flow < 33% of predicted
- Arrange immediate hospitalization

Treatment

- Set up IV line and rehydrate with 0.9% Normal
 Saline
- Give High flow oxygen 5//min

- Give Salbutamol Nebuliser solution 5 mg by nebuliser repeated initially as required, then every 6 hours or Salbutamol 4 puffs inhaled via spacer device and
- Give Hydrocortisone 200 mg IV every 8 hours for 3 – 5 days then
 - Change to Prednisolone 40mg orally as a single daily dose in the morning, with food for another 5 -7days
 - If you need to use steroids for more than 10 days, gradually taper the dose for steroids 10mg per week initially, and decrease by 5mg until you stop.
- Give an appropriate antibiotic therapy:
 - Amoxycillin 500mg tds for 5 days or Doxycycline 200 mg on first day followed by 100 mg od for a further 4 days
- Repeat Salbutamol dose via spacer or nebuliser every 15 -30 minutes until improved or reassess after 1hour
 - If no improvement add only if not already taking Theophylline
- Give Aminophylline 250mg slowly over 10 minutes. Beware of acute Aminophylline toxicity including cardiac arrhythmias and seizures.

- If no response:
 - ➢ Give Aminophylline 250 500mg an IV infusion in 1L of 5% Dextrose or 0.9% Sodium Chloride over 12 hours
 - ➢ Give Magnesium Sulphate 1.2 − 2.0g IV over 20 minutes
 - ➤ Give **Adrenaline** 0.5-1.0ml of 1:1000 slowly nebulised or IM
 - You may need to rescue the exhausted patient with assisted ventilation
 - Reassess continuously until the patient settles

16.2.3 Milder Asthma Attacks

General measure

 Carefully teach and monitor correct inhaler use technique and recommend a spacer device

Treatment

Adults:

- Give Salbutamol inhaler 2 puffs via a spacer device repeated as required initially, then every 8 hours
- Keep under observation at least for 24 hours after attack

Alternatively:

Give Salbutamol 2 - 4mg orally tds

Note: Antibiotic therapy may be indicated if signs of infection

Maintenance and preventive treatment of asthma - the stepwise approach

- An environment free from cigarette and wood smoke can reduce attacks
- Check compliance and inhaler technique at each step before progressing
- Step up where required due to frequency of exacerbations
- Step down where possible:
 - > STEP 1: Initial treatment should be with Salbutamol inhaled via a spacer device (see above) as required or tablets if inhalers are unavailable
 - > STEP 2: If required more than once every day add preventive therapy-inhaled steroid eg Beclamethasone 2 puffs (200mcg) twice a day via a spacer. Increasing to 4 puffs twice a day as required
 - > STEP 3: Refer for specialist care if no control with steps 1 & 2

Alternatively (to be used only if the above are NOT available)

- Give Salbutamol tablets 2-4 mg 3-4 times daily.
- Give Aminophylline 100mg twice daily.
- Give Prednisolone 5 mg orally once daily.

Paediatrics

Treatment for Mild Asthma

- Use Salbutamol inhaler via spacer (and facemask if <3 years)
- Puffs 4 hourly for 2-3days

- Discharge with advice:
 - > Inhaler and spacer technique
 - Avoid allergens
 - Return if worse

Treatment for Moderate Asthma

- Use Salbutamol inhaler via spacer (and facemask if <3 years)
- <4yrs 5 puffs every 20min × 3
- ≥4yrs 10 puffs every 20mins × 3
- Give **Prednisolone**:
 - <5yrs 20mg od for 3 days</p>
 - ≥5yrs 30mg od for 3 days
- Clinician to review child after three treatments
 - Admit if no improvement
 - look carefully for severe or lifethreatening features
 - Continue 2-4hourly salbutamol via spacer.

Treatment for Severe or Life-Threatening Asthma

- Give Oxygen
- Salbutamol Nebuliser
 - <4yrs 2.5mg every 20-30mins</p>
 - ≥4yrs 5mg every 20-30mins
- Steroids:
 - Give Prednisolone or IV Hydrocortisone if not taking orally
 - ✓ <5yrs 20mg Prednisolone od or 50mg Hydrocortisone IV single dose
 - ✓ ≥5yrs 30mg Prednisolone od or 100mg Hydrocortisone IV single dose

- Regular reassessment and consult
- 2/3 maintenance IV fluids
- If not improving
 - Consider IV Aminophylline loading dose = 5mg/kg over 20mins then 1mg/kg/hr only with senior input
 - Consider Chest X-Ray to rule out Pneumothorax and Foreign body then consider ITU

Aminophylline - CAUTION

- If the child has been on maintenance
 Theophylline or has been taking Erythromycin,
 the loading dose of Aminophylline should be omitted.
- These children will need very careful monitoring.
 Stop the Aminophylline immediately if the child vomits, has a pulse rate of > 180/min, develops a headache or has a convulsion.
- Supportive Care
 - Ensure that the child receives daily maintenance fluids appropriate for his/her age. This can be orally or IV.
 - Encourage continued breast feeding.
- On Discharge
 - Prescribe Salbutamol via spacer 2-4puffs 4 hourly for 2-3 days, then as required.
 - Advise completion of 3 days of Prednisolone
 - Teach inhaler and spacer technique
 - Avoid allergens

Chronic / Ongoing Treatment of Asthma Medication

- All children (except mildest wheeze) need a salbutamol inhaler on discharge for when the child is acutely breathless or wheezy. 2 puffs (200mcg) up to 4 times daily.
- If >1 asthma attack consider daily

Beclomethasone

- > years 1 puff (100mcg) twice a day
- 2-12 years 2 puffs (200mcg) twice a day
- ➤ 12-18
- years 2-4puffs (200-400mcg) twice a day.

Note: 'Interval symptoms' (cough/wheeze at night or after exercise, prolonges recovery from URTIs) are guides to poorly controlled asthma.

Guardian Education

- How and when to take inhalers (this will need to be observed)
- How to use a spacer
- When to seek help (eg breathlessness not controlled byinhalers, sudden increase in the need for 'relievers')
- Possible precipitating factors
- Follow up monthly

Note:

Use for the shortest time possible before reverting to preferred Agents

- Where asthma is mainly a problem at night, Salbutamol before bed may be sufficient
- If asthma is brought on by exercise, adults can take 2 puffs of Salbutamol inhaler via a spacer device a few minutes before games or sports
- Exercise induced asthma is usually a sign of poor control. If possible introduce an extra level of medication according to the stepwise approach above.
- If asthma is brought on by exercise, older children and adults can take 2 puffs of **Salbutamol** inhaler via a spacer device 30-60 minutes before games or sports

16.3 Community Acquired Pneumonia (CAP)

 For all forms of pneumonia HIV testing is required.

16.3.1 Mild to Moderate Pneumonia

Usually caused by pneumococcus (sudden onset)

Treatment

- Give **Amoxycillin** 500mg tds for 5-7 days *If Penicillin allergic*
 - Give Erythromicin 500mg qid for 5-7 days
 - Give **Doxycycline** 100mg bd for 5 7 days

Note: If the patient does not improve, consider alternative diagnoses including

16.3.2 Atypical Pneumonia

- Caused by Mycoplasma pneumoniae and Chlamydia pneumoniae
- Suspect in previously healthy young adult not responding to treatment

Treatment

• Give **Erythromycin** 500mg qid for 5 days

16.3.3 Nosocomial Pneumonias

 Caused by Staphylococcus aureus, gram negative rods and Pneumococcus

Treatment

- Give **Co-amoxiclav** 1.2g IV q8h or **Ceftriaxone** 2g IV q24h
- Followed by oral Co-amoxiclav 625mg tds for 7 days

16.3.4 Severe Pneumonia

Signs and Symptoms

- Respiratory rate > 30/min, shock (- BP <90/60mmHg), confusion / drowsiness central cyanosis
 - Refer to district hospital

Treatment

• Give **Ceftriaxone** 2g IV q12h plus

Azithromycin or **Erythromycin** 500mg qid *Alternatively*

• Give Co-amoxiclav 1.2g IV q8h

If aspiration

Add Clindamycin 600mg qid or Metronidazole
 400mg tds 1.2g IV q8h

Erythromycin 500mg qid for 7 days *Or* Give **Doxycycline** 100mg od for 7 days

16.4 Suppurative Lung Disease

16.4.1 Lung Abscess

Give Co-amoxiclav 625mg tds

plus

Metronidazole 400 mg tds

Alternatively

• Give **Erythromycin** 500mg qid

or

• Give Doxycycline 100mg bd

Note: Continue treatment for 21 to 28 days

16.4.2 Empyema

- Carry out surgical drainage
- Continue antibiotic therapy as for CAP for 21 to 28 days
- Rule out TB

16.5 Bronchiectasis

Signs and Symptoms

 Chronic cough and sputum production years after treatment of pulmonary TB

Treatment

- Physiotherapy to aid postural drainage of secretions every morning
- Give prompt antibiotic therapy when secondary infection occurs
- Give Co-amoxiclav 625mg tds

Alternatively

- Give Doxycycline 200mg stat then Doxycycline 100mg od
- If sputum is foul smelling, add Metronidazole 400mg tds

Note: Treat for 14 days

16.6 Chronic Lung Disease / COPD

- Acute exacerbations may present in a similar way to asthma, however the patients will be older, and will often have smoked or been exposed to wood smoke, e.g. cooking at home.
- Not always related to an infection.

Treatment

- Give Salbutamol as required.
- Always use a spacer device as described in the section on asthma above.
- Give **Prednisolone** 40mg od for 10 days *If there is fever and purulent sputum then*
 - Give **Amoxycillin** 500mg tds for 7 days

Notes:

 If a patient has more than 4 exacerbations in a year a trial of regular inhaled
 Salbutamol via a spacer 2 puffs four times daily, and as required in between can be tried Long term use of oral steroids is NOT recommended.

16.7 Pulmonary Embolism

- Treat with 6 months Warfarin therapy if available, but only if INR monitoring and specialist supervision are available.
- Long term treatment may be necessary in certain circumstances, including recurrent cases.
- Look for a triggering cause

17.0 Sexually Transmitted Infections (STIs)

- Refer to the Management of Sexually Transmitted Infections using Syndromic Management Approach, Guidelines for Service Providers, MoH, 3rd edition May 2008 for more detailed information.
- All patients who present with STI symptoms should be offered HIV Testing and Counselling.

Note: Prompt and effective treatment of STIs helps prevent spread of HIV infection

General Management

- Ensure adequate privacy in patient management.
- Establish a correct diagnosis whenever possible.
- Make efforts to trace, treat and counsel all sexual contacts.
- Provide health education and counseling on each return visit.
- Advice on 'safer sex' practices to prevent reinfection, i.e. abstinence, correct use and
 storage of condoms, mutual faithfulness of
 uninfected partners, decrease in number of
 sexual partners, use of non-penetrative sexual
 techniques and the importance of partner
 notification and treatment.
- Offer a supply of condoms at each patient's visit.

Note: Periodically check the patient's understanding of the above issues by asking him/her to repeat the information given

17.1 Syndromic management of STIs

- The syndromic approach is based on the fact that most common causes of an STI infection generally present with certain groups of signs and symptoms (syndrome) and treatment given is supposed to target the commonest possible causes of that syndrome.
- Common STI syndromes:
 - Genital ulcer disease (GUD)
 - Urethral discharge (UD)
 - Genito-urinary symptoms in women (GUS)
 - Lower abdominal pain (women)(LAP)
 - Acute scrotal swelling
 - Enlarged inguinal lymph nodes (bubo)
 - Balanitis/balanopostitis

17.1.1 Genital ulcer disease (GUD)

Common Causes

- Genital herpes, Chancroid and Syphilis may be present concurrently.
 - Genital herpes is the most prevalent amongst the three.
- Treat patients with GUD for the above three infections

General Management

- Aspirate fluctuant lymph nodes (buboes) through adjacement normal (i.e. not inflamed) skin.
- Do not incise.
- Ask patients to return if non-fluctuant nodes become fluctuant

Treatment

- Give Ciprofloxacin 500mg orally stat and
- Give **Benzathine penicillin** 2.4 MU IM stat
- Give Acyclovir 800mg bd for 7 days
- Tell patient to return for follow-up care in 7-10 days, see below

Note: Acyclovir is indicated only in symptomatic GUD clients

If patient allergic to penicillin/Ciprofloxacin and pregnant or lactating:

- Give **Erythromycin** 500mg qid for 15 days plus
- Give Acyclovir 800mg bd for 7 days

Infants born to mothers treated for GUD with Erythromycin alone:

 Give Benzathine Penicillin 500,000 IU/kg as a single dose

Follow-up care of GUD

- Inform the patient to return 7-10 days after starting treatment.
- If the ulcers have not healed or are getting worse, repeat GUD treatment if there is evidence of noncompliance.
- If the client complied fully and there is no improvement:
 - ➤ Give **Azithromycin** 2g stat.
 - > Review in further 7-10 days
 - If no improvement, refer for specialist opinion
 - If improved, follow patient's progress until completely healed
 - No further antibiotics are required at this time
- If the ulcers have improved but not completely healed:
 - Repeat chancroid treatmentCiprofloxacin 500mg single dose
 - Review in further 7-10 days

- Subsequent action as above
- If the ulcers have completely healed:
 - Reinforce counseling and patient education
 - Promote/provide condoms

17.1.2 Urethral Discharge/Urethritis in Men (UD)

Signs and Symptoms

Discharge or dysuria

Common causes

- Neisseria gonorrhoea, Chlamydia trachomatis and trichomonas vaginalis.
 - Common in males

Note: If there is dysuria but no discharge and no sexual contact in the last 2 weeks, seek other causes, like urinary tract infection, prostitis or schistosomiasis

Treatment

- Give **Gentamycin** 240mg IM stat plus
- Give Doxycycline 100mg bd with food for 7 days
- Give **Metronidazole** 2g stat

Alternative to Doxycline in pregnancy/lactation:

- Give Erythromycin 500mg qid for 7 days
- Review after 7 days
- If symptoms persist or recur:
 - Rule out re-infection
 - Assess compliance
 - Retreat as above if noncompliant or there is evidence of reinfection

- > Review after further 7 days
 - ✓ If symptoms still persist, refer for further investigation

17.1.3 Abnormal Vaginal Discharge in Women (AVD)

Causes

- Vaginal infection, cervical infection, endometrial infection/pelvic inflammatory disease (PID)
 - > Common causes of vaginal infections
 - Trichomonas vaginalis, candida albicans and bacterial vaginosis.
 - Common Causes of cervical infections
 - ✓ Neisseria gonorrhoeae and chlamydia trachomatis.

Note: Vaginal discharge is normal during and after sexual activity; at various points through-out the menstrual period; and during pregnancy and lactation.

General Management

- Do risk assessment to identify women at risk of cervical infection
 - treat for vaginitis to those with negative risk assessment
 - > treat for cervicitis and vaginal infection to those with positive risk assessment.

- Treat all women with vaginal discharge and a positive risk assessment for gonococcus and Chlamydia infection, plus trichomoniasis and bacterial vaginosis.
 - If the discharge is white and curd-like also treat for *candidiasis*.
- Treat all women with vaginal discharge and a negative risk assessment for trichomoniasis and bacterial vaginosis
 - ➤ If the discharge is white and curd-like, also treat for *candidiasis*.

- If vaginal discharge is present and the risk assessment is positive:
 - ➤ Give **Gentamycin** 240mg IM stat plus
 - Give Doxycycline 100mg bd for 7 days, plus
 - ➤ Give **Metronidazole** 2g orally single dose
- If the discharge is white or curd-like add 1
 Clotrimazole Pessary 500mg inserted intravaginally stat
- If vaginal discharge is present and risk assessment is negative:
 - Give Metronidazole 2g orally single dose stat
- If the discharge is white or curd-like add 1
 Clotrimazole Pessary 500mg inserted intravaginally stat
- If no discharge is found and risk assessment is positive:
 - ➤ Give **Gentamycin** 240mg IM stat plus
 - Give Doxycycline 100mg bd for 7 days

- If no discharge is found and risk assessment is negative:
 - Reassure client, counsel, educate and provide condoms.
 - Advise client to come back if symptoms persist.
 - Offer HIV testing after providing information and counselling

Note: Examination of GUS in women should never be omitted only for convenience of the health worker

17.1.4 Lower abdominal pain in women (LAP Syndrome)

 A serious condition which should be considered in every woman with lower abdominal pain.

Notes:

- Not every woman with lower abdominal pain has PID.
- Be sure to include any conditions which require immediate surgical or gynaecological treatment

Signs and symptoms

 Fever, lower abdominal pain, pain on discharge, vaginal discharge, cervical tenderness, and often adnexal tenderness or masses on bimanual examination. Signs and symptoms of acute illness requiring immediate gynaelogical/surgical attention:

 Missed or overdue or delayed period; recent abortion, delivery or miscarriage; metrorrhagia; abdominal guarding or rebound tenderness; Active vaginal bleeding.

General Management

- If the patient has a missed/overdue period or abnormal vaginal bleeding:
 - Consider referral
 - ✓ When referring, ensure patient's general condition is stable
- If the patient is very ill, bleeding heavily or in shock:
 - Set up an iv drip and commence resuscitation measures
- If the patient does not have missed/overdue period or abnormal vaginal bleeding but does have any of the following:
 - Recent delivery; Recent/suspected abortion; Rebound tenderness; Abdominal guarding
 - ✓ Give first dose of treatment for PID.
 - ✓ Refer immediately for hospital admission after resuscitating the patient should this be required.

✓ Admit if the patient: is obviously sick; is pregnant; vomits oral medication or if adequate follow-up care cannot be provided.

- If the patient does not have missed/overdue period or abnormal vaginal bleeding and does not have any of the signs/symptoms listed on page 404 but does have cervical excitation tenderness or fever:
 - Give Gentamycin 240mg IM stat,
 - Give Doxcycline 100mg bd and
 - ➤ Give **Metronidazole** 400mg bd for 10 days.
 - Remove any IUCD if any and offer other means of contraception
 - Treat partner for gonococcal and chlamydial infection
 - > Review patient after 72 hours:
- *If improved,* complete 10-day course of treatment for PID
- If not improved, refer for gynaecological or surgical consultation

- If the patient does not have missed/overdue period or abnormal vaginal bleeding and does not have any of the signs/symptoms listed on page 404, and not have cervical excitation tenderness or fever:
 - Determine whether the patient has any other genitourinary complaint/syndrome and manage as per appropriate syndrome:
 - Ask her to return if the abdominal pain persists

17.1.4.1 Lower Abdominal Pain in Women (PID): In-patient treatment

Signs or symptoms of acute PID requiring admission

- Failure to respond to syndromic treatment regime within 72 hours
- Presence of tender pelvic mass which may be an abscess or an ectopic pregnancy
- History or suspicion of recent induced abortion, delivery or miscarriage
- Active vaginal bleeding
- Missed, overdue or delayed period
- Pregnancy
- Metrorrhagia
- Vomiting of oral medication.

Note: The patient should be admitted if follow-up care cannot be guaranteed

Treatment

- If toxic:
 - > IV fluids
 - Parenteral antibiotics.
 - ✓ Gentamycin 1.5 mg/kg slow IV or IM q8h plus
 - ✓ Chloramphenicol 500mg IV q6h
 - ✓ Metronidazole 500mg IV q8h
- When improved and able to swallow:
 - add Doxycycline 100mg every 12 hours and
 - switch from parenteral to oral
 Metronidazole 400mg every 12 hours for 10 days
 - Analgesic
- If pain is severe:
 - Give Pethidine 100 mg IM or orally
 - Repeat every 3-6 hours, as required

Notes:

- Acute PID may be due to puerperal or post-abortion sepsis.
- Admit treat with parenteral antibiotic therapy.
- **Evacuate the uterus** within 12 hours of antibiotic therapy regardless of the patient's temperature
- Provide supportive care such as blood transfusion, iv fluids and
- Closely monitor vital signs.

17.1.5 Acute scrotal swelling or pain

General Management

- Distinguish from scrotal swelling/pain due to:
 - Other long-standing causes, e.g. Scrotal hydrocele, varicocele, inguinal hernia
 - Recent/acute illness, e.g. Testicular torsion or trauma, inguinal hernia

Note: Thorough history and physical examination are necessary to exclude potentially lifethreatening conditions and to determine whether immediate surgical attention is required

- Even if the presumptive diagnosis is an STI, treat all patients and partners for gonorrhoea and chlamydia infection:
- Give Gentamycin 240mg IM stat and
- Give **Doxcycline** 100mg bd for 7 days.
- In pregnant or lactating mothers
 Doxcycline should be replaced with
 Erythromycin 500mg qid for 7days.
 - Additional therapy for the patient:
 - bed rest with the scrotum elevated
 - > cold compresses to help reduce swelling
- If there is no evidence of painful and/or swollen scrotum, look for signs of another STI and if present treat appropriately

17.1.6 Enlarged inguinal nodes (bubo)

- Both chancroid and lymphogranuloma venereum (LGV) can cause bubo.
- Exclude the following conditions which may also cause enlarged inguinal lymph nodes: septic skin lesions on thigh, foot, leg, toes, buttock, anus, perineum, scrotum, penis, labia, vulva and vagina, systemic infections e.g. Hepatitis B. HIV infectious, mononucleosis, syphilis, TB, other infections e.g. bubonic plague, cat scratch fever, trypanosomiasis, lymphoma, leukemia, Kaposi's sarcoma.
- Exclude other conditions which may cause groin swelling unrelated to enlarged lymph nodes including: inguinal hernia, lipoma, a boil in overlying skin.
- Confirm presence of bubo by careful examination

Note: All patients with bubo should be carefully examined for signs of other STIs

- If bubo present and genital ulcer present: treat as for genital ulcer disease syndrome
- If bubo present, and painful, fluctuant or recent onset (under 2 weeks) and no genital ulcer present: treat patient and partner for LGV

Give Doxycycline 100mg bd with food for 14 days

Alternatively in pregnancy/lactation

- Give **Erythromycin** 500mg qid for 14 days
- If bubo fluctuant, aspirate through adjacent normal skin (do not incise)
- If enlarged inguinal lymph node present, but not painful, fluctuant or of recent onset (under 2 weeks) and no genital ulcer present: look for other causes of inguinal swelling:
 - e.g. generalized lymphadenopathy (rule out secondary syphills and HIV), hernia, tumour.
- Refer for biopsy if indicated
- If bubo not present but other signs of STI found, treat accordingly
- If bubo not present and other signs of STI not found, reassure, educate/counsel the patient
- Promote/provide condoms

17.1.7 Balanitis/Balanopostitis (BA Syndrome)

Cause

- Fungal infection, trichomonal infection and medicine reactions
 - Common and persistent in persons with diabetes and with immunosuppression caused by HIV infection

Note: The most common reason for balanitis is poor genital hygiene

General Management

- Ask patient about any recent topical application of medicines (including traditional medicines)
- Ask patient if partner/s have vaginal itching or discharge – this may indicate candidiasis or trichomoniasis
- Examine the patient carefully for presence of genital ulcers and urethral discharge

- If the foreskin is retractable and ulcer/s present, or if the foreskin is not retractable:
 - Treat as for genital ulcer Disease syndrome.
- If the foreskin is retractable, no ulcer, but urethral discharge present:
 - Treat as for urethral discharge

- If the foreskin is retractable, no ulcer, no urethral discharge, but erythema or erosions present:
 - > Treat for candida infection
 - ✓ Apply Gentian Violet 1% aqueous solution to glans penis daily for 7 days.
 - Treat partners with Clotrimazole pessary 500mg inserted vaginally once.
 - Advise on genital hygiene, including frequent washing of penis with soap and water
 - Review patient in 7-10 days:
- If symptoms persist:
 - Treat patient and partner for trichomoniasis with
 Metronidazole 2g stat.
 - Advise client to avoid alcohol during treatment and for 48 hours after dose
- If symptoms resolved and no signs of STI:
 - > Reassure, educate/counsel
 - Promote/provide condoms
- If foreskin is retractable, no ulcer, no urethral discharge, no erythema or erosions and no signs of STI:
 - Reassure, educate/counsel
 - Promote/provide condoms

17.2 Genital warts

- Distinguished from condyloma of secondary syphills, and molluscum contagioum
- Besides local caustic applications, surgical removal or electrocautery may be used for treatment:
 - For more extensive growth
 - When topical applications have failed
 - When topical application are contra-indicated

Treatment

- Apply Compound Podophyllin Paint to the lesions at weekly intervals
- Apply Yellow Soft Paraffin to avoid normal tissue damage
- Use only for scattered growth
- When applied to vulval mucosa or to meatal warts, allow to dry before coming back into contact with normal epithelium
- Remove the paint by washing off after 1-4 hours

Note: Do not use this therapy during pregnancy

 If no effect after 4-6 weeks, stop treatment and consider alternative methods of removal

Alternatively to Podophyllin Paint, and for treating vulvar warts:

• Apply Silver Nitrate Stick (pencil) once daily

17.3 Neonatal Conjuctivitis

- These serious conditions rapidly progress and threaten sight
- Admit patient to hospital
- Closely monitor until the infection has resolved
- Give antimicrobial therapy without delay
- Irrigate the eyes frequently with normal saline

Note: At birth give all neonates a single prophylactic application of tetracycline eye ointment

Treatment

Adults and all parents of infected babies:

- **Gentamycin** 240 mg IM single dose *Infants with signs of conjunctivitis:*
 - Isolate immediately
 - Institute a rigorous system of barrier nursing with careful attention to hygiene
 - Give **Gentamycin** 5mg/kg IM once (7.5mg /kg if the infant is older than 7 days) and **Erythromycin** 12.5mg/kg qid for 14 days.

Alternatively for Gentamycin,

- Give Cefotaxime 50mg/kg IM as a single dose (maximum 125mg)
- Give Tetracycline eye ointment 1% qid for 3 days applied in each eye every 6 hours for 3 days
 - Clean away any discharge before application

- Wash eyes with clean water/saline ideally every 2 hours until the purulent discharge is cleared
- Treat Father with
 - ➤ **Gentamycin** 240mg IM stat, and
 - Doxycycline 100mg bd for 7 days
- Treat Mother with:
 - Gentamycin 240mg IM stat, and
- ➤ **Erythromycin** 500mg qid for 7 days *Alternative topical agent:*
 - ➤ **Gentamycin eye drops 0.3%**, 1.2 drops into each eye every 2 hours
- Reduce dose frequency as the infection is controlled
- Continue for 48 hours after healing

17.4 Syphillis

- Use this regime for patients with syphilis confirmed by laboratory testing
- In treatment of secondary syphilis, a
 Herxheimer reaction (malaise, fever,
 headache, rigors) may sometimes occur
 within 6-12 hours of initial treatment

Treatment

Treat this with Aspirin 600mg every 6 hours

17.4.1 All syphilis except neurosyphilis

Includes:

- Early syphilis: primary (ulcer), secondary (generalized skin rashes, condylomata lata) or latent syphilis of not more than 2 years duration
- Late syphilis: benign, cardiovascular and latent syphilis of more than 2 years; syphilis of indeterminate duration
- ➤ congenital syphilis in children

 Note: Treat as late syphilis all patients with a positive RPR or VDRL and no documented syphilis serology in the last 2 years.

17.4.1.1 Early syphilis in adults

Treatment

- Benzathine Penicillin one dose of 2.4 MU IM
 - Divide as 1.2 MU into each buttock

Alternatively, if hypersensitivity to penicillin:

• **Doxycycline** 100mg bd for 15 days

Note: In pregnancy/lactation, substitute **Doxycycline** with **Erythromycin** 500mg qid for 15 days

17.4.1.2 Late syphilis in adults

Treatment

- Benzathine Penicillin 3 doses of 2.4 MU IM at weekly intervals
 - Divide each weekly dose 1.2 MU into each buttock: total (3 doses) is 7.2 MU

Alternatively, if hypersensitivity to penicillin:

Doxycline 100mg orally every 12 hours for 30 days

Note: In pregnancy/lactation, substitute Doxycycline with Erythromycin 500 mg every 6 hours for 30 days

Notes for pregnant patients

- Any history of penicillin hypersensitivity must be reliable as these patients are put at serious disadvantage because they cannot be given tetracyclines
- The child must be treated for congenital syphilis at birth as Erythromycin does not readily cross the placenta

17.4.2 Congenital syphilis in children

 Treat with a single dose of Benzathine penicillin 50 000 IU/kg IM in all infants born to sero-positive mothers whether or not the mothers were treated during pregnancy (with or without penicillin) unless they have features of congenital syphilis.

- Thoroughly examine for congenital syphilis all infants born to women with reactive serologic tests: look for ascites, oedema, jaundice, hepatosplenomegaly, rhinitis, nasal discharge, hoarse cry, skin rash, and/or pseudoparalysis of any extremity.
- Treat infants with these symptoms as early congenital syphilis:

Treatment

- Give Benzylpenicillin 50 000 IU/kg/dose IV q12h, during the first 7 days of life and every 8 hours thereafter for a total of 10 days,
- Children with late congenital syphilis (more than 2 years) are treated as follows:
 - Give Benzylpenicillin 50 000 IU/kg/dose IV every 4 to 6 hours 10 to 14 days,

Alternatively, in penicillin allergic children

 Give Erythromycin syrup 12.5mg 6 hourly for 30 days.

Note: The risk of penicillin hypersensitivity in the 1st month of life can be safely discounted

17.4.3 Neurosyphilis

 Higher penicillin doses are necessary to ensure that levels in the CSF do not fall below required amount throughout the course of treatment

Treatment

Adults:

- Give Benzylpenicillin 4MU IV q6h for 14 days then
- Give Benzathine Penicillin 2.4 MU IM once weekly for 3 consecutive weeks

Alternatively if confirmed hypersensitivity to penicillin:

Doxycycline 200mg every 12 hours for 30 days
 Note: In pregnancy: substitute Doxycycline with
 Erythromycin 500mg every 6 hours for 30 days

17.5 Trichomoniasis, vaginal

- Coincident bacterial vaginosis reduces the effectiveness of single dose Metronidazole treatment
- Asymptomatic male partners should also be treated

Treatment

Adults:

 See AVD Syndrome
 Infants with symptomatic trichomoniasis or urogenital colonization persisting after the 4th month

 Give Metronidazole 5 mg/kg every 8 hours for 5 days

18.0 Skin conditions

18.1 Bacterial skin infections

18.1.1 Impetigo

Signs and Symptoms

- Pruritus
- Crusts

Investigation

Pus Swab

Causes

 Staphylococcal aureus and Streptococcal pyogenes

General management

- Keep infected areas clean- wash daily with soap and water
- For children, instruct the mother
- Prevent spread to others take care with towels and clothes and change/clean bedding frequently
- Remove crusts with warm water

Treatment

Apply Gentian Violet paint 0.5 % on wet lesions

Alternatively

Adults

 Give 5% Salicylic acid with 5% Sulphur ointment

Children

 Give 2% Salicylic acid with 5% Sulphur ointment

- If extensive or involving hairy areas: Give
 Flucloxacillin 125-500mg qid for 5-7 days or
- Give **Erythromycin** 125-500mg qid for 5-7 days *If no response, refer to next level of care*

18.1.2 Folliculitis

Bacterial infection of the mouth of the hair follicle with staphylococcus aureus

Symptoms

• Pain on affected area, pruritis

Signs

Swelling on affected area (pustules)

Investigations

Biopsy

18.1.3 Furunculosis and Carbuncles

Signs and Symptoms

- Pain
- Swelling of affected area

Investigation

Pus Swab (biopsy at central hospital level)
 Red Flags

 If affected area is wide (spread out) then refer to next level of care for Incision and drainage (I & D)

Treatment

- Clean the affected area with water and soap
- Do incision and drainage if fluctuant
- Give systemic antibiotics
 - ➤ Give Flucloxacillin 125-500mg qid for 5-7 days *or*
 - Give Erythromycin 125-500mg qid for 5-7 days

18.1.4 Ecthyma

Signs and Symptoms

- Pain on affected area
- Formation of crust and ulceration

Investigations

- Pus swab and biopsy
- An ulcerative streptococcal pyogenes skin infection which can easily be confused with impetigo
- Only recognized upon removal of a scab where a punched out ulcer may be seen

General Management

- Remove crusts with warm water
- Give systemic antibiotics (Erythromycin or Flucloxacillin and alternative treatment is Amoxicillin) for 14 – 28 days

18.1.5 Neonatal Pustulosis/ Bullae

Signs and Symptoms

- Pain on affected area, pruritis
- Pustules on affected area

Investigations

 Check Venereal Disease Research Laboratory (VDRL) status

Causes

 Staphylococcal aureus or Streptococcal pyogenes, syphilis (Treponema pallidum)

Treatment

- Give Potassium permanganate baths twice a day for two weeks
- Give Benzylpenicillin 50,000 units/kg IM or IV q12h for 5 days plus Gentamycin
 2.5mg/kg IM or IV q12h for 5 days
- Check the mother's VDRL or TPHA if positive, then treat

Red Flag

• If neonate shows signs of severe illness (e.g. not feeding)

18.1.6 Erysipelas

Signs and Symptoms

- Pain on affected area
- Well demarcated (defined), erythematous superficial skin lesion with some blisters

Causes

Commonly caused by a beta- haemolytic group-A streptococci

Investigations

Biopsy performed at central hospital level

Treatment

- Give Benzyl Penicillin 2MU IM q6h
- When temperature drops or when condition improves change to Erythromycin 500mg qid for 5-7days
- Apply GV paint bd

18.1.7 Cellulitis

Signs and Symptoms

- Pain on affected areas
- Poorly defined erythematous lesion
- This is bacterial infection of the deeper part of the dermis and the upper part of the subcutaneous tissue

Investigations

Biopsy performed at central hospital level

Treatment

 Give Flucloxacillin 125 -500mg qid for 7 – 10 days

If penicillin hypersensitive

Give Erythromycin 500mg qid for 7 -10 days

If the patient is systemically unwell

- Give Benzyl Penicillin 2 MU IM qid for 7 –
 10 days or oral antibiotics
- Elevate the leg(s) on a pillow to reduce swelling

18.1.8 Staphylococcal Scalding Skin Syndrome – Lyell's disease

Signs and Symptoms

- Pain on affected area
- A febrile, rapidly evolving generalized and blistering desquamative skin condition in which the skin exfoliates in sheets
- Commonly affects neonates and young children, rare in adults unless one is immune compromised

Investigations

- Biopsy performed at central hospital level General Supportive Measures
 - Replace fluid
 - Give **Flucloxacillin** *or* **Cloxacillin** 250 500 mg qid for 7days

Note: Cortico-steroids are contraindicated

18.2 Erythema Multiforme/Steven Johnson's Syndrome

18.2.1 Erythema multiforme – Minor (Iris Type)

Signs and Symptoms

- Usually asymptomatic
- Prodromal symptoms are generally absent and there is relative sparing of the mucus membrane and the trunk
- Target lesions (Iris) are characterized by dark centre and inner pale ring and an erythematus type outer border

Treatment (Symptomatic)

- Give Potassium Permanganate baths bd
- Apply Calamine lotion if vesicular origin or popular eruptions
- Identify the cause and treat. If the cause is medicine, stop the medicine
- Give Chlorpheniramine 4mg tds for 7 days

18.2.2 Steven Johnson's Syndrome/Vesicobullous type

- Potentially fatal condition (to be managed at central hospital level)
- Often caused by hypersensitivity to medication
- Always ask for use of antiretroviral therapy (nevirapine), sulphur drugs, and penicillines.
 These need to be stopped immediately

Signs and Symptoms

- Pain and burning sensation on affected area
- Peeling skin and ulceration of mucous membrane

Treatment (Symptomatic)

- Give Normal Saline
- Add Dextrose 50%
- Identify the cause and treat. If the cause is medicine, stop the medicine
- Keep patient warm
- Give Potassium Permanganate baths bd
- Apply **GV Paint** bd on fresh lesions
- Give Acyclovir 800mg tds for 5 days in adults if of viral origin
- Give **Ceftriaxone** 2g IV q24h for 5 days
- High protein diet is recommended

18.3 Toxic Epidermal Necrolysis (TEN)

- Often caused by hypersensitivity to medication but unlike in Steven Johnsons Syndrome, the mucous membrane is not affected
- Always ask for use of antiretroviral therapy (nevirapine), sulphur drugs, and penicillines.
 These need to be stopped
- Refer to the hospital

Treatment

- Give Prednisolone (step-wise treatment)
 80mg od for 1 week; 70mg od for 3 days;
 60mg od for 3 days; 50mg od for 3 days;
 40mg od for days; 30mg od for days; 20mg
 od for days; 10mg od for days; 5mg od for 3
 days
- Give Normal Saline
- Identify the cause and treat
 - Give Ceftriaxone 2g IV once daily for 5 days
 - High protein diet is recommended

18.4 Fixed Medicine Eruption

- Diagnosis is clinical and in relation to history
- Stop the offending medicine

Signs and Symptoms

- Itching and/or pain on affected area
- Erythamotous skin, blistering and hyperpigmentation on affected area

Treatment

Use potent topical steroids e.g.
 Betamethasone 0.05% cream bd

If severe:

 Give Chlorpheniramine 4-8mg bd for 7 -14 days

18.5 Urticaria

It is called angioedema if deep dermis and subcutaneous tissues are involved

Signs and Symptoms

- Extreme pruritus
- Wheals on affected area

Red Flags

If it is angioedema, refer

General Management

- Explain the condition to the patient
- Remove the cause if known

Treatment

- Give antihistamines
 - Figure 4.25 Give Promethazine 25 mg IM or orally at night or every 8 hours or at night for 1 to 2 weeks or till 48 to 72 hours after submission of the wheals.

 Anthistamines can be given for 1-3 months

Alternatively

- Give Chlorpheniramine 4-8mg at night or every 8 hours for 1-2 weeks (maximum of 3 months)
- Give Cetrizine 5-10mg od (if severe) for maximum of 3 months
- Give Albendazole 400mg stat
- Apply Calamine lotion at night or twice daily
- Give Adrenaline 0.5mg stat dose subcutaneously in severe urticarial with tracheal angioedema

18.6 Eczema (dermatitis)

Signs and Symptoms

- An inflammatory itchy skin condition
- Erythamotous skin, oozing, vesicles, bullae, crusts, lichenification

- Determine the type of eczema
- Types: Atopic (seborrhoeic [indogenous]),
 Allergic (irritant contact eczema [exogenous])
- Eczemas are often secondarily infected (impetigo), systemic antibiotic treatment should be added if indicated
- In HIV (+) children, extensive seborrhoeic dermatitis may occur
 - Give Potassium Permanganate baths bd
 - And topical steroids e.g. Give Hydrocortisone 1% cream or ointment or Sulphur 5% with emulsifying ointment bd
 - If no response, give second line topical steroids or group 3 e.g.
 Betamethasone 0.05% creams
- If secondary bacterial infection occurs:
 - Treat with systemic antibiotics: Give Flucloxacillin or Erythromycin 500mg qid for 5-7 days in adults

General Management

- Counsel the patient on the condition
- Treatment depends on the texture of the affected skin
- Remove any obvious precipitating factors (allergic or contact eczema) ask about soaps detergent, vaseline, cosmetics, clothings etc.
- Use antihistamine to relieve itching.

Treatment

Adults:

Give Promethazine 25 mg orally at night for 7 days

Children:

- Give Promethazine 1 mg/kg at night for 7 days
 Alternatively
- Give **Chlorpheniramine** 4-16mg at night for 2 weeks
- Use antibiotics for secondary infection Give systemic antibiotic treatment only if lesions are infected or signs of systemic
 - infection are present

Note: First line treatment failures can then be treated with Betamethasone cream/ointment or Crude coal tar 5 % ointment.

18.6.1 Acute eczema

 Sudden eruption with erythema, vesicles and sometimes bullae, often with serous exudates (wet appearance)

General Management

- Wet or oozing lesions dry them first with
 Calamine lotion 2 times a day
- Give **Chlorpheniramine** 4mg at night for 7 days

18.6.2 Subacute eczema

 Lesions take several days to erupt, are red but not wet or may be slightly wet. Few vesicles.

General Management

- Normal or slightly wet lesions- use
 Hydrocortisone 1% cream twice a day
- Give **Chlorpheniramine** 4mg at night for 7 days

18.6.3 Chronic eczema

 Develops after months/years, thickened dry and scaly skin, (lichenification), deep cracks (can bleed) scratch marks, sometimes infected

General Management

- Dry skin lesions
 - > Apply **Hydrocortisone ointment 1%** bd

Alternative

• Give potent steroid e.g. **Betamethasone 0.1%**

18.7 Fungal skin infections

18.7.1 Tinea

- Types: Capitis, corporis, pedis, cruris, unguium.
- Instruct patients on the importance of treatment compliance in order to eradicate the infection

Signs and Symptoms

- Pruritus
- Well-defined lesions and hair loss in tinea capitis
- In tinea corporis and cruris, there are welldefined lesions and scales
- Scaling and ulceration occur inter-digitally in tinea pedis

Treatment

- For wet lesions
 - Dry by soaking or mopping with Potassium Permanganate bd for 2 weeks

Alternatives

- Apply Calamine Lotion or Gentian violet Paint
 - Then, apply Compound Benzoic Acid Ointment 6% and Salicylic Acid 3% or Clotrimazole Cream 1%
 - In chronic or extensive cases and those involving hairy areas:
 - ✓ Add Griseofulvin 500mg orally daily with food, single dose or in 2 divided doses, 4-6 weeks
 - ✓ Children: 10mg/kg/dose

Notes:

- Tinea corporis, cruris, pedis treat for 4 weeks
- Tinea capitis treat for 6 weeks
- Tinea unguium of the fingernails treat for 12 months
- Tinea unguium of the toe nails treat for 2 years

Alternative systemic antifungals

Give Ketoconazole or Fluconazole

Red Flag

- Ketoconazole may cause liver toxicity when used for prolonged periods
- In HIV (+) patients all Tineas may be extensive

18.8 Viral skin infections

18.8.1 Herpes simplex

Types: Type 1 (affects lips), Type 2 (affects genitals but can interchange due to oral sex)

Signs and Symptoms

- Pain and pruritus on affected area
- Grouped vesicles on affected area

- In case of bacterial superinfection
- Use salt mouth wash or Chlorhexidine Solution bd for 7 days
- Give Acyclovir Cream or GV Paint or Silver
 Sulphadiazine Cream Application bd for 1 2weeks

- Give Aspirin 600mg tds or Paracetamol 1g tds for 3 days (avoid Aspirin in children as it may cause Reye's syndrome)
- In severe conditions give Acyclovir 200-400mg
 tds for 5 to 7 days and consider testing for HIV

18.8.2 Varicella (Chicken pox)

Refer to Section 9.8

18.8.3 Herpes zoster (shingles)

- Causes: varicella zoster virus
- A common presentation in HIV(+) patients

Sign and Symptoms

- Painful lesions
- Grouped vesicles, then development of blisters may occur

- Give Acyclovir 800mg 5 times a day for 5-7 days; best within 24 -72hours from onset of the lesions
- Apply Calamine lotion or Acyclovir cream bd on intact lesions until they break or GV paint or topical antibiotic cream bd
- Give analgesics for pain relief
- For ophthalmic herpes zoster treat as above and refer to eye department
- Refer for HIV Testing

18.9 Prurigo/Pruritus

Sign and Symptoms

Itching

Causes

 Iron deficiency, anaemia, lymphoma, leukaemia

Treatment

- Adults and Children (symptomatic treatment):
- Apply Calamine Lotion 2-3 times daily
- Give **Promethazine** 25mg single dose at night

or

- Give Chlorpheniramine 4-8mg at night for 2 weeks
 - Children: 1 mg/kg/ dose
- Give Emolients like Emulsifying plain
 ointment applications or baths twice a
 day may help reduce itching
- Investigate the cause, and treat accordingly

18.10 Scabies

Signs and Symptoms

- Pruritus
- Burrrows and papules on affected area
- Secondary infection is common and may mask the condition.
- Treat the whole family and any other close contacts

Treatment

- Wash the whole body with mild soap and water, preferably at night, and dry up
- Apply Benzyl Benzoate Application 25% to the whole body from the neck down
- Ensure all parts of the skin are covered
- Allow the medication to dry and to remain on the skin for at least 10 hours or over night

Next morning wash off the application with

- soap and water
- Wash all contaminated clothes, beddings and towels and use already washed clothes
- Repeat the above treatment on day 5

 Alternative for benzyl benzoate application
 - Apply Lindane 1% lotion, single dose applied as above

Note:

- Avoid Lindane in children less than 2 years old
- In children under 1year, also treat the face (except the area surrounding the eyes)
 - For children under 5 years, Use 12.5% benzyl benzoate application
 - Prepare this by diluting one part 25% benzyl benzoate application with an equal part of water

In cases of severe or extensive infection, especially with secondary bacterial infection:

- Give a systemic antibiotic see Section 18.1
 - Apply 5% Salicylic acid and sulphur ointment bd

- ✓ Consider checking HIV status *If itching is problematic*:
 - Give reassurance
- ➤ Itching may persist for up to 2-3 weeks *If severe:*
 - Chlorpheniramine 4-8 mg at night for 2-3 weeks

18.11 Tropical ulcer

Signs and Symptoms

Painful ulcers

General Management

Improve nutrition and diet

Treatment

 Clean ulcer with Hydrogen peroxide solution bd for two weeks

Alternatively

- Apply Cetrimide 15% +Chlorhexidine solution 1.5% diluted 1 in 20 or Potassium permanganate soaks 1ml in 10,000mls of water
- Debride the wound if necrotic
- Daily Potassium permanganate soaks or cleaning
- Dress the wound with Zinc paste with sulphur regularly till healed
- Rest with leg elevated
- Do a skin graft if wound is clean and shows granulation

If local infection presents:

- Give Amoxicillin 500mg every qid for 7 days or
- **Erythromycin 500mg** qid for 7 days
 - Children 12.5 mg/kg body weight in 4 divided doses
- If possible, carry out culture and sensitivity testing to determine suitable antibiotic therapy
- Refer the patient to the hospital

18.12 Onchocerciasis

Refer to Section 15.2

18.13 Buruli Ulcer

Sign and Symptoms

- Painless ulcers
- Papules, nodules, plaques, ulcers
- This is an ulcerative skin condition caused by Mycobacterium ulcerans

Treatment

- Depends on stage of condition, but if ulcerated
 - Daily wound dressing with Normal Saline 0.9%
 - Medicine therapy doesn't usually work
 - Surgery

18.14 Leprosy

Refer to Section 9.2

19.0 Vaccinations

• For tetanus toxoid vaccination (TTV) see Section 12.3

Vaccination schedule for children

At birth: **BCG** 0.05ml intradermally

Children > 1 year 0.1 ml

Polio 0: 2 drops orally

(= "zero dose")

6 weeks: **PENTAVALENT** 1: 0.5 ml IM

Polio 1: 2 drops orally

ROTA 1: 1ml oral

PCV 1: 0.5ml IM

10 weeks: **PENTAVALENT** 2:0.5 ml IM

Polio 2: 2 drops orally

ROTA 2: 1ml pral

PCV 2: 0.5ml IM

14 weeks: **PENTAVALENT** 3:0.5 ml IM

Polio 3: 2 drops orally

PCV 3: 0.5ml IM

6 months: Oral Vitamin A

9 months: MEASLES: 0.5 ml deep S/C

12 months: Oral Vitamin A

PENTAVALENT = **DPT**, **Hepatitis B** and **Heamophilus**

influenza type B

PCV = Pneumococcal Conjugate Vaccine

Notes:

- Aim to complete this schedule within the first year of life
- BCG vaccination: give this as early as possible in life, preferably at birth – complications are uncommon. BCG is contraindicated in symptomatic HIV infection
- Measles vaccination: normally give this when a full 9 months of age is reached
- Can give an extra dose which is recommended for groups at high risk of measles death, such as children in refugee camps, HIV- positive infants and during outbreaks of measles
- Pentavalent/polio/ROTA/PCV: the minimum interval between doses is 4 weeks
- Tetanus toxoid vaccination: give a full course of this:
 - To all women see Section 12.3
 - After administration of anti-tetanus serum (ATS) to any previously unimmunised patient
 - ➤ If over 10 years has elapsed since the last booster dose

20.0 Bites, Burns and Wounds

20.1 Animal Bites

- Avoid suturing any kind of bite wound
- Thorough cleansing and debridement of the wound is essential
- The combination of local wound treatment, passive immunization with rabies immunoglobulin (RIG), and vaccination with anti-rabies vaccine is recommended for all severe exposures to rabies (see Table on Section 20.1.2)
- Since prolonged rabies incubation periods are possible, persons who present for evaluation and treatment even months after having been bitten should be treated in the same way as if the contact occurred recently

Thorough prompt local treatment of all bite wounds and scratches which may be contaminated with rabies virus is very important as elimination of the rabies virus at the site of infection by chemical and physical means is the most effective method of protection

 As part of local treatment in all cases of possible exposure, carefully instill rabies immunoglobulin RIG, if available, in the depth of the wound and infiltrate around the wound. See *Table on Section 20.1.2* for dose information Avoid contact with the patient's saliva which is potentially infective. If possible, wear eye protection as patients may spit and infection through the conjunctiva can occur

Treatment

Adults and Children:

- Give Tetanus Toxoid Vaccination (TTV) see
 Section 12.3
- Flush and cleanse (scrub) the wound with cetrimide 15% + chlorhexidine solution 1.5% diluted 1 in 20 with water

Alternatively:

- Wash with Hydrogen peroxide solution (10 vol) or soap or detergent
- Rinse with Normal Saline and dress with a weak lodine Solution or lodine Cream
- Give Anti-Rabies Vaccine, only if necessary according to the recommendations in Section 20.1.2
- Give Co-amoxiclav 625mg tds plus
 Metronidazole 400mg tds
- If possible, capture and observe the animal for 10 days. If the animal is still alive after this time period, it does not have rabies

 Human bites should be managed as animal bites except for the use of anti rabies vaccine

20.1.1 Post- exposure immunization

 Give anti-rabies vaccines to all patients unvaccinated against rabies, together with local wound treatment, and in severe cases, rabies immunoglobulin (see recommendation in Section 20.1.2)

20.1.2 Administration of anti-rabies vaccine

- Use intra-dermal injection regimes for Anti Rabies Vaccine whenever possible
- Give a 0.1 ml dose of Anti Rabies
 Vaccine intradermally in either the forearm or upper arm, on days 0, 3 and 7
- Then give 0.1 ml of **Anti Rabies Vaccine** at one site on days 30 and 90

Alternative intramuscular regime

• Give one 1 ml dose of Anti Rabies vaccine IM on days 0, 3, 7, 14 and 28

Suitable injection sites

- In adults: always inject the anti-rabies vaccine into the deltoid area of the arm
- In young children: the anterolateral area of the thigh may also be used
- Never use the gluteal area for vaccination as it is then much less effective

Recommendations for Anti-Rabies Vaccination

| NATURE OF EXPOSURE | CONDITION | N OF ANIMAL | RECOMMENDED ACTION | | |
|---|--|------------------|--|--|--|
| OF EXPOSORE | At time of exposure | 10 days later | ACTION | | |
| 1. Saliva in contact with skin, but no skin lesion | Healthy | Healthy | Do not vaccinate | | |
| | | Rabid | Do not vaccinate | | |
| | Suspect | Healthy | Do not vaccinate | | |
| | | Rabid | Do not vaccinate | | |
| 2. Saliva in | Healthy | Healthy | Do not vaccinate | | |
| contact with skin that has lesions, minor bites on trunk or proximal limbs | | Rabid | Vaccinate | | |
| | Suspect | Healthy | Vaccinate, but stop course if animal healthy | | |
| | | Rabid | Vaccinate | | |
| | | unknown | Vaccinate | | |
| 3. Saliva in contact with mucosae, serious bites (face, head, fingers, or multiple bites) | Domestic or wild rabid animal or suspect | | Vaccinate, and give antirabies serum | | |
| | Healthy domestic animal Vaccinate, but stop course if animal healthy after 10 days | | | | |

20.1.3 Post-exposure immunization in previously vaccinated patients

- In persons known to have previously received full pre-or post-exposure treatment with rabies vaccine within the last 3 years
- Give one booster dose of 0.1 ml Anti Rabies Vaccine intradermally on days 0 and 3

Alternative intramuscular regime:

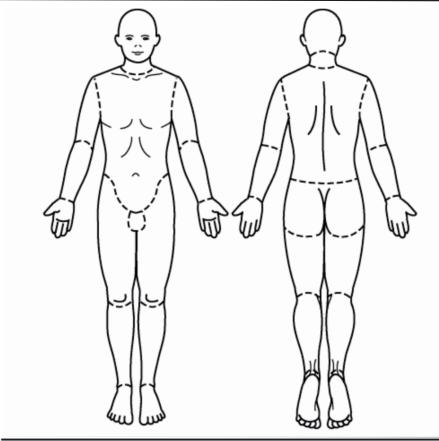
- Give one booster dose of 1 ml Anti Rabies
 Vaccine IM as above
- If completely vaccinated more than 3 years before or if incompletely vaccinated, give a complete post-exposure vaccination course

20.2 Burns

- Remember the ABCs of life support
 - If evidence of inhalation, such as singed nasal hairs, soot in nose, refer to tertiary hospital
 - Pain relief (refer to Palliative Care section)
- Assess the severity of the burn (see table in section 20.2.1)
- Refer patients with burns of more than 15% (children: > 10%) of body surface area (BSA) to hospital on iv fluid therapy for resuscitation and burns dressing

- Refer all deep burns or burns of the face, neck or hands and perineum for further assessment
- Burns across joints should be immobilized and later encourage passive movement to prevent from contractures
- If the burn is more than 40%, mortality is almost 100% therefore referral to a tertiary hospital may not be necessary
- Circumferential burns of the limbs and trunk require immediate bed side escharotomy (see diagram of the site of incisions)
- Burns by nature are usually initially sterile.
 The aim of treatment is to speed healing while minimizing the risk of infection
- In the sick burned patient (fever+diarrhoea+/a rash):
 - Have a high index of suspicion of toxic shock syndrome
 - Naso gastric tube insertion is helpful as gastric dilatation is common
 - ➤ Give anti-acids to prevent gastric stress ulcers (see Section 7.6)
 - Do a blood culture and malarial parasites
 - Start intravenous flucloxacillin

20.2.1 Calculation of Body Surface Area Affected



| | Birth 1 yr. | 1–4 yrs. | 5–9 yrs. | 10–14 yrs. | 15 yrs. | Adult | Burn size estimate |
|-----------------|----------------|-------------|-------------|---------------|------------|-------|--------------------------|
| Head | 19 | 17 | 13 | 11 | 9 | 7 | |
| Neck | 2 | 2 | 2 | 2 | 2 | 2 | |
| Anterior trunk | 13 | 13 | 13 | 13 | 13 | 13 | |
| Posterior trunk | 13 | 13 | 13 | 13 | 13 | 13 | |
| Right buttock | 2.5 | 2.5 | 2.5 | 2.5 | 2.5 | 2.5 | |
| Left buttock | 2.5 | 2.5 | 2.5 | 2.5 | 2.5 | 2.5 | |
| Genitalia | 1 | 1 | 1 | 1 | 1 | 1 | |
| Right upper arm | 4 | 4 | 4 | 4 | 4 | 4 | |
| Left upper arm | 4 | 4 | 4 | 4 | 4 | 4 | |
| Right lower arm | 3 | 3 | 3 | 3 | 3 | 3 | |
| Left lower arm | 3 | 3 | 3 | 3 | 3 | 3 | |
| Right hand | 2.5 | 2.5 | 2.5 | 2.5 | 2.5 | 2.5 | |
| Left hand | 2.5 | 2.5 | 2.5 | 2.5 | 2.5 | 2.5 | |
| Right thigh | 5.5 | 6.5 | 8 | 8.5 | 9 | 9.5 | |
| Left thigh | 5.5 | 6.5 | 8 | 8.5 | 9 | 9.5 | |
| Right leg | 5 | 5 | 5.5 | 6 | 6.5 | 7 | |
| Left leg | 5 | 5 | 5.5 | 6 | 6.5 | 7 | |
| Right foot | 3.5 | 3.5 | 3.5 | 3.5 | 3.5 | 3.5 | |
| Left foot | 3.5 | 3.5 | 3.5 | 3.5 | 3.5 | 3.5 | |

Total BSAB _____

Adults and Children:

- Give IV fluid replacement according to the calculation below
- With mild burns (< 15% adults BSA burned or < 10% children) give oral fluid replacement therapy using as much ORS as the patient will tolerate

Adults:

- With 15% or more and children with 10% or more require IV fluid resuscitation
- Calculate according to Parkland formula:-Adults: 4mls per /kg body wt/ TBSA

Children:

- 3mls per kg body wt/TBSA + maintenance
- Maintenance for children

20.2.2 Calculation of IV fluid replacement

- The object is to maintain normal physiology as reflected by urine, vital signs and mental status
- Use Ringer's Lactate IV infusion or (if this is not available) Normal Saline
- A general formula and dosage schedule that may be used for the first 24 hours is:
 - Total volume of IV infusion required (before above additions) = 4 ml/kg x % BSA burned plus normal daily requirement

- Give 50% of this total in the first 8 hours calculated from the time of burn
- Give 50% in the next 16 hours
- Give analgesic treatment (Section 24.1)
 - Strong analgesia (e.g. Morphine or Pethidine) will be required for the first 48 hours
- Under aseptic conditions, gently cleanse the lesion with Cetrimide 15% +
 - **Chlorhexidine 1.5%** diluted 1 part in 20 parts water

Alternative:

- Cleanse with Hydrogen Peroxide Solution (10 vol) or soap and water
- Never use alcohol-based solutions
- Repeat the cleansing each day debriding the lesion and removing necrotic tissue as necessary
- Give tetanus toxoid vaccination (see Section 12.3) If patient is developing signs of tetanus
 - Give Tetanus Antitoxin (ATS) 1,500 units s/c or IM
- Once the lesion is clean/clear of necrotic tissue:
 - Refer for skin grafting, if necessary, otherwise
 - Dress the burn with Paraffin GauzeDressing

- Cover this with dry gauze dressing thick enough to prevent seepage through to the outer layers
- Change the dressing after 2-3 days, and then as necessary

If the burn becomes infected:

- Apply Silver Sulphadiazine Cream 1% bd
 - Before application, completely remove any old topical medication
 - Cover with sterile gauze

If the patient becomes ill after burn infection:

- Carry out culture and sensitivity testing on the exudates
- Treat with systemic antibiotic(s) according to findings

20.3 Open wound

General Management

- Appropriately suture fresh wound (i.e. < 6 hours old)
- Do not suture penetrating wound
- Refer to the next level of care

Note:

- Any skin damage overlying a fracture makes it an open fracture
- It is important to ascertain the cause of the wound

Alternatively

- Clean with Hydrogen Peroxide Solution (10 vol)
- For local infiltration or as a peripheral nerve block use Lignocaine Hydrochloride injection 1% maximum dose: adults 4mg/kg body weight; children 4mg/kg
 - Or use Lignocaine 1% (10mg per 1ml)+ adrenaline 1:200,000 injection
 - ✓ Maximum dose: adults7mg/kg, children 7mg/kg

Note: Do not use anaesthetics containing adrenaline for anaethesia in digits or appendages due to risk of ischaemic necrosis

- Always remember to give Tetanus Toxoid
 Vaccination (Section 12.3)
- If patient has signs of tetanus give Tetanus
 Antitoxin Serum (ATS) 1,500 IU s/c or IM
- If wound is grossly contaminated give
 Tetanus Antitoxin Serum 3,000 IU s/c or IM (alongside copious washing)

20.4 Insect stings and bites

20.4.1 Envenomation by insects

- Bees, wasps: Usually benign, but may provoke either laryngeal oedema or anaphylactic shock (see Section 5.1.1)
- Spiders, scorpions: The majority of spiders are benign. If a truly toxic species is thought to be responsible apply first aid and supportive measures as for snake bite (see Section below)

20.4.2 Snake bite

 Venom diffuses mainly via the lymphatics, not via blood vessels, tourniquets are thus of little use

First Aid Treatment

Clean the would with Cetrimide +
 Chlorhexidine solution 15% + 5% diluted 1
 in 20

Alternatively

- Clean with Hydrogen Peroxide Solution (10 vol)
- Apply film constant pressure to the site of the bite
- Apply a crepe bandage firmly to the entire limb
- Immobilize the patient for 12 hours observation
- Give reassurance

Note: Not all patients with snake-bite should be given **Anti-venom**

Administration of Snake Anti-venom

- Ensure that the Anti-Venom Solution is clear
- Give 0.5 ml of Adrenaline 1/1000 subcutaneously if needed
- Give 100 ml of the Polyvalent Antivenom as an iv infusion, diluted in 300 ml of Normal Saline
- Children:
 - ➢ Dilute in 0.4 ml/kg of Saline
 - Give the infusion slowly for the first
 15 minutes (most reactions occur within this period
 - ➤ Thereafter increase the rate gradually until the whole infusion is completed within 1 hour

If there is a history of allergy:

 The patient may still need to be given anti-venom because of systemic poisoning, but take particular care

If a reaction occurs:

• **Hydrocortisone** may need to be administered in addition to **Adrenaline**

If there is no clinical improvement by the end of the infusion:

Repeat the same dose as above

Note: Reserve Polyvalent Snake Antivenom (anti-snakebite serum) for patients with one or more of these signs and symptoms:

- > Hypotension
- Vomiting
- Neurotoxicity
- Haemotoxicity

Supportive Therapy

- Give reassurance most snake bites are not dangerous
- Treat shock if any (see Section 5.1.1)
- Give an antihistamine:

Adults:

Give Promethazine 25 mg/day up to 75 mg/day in equal doses

Children > 6 months:

- Give Promethazine 1 mg/kg/day given in divided doses every 12 hours
- Give **Tetanus Toxoid Vaccination** (Section 12.3) If patient is developing signs of tetanus:
- Give Tetanus Antitoxin (ATS) 1,500 IU s/c or IM
- Give Benzyl penicillin 2.4 MU once daily for 5 days
 - > 25,000 units/kg daily
- Eventually excise sloughs and graft skin early

20.5 Superficial injury, bruise, minor cut

General management

Clean the wound with Cetrimide 15% +
 Chlorhexidine Solution 1.5% diluted 1 in 20
 parts water

Alternatively

- Use **Hydrogen Peroxide Solution (10 vol)**If the patient is unimmunised or not fully immunized and the wound is grossly contaminated:
 - ➤ Give **Tetanus Toxoid Vaccination** (Section 12.3) If the wound is dirty:
 - ✓ Adults: Give Benzylpenicillin 2.4 MU IM stat
 - ✓ Children: Give Benzylpenicillin 25,000 units/kg

Then

- ✓ Adults: Give Amoxicillin 500mg every8 hours for 5 days
- ✓ Children: Give Amoxicillin 15mg/kg

 If Penicillin hypersensivity:
 - ✓ Give Erythromycin 250 mg every 6 hours for 5 days

21.0 Renal Conditions

 Starts with upper complicated then acute lower urinary tract conditions

21.1 Cystitis/Urethritis

Signs and Symptoms

- Pain when passing urine
- Fever
- Blood in urine

Note: Think of Bilharzia; Diabetes Mellitus and cancer of the urinary bladder also in patient with similar presentation and symptoms are recurrent

Treatment

- Ensure adequate fluid intake
- For acute uncomplicated (all non-pregnant women, symptoms duration less than 1 week, not men or catheterized patients) give
 Ciprofloxacin 500mg bd for 5 days

Alternatively

- Give Nitrofurantoin 100mg qid with food for 7 days
- Consider urine microscopy, culture and sensitivity if no response or recurrent infections to guide treatment

21.2 Complicated Urinary Tract Infections

 Includes men, pregnant women, catheterized patients, patients with abnormal urinary tracts and those with symptoms > 1 week

Treatment

- Give Ciprofloxacin 500mg orally bd for 5 days or
- Give Co-amoxiclav 375mg tds or 625mg bd for 5 days

Alternatively

- Give Gentamycin 240mg IM or IV stat
- Followed by any of the above oral antibiotics
- Consider urine microscopy, culture and sensitivity to guide treatment if no response to above treatment or recurrent infections

21.3 Upper Urinary Tract Infections

21.3.1 Pyelonephritis

Sign and Symptoms

- Significant fever, rigors, vomiting, flank pain Treatment
 - Give Ciprofloxacin 500mg orally bd for 10-14 days or
 - Give Co-amoxiclav 375 mg tds for 10-14 days
 - IV fluids if clinically indicated

Alternatively

- Give Gentamycin 240mg IM or IV stat
- Followed by any of the above oral antibiotics for 10-14 days
- Refer patients with recurrent UTI for further investigations

21.4 Urinary Tract Infections (UTIs) in Children

Sign and Symptoms

- In young children, UTI often presents with nonspecific signs, on examination, such as vomiting, fever, irritability, or failure to thrive
- Older children may present with more specific signs such as abdominal pain, pain on passing urine or increased frequency of passing urine

Investigations

- Try to get a clean catch sample of urine
- In sick infants, suprapubic aspiration of urine may be required
- Dipstick the urine looking for leukocytes and nitrites
- Do microscopy if positive dipsticks result and consider UTI if more than 5 white cells per high power field

Treatment

| <3 months | Admit the child and treat with: Gentamicin and Benzylpenicillin, Ceftriaxone as second line; consider Blood Culture and Lumbar Puncture | |
|---|---|--|
| >3 months or older child with signs of systemic illness (T>38C, rigors, renal angle tenderness) | Admit and treat with Gentamicin and Benzylpenicillin or Ceftriaxone Change to oral antibiotic when fever settled and improving to complete 10 days treatment | |
| >3 months or older child with no signs of systemic illness | Treat with oral antibiotics Cotrimoxazole 10mg/kg bd or Amoxycillin, Ciprofloxacin, Nitrofurantoin 1.5mg/kg qid | |

- Refer to hospital for further investigations and follow up in the following:
 - > Atypical UTI:
 - ✓ Seriously ill
 - ✓ Poor urine flow
 - ✓ Abdominal or bladder mass
 - ✓ Raised creatinine if measured
 - ✓ Septicaemia
 - ✓ Failure to respond to treatment with suitable antibiotics within 48 hours

Recurrent UTIs:

- ✓ 2 or more episodes of UTI with acute pyelonephritis/upper urinary tract infection
- ✓ One episode of UTI with acute pyelonephritis/upper urinary tract infection plus one more episode of UTI with cystitis/lower urinary tract infection
- ✓ Three or more episodes of UTI with cystitis/lower urinary tract infections
- All males > 1 year of age with UTI
 - They should all have a renal tract ultrasound scan to look for structural abnormalities
 - Children with abnormalities should be followed up in clinic at the hospital

21.5 Acute Nephrotic Syndrome

- Most often occurs as a complication of a streptococcal infection
- Usually manifests itself 1-5 weeks after an episode of pharyngitis, impetigo, or infected scabies
- Affects mainly children <3 years old, and adults, who should be referred to a doctor

Note: Renal diseases can easily be mistaken for malnutrition

General Treatment

- Monitor BP, urine output, weight
- Urine dipstick and microscope
- Avoid added salt
- Treatment is usually supportive

Adults:

 Give Phenoxymethylpenincillin 500 mg every qid for 7 days

If oedematous:

Give Frusemide 40-80 mg od

If hypertension is present:

Treat accordingly (Section 2.3)

21.6 Nephrotic syndrome

Diagnosis

- Proteinuria
- Hypo-albuminaemia
- Oedema
- High cholesterol
- Investigate a possible cause
 - Blood pressure
 - Urinalysis
 - Urine microscopy (look for casts and check for schistosoma ova)
 - MPs and PCV
 - Electrolytes
 - Urea and Creatinine
 - Imaging on individual basis: renal USS, CXR, cardiac echo

General Management

- Adequate protein intake
- Restrict salt intake
- Monitor fluid intake and output
- Daily weight measurements

Treatment

Adults

- Give **Frusemide** 40-80 mg as a single dose each morning
- Give **Enalapril** 10-20 mg od (use with caution, stop if renal function deteriorates)
- A trial of steroids is indicated (responsiveness to steroids in adults is less than in children). Give **Prednisolone** 50-60 mg od for up to two months, tapering off is required after response

If Schistosomiasis is diagnosed or suspected as cause

• Give Praziquantel 40 mg/kg single dose

Children

- Give Prednisolone 2mg/kg/day or 60mg/m2/day for at least 4 weeks then a reducing regimen
- Should be reviewed at hospital clinics before reducing dose
- Only use steroids if picture consistent with minimal change disease (heavy proteinuria, NORMAL BP, NO haematuria

- Consider Frusemide 1-4mg/kg/day oral or IV when hypertensive. Review use of diuretics on a daily basis
- **Prophylactic Penicillin** or **Amoxicillin** oral to prevent infections.
- Encourage mobilization as normal

If presentation is acute:

- Give Phenoxymethlyl Penincillin 500 mg qid for 7 days
- Refer to Nephrologist

21.7 Renal Colic

Treatment

- Intravenous fluids
- Give Morphine or Pethidine. (Section 24.9)
- Give Hyoscine butylbromide 20 mg deep IM stat
- Repeat after 30 minutes if necessary
- Ensure fluid intake of 3-4 litres/day after the crisis
- Intravenous Pyelography is usually indicated

21.8 Renal Failure (acute)

- Patients with acute renal failure should be referred to a hospital
- Carefully check the use of any drug in renal failure and reduce drug doses where required, see below

Signs and Symptoms

- Oliguria
- Oedema

- Vomiting
- Look for clues for the cause of renal failure, which include:
 - Shock
 - Acute glomerulonephritis
 - Use of herbal remedies containing nephrotoxins

General Management

- Assess the hydration status of the patient
- Patients who are dehydrated will need fluid resuscitation
- Avoid Ringer's Lactate fluids
- Patients who are fluid overloaded will need fluid restriction and/or diuretics
- Restrict salt intake
- Weigh the patient daily
- Carefully monitor fluid intake and output on a chart
- Reduce the rate of rise of urea:
 - Give adequate calories
 - > Restrict protein in the diet
 - Treat hyperkalaemia:
 - ✓ Restrict potassium intake by restricting fruits, vegetables, meat and feezy drinks
 - ✓ If potassium is > 6.5mmol/l give Insulin 10 Units in 50ml of 50% Dextrose infusion over 30 minutes
 - ✓ Give a Potassium Binding Resin 30-60g orally

 Refer patient to Central Hospital for further management and consideration for dialysis if not responding to measures above

Indications for dialysis include:

- Hyperkalaemia refractory to insulin shifting
- Fluid overload not responsive to diuresis
- Metabolic acidosis
- Pericarditis
- Uremic symptoms and signs (encephalopathy, haemorrhagic pericardial effusion bleeding)
- Lithium and theophylline overdose

Note:

- Treat complications of renal failure such as convulsions, hypertension
- Do an HIV and Hepatitis B test before referral for dialysis

21.8.1 Use of medicines in renal failure/impairment

Note: Take great care when prescribing any medicine and carefully check medicine prescribing information (e.g. in BNF, MNF) regarding use in renal failure/impairment

Usually safe medicines:

- Doxycycline
- Erythromycin
- Penicillin
- Phenytoin
- Rifampicin

Use with care in reduced doses:

- Amoxycillin
- Chloramphenicol (avoid in severe impairment)
- Cotrimoxazole
- Diazepam
- Digoxin
- Insulin
- Isoniazid-containing medicines
- Pethidine (increase dose interval, avoid in severe impairment)
- Phenobarbitone
- Propranolol
- Antiretroviral medicines

Avoid using:

- ACE inhibitors (eg. Captopri)
- Aspirin and other NSAIDS (eg. Ibuprofen, Indomethacin)
- Codeine
- Ethambutol
- Gentamycin
- Nalidixic acid
- Nitrofurantoin
- Streptomycin

22.0 Poisoning

22.1 General principles of treatment

- Determine details of the poisoning:
 - What was the poison?
 - When did the poisoning take place?
 - What kind of poisoning took place? e.g. by swallowing, inhalation, contact with the skin or eyes
 - How much was taken?
- Prevent further exposure to the poison (if possible)
 - Remove contaminated clothing
 - Wash contaminated skin with soap and lots of cold water

Note: In all cases of poisoning in children, the social workers must be involved to assess the social circumstances and take the necessary steps to ensure the child is not at risk of further incidences

General Management

- Conserve body heat (if necessary)
- Maintain respiration
- Clear the airway
- Breathing-Maintain ventilation use artificial respiration if necessary, patient may need ventilation

- Maintain BP/treat shock
 - Correct hypotension
 - ➤ Elevate the legs
 - Correct hypertension
 - In refractory shock discuss with anesthetist
- Maintain fluid balance
- Monitor fluid intake and output

22.2 Swallowed poisons

Treatment

- Prevent gut absorption
- Empty the stomach (if appropriate)
 - Only do this if within 4 hours of the poisoning and if the patient is conscious
- Do not empty the stomach if:
 - A corrosive substance was swallowed,
 e.g strong acid or alkali, bleach
 - Paraffin or a petroleum product was swallowed
 - The patient is unconscious or convulsing
 - > The subsistence is not known

- Treat any complications as necessary
 - E.g. hypothermia, hypoglycaemia, convulsions, electrolyte or acid/base disturbances

22.2.1 Methods of emptying the stomach

- Induction of vomiting
 - Give Ipecacuanha emetic mixture
 - ✓ Children < 18 months: 10ml
 - ✓ Children > 18 months: 15ml
 - ✓ Adults: 15-30 ml
 - > Repeat after 20 mins if ineffective
 - Follow this with 15 ml/kg of water

Note: It is essential to prevent any vomit from entering the lungs

- Stomach wash-out (gastric lavage)
 - Should only be done by staff familiar with the procedure
 - Lie the patient head down on the left side
 - Pass a wide gauge soft rubber tube(Ryle's tube) into the stomach
 - Tube should be wide enough to allow large particles to pass through. e.g. tablets
 - Pour 300ml tap water down the tube
 - For children > 5years: use 100-200 ml water
 - For children < 5 years: use normal saline instead of water

- Aspirate with the patient in the head down position, taking special note of the airway
- Repeat lavage until aspirated fluid is clear

22.2.2 Use of activated charcoal

- 50-100g Activated Charcoal will prevent absorption of most medicines given within 1 hour of ingestion
- Only effective if given within 4 hours of poisoning when most of the poison is still in the stomach
- Only give Activated Charcoal after vomiting (induced or otherwise) has ceased
- Do not use ordinary charcoal it will have no effect
- Do not use activated charcoal in the following situations:
 - ➤ If the patient is unconscious, drowsy or having fits, because of the risk of choking
 - At the same time as, or just before giving, ipecacuanha or any oral antidote as it may bind these and prevent them working
 - For poisoning by acids, alkalis, alcohol, iron and petroleum products

22.2.2.1 Administration of activated charcoal

- Add 50 g (children: 1 g/kg) to 400 ml water in a bottle
- Mix well by shaking until all the powder is wet
- Administer by the gastric lavage tube (unless the patient agrees to drink the charcoal slowly)
- Repeat if required after 4-6 hours

22.3 Paraffin, petrol and other petroleum products

- Includes paint thinners, organic solvents, etc
- The main danger from these is damage to lung tissue and liquid pneumonitis following aspiration

General Measures

- Take great care to prevent the substance entering the lungs
- Do not make the patient vomit
- Do not do gastric lavage except:
 - Where the amount of paraffin, etc, swallowed was high (> 10 ml/kg) as these levels may cause brain damage
 - Only after endotracheal intubation under anaesthesia
- Treat any pulmonary oedema and pneumonia as required
- Giving un absorbable oral liquid, e.g medicinal liquid paraffin

22.4 Iron poisoning

Treatment

- A dose of 20 ml/kg of Iron Syrup or 2-3 iron tablets/kg may be fatal in children.
 Abdominal X-ray may show the number of tablets swallowed
- In severe cases there are risks of vomiting and gut haemorrhage in the acute stage and liver necrosis and shock after 1-2 days.
 Therefore observe the patient for at least 48 hours
- Remove any tablets by inducing vomiting and/or by gastric lavage

In less serious cases:

 Give Desferrioxamine 5-10g orally or by nasogastric tube in 50-100 ml of Sodium Bicarbonate solution 5%

In serious cases:

- Give Desferrioxamine 15 mg/kg/hour by IV infusion in Dextrose 5% or Normal Saline solution
 - Max: 80 mg/kg in each 12 hour period
 - Continue until free of symptoms for 24 hours

22.5 Salicylate (Aspirin) poisoning

- Gastric empting is delayed
- Always empty the stomach
- Give repeated doses of **Activated Charcoal** to delay absorption of any remaining poison (See Section 22.2.2.1)

- Watch for and treat hypoglycaemia, convulsions and metabolic acidosis
- In severe cases:
 - Darrow's ½ strength in Dextrose 5% infusion with added Sodium Bicarbonate (30 mmol/litre) or Ringers Lactate may be needed to increase renal excretion

22.6 Organophosphate or Carbamate Poisoning

- Very toxic chemicals found in insecticides and pesticides, eg. Some rat poisons
- Poisoning may be by ingestion, inhalation or absorption through the skin
- Presents with anxiety, restlessness, small pupils, increased secretions and bradycardia

General Management

- Remove any contaminated clothing
- Establish and maintain airway
- Artificial respiration with oxygen may be required at any stage during the first 24 hours after poisoning
- Empty the stomach if poison swallowed
- If there is skin contact with the poison, wash them thoroughly
- Wear rubber gloves to prevent contamination
- Do not rub the skin.
- Shave hair if heavily contaminated
- Give Atropine 1.2mg IV or IM (children: 0.05 mg/kg)

 Then give 0.6 mg (children: 0.05 mg/kg) every 10 minutes as required to achieve and maintain atropinisation (hot dry skin, dry mouth, widely dilated pupils, fast pulse)

In severe organophosphate poisoning:

- Give an initial Atropine dose of 4-6mg (children: 2 mg)
- Repeat 2 mg every 10 minutes as required to achieve and maintain full atropinisation
- Total needed in first 24 hours is usually < 50 mg
- High dose **Atropine** may be required for several days

In severe organophosphate poisoning only and in cases not responding to atropine:

Give Pralidoxime Mesylate 1-2g concurrently with Atropine

Children: 20-40 mg/kg

- Administer by slow IV (over 15-30 minutes)
 as a 5% solution in water for injections
- If IV not possible: give IM or s/c and repeat once or twice at 4-6 hour intervals if needed
 - If possible: monitor treatment by determination of bloodcholinesterase concentrations

Note: Do not give **Pralidoxime** (or other oximes) in carbamate poisoning

22.7 Paracetamol poisoning

- Paracetamol is an ingredient of many over the counter pain cold flu remedies
- A dose over 150 mg/kg (ie. approx 10 g in an adult) may cause severe liver and (less frequently) kidney damage within hours of ingestion
- In the first 24 hours there may be nausea and vomiting or there may be no sign of poisoning
- Persistence of these symptoms and associated right subcostal pain and tenderness usually indicates hepatic necrosis
- Liver damage reaches a maximum 3-4 days after poisoning and may be fatal

Even if there are no significant early symptoms, overdose patients should be urgently transferred to hospital

If overdose occurs within 4 hours of admission:

- Empty the stomach to remove ingested medicine
- If respiration is depressed: do not use emesis –
 use airway protected gastric lavage instead
- Keep patient warm and quiet
- Observe carefully for at least 3-4 days
- Monitor fluid, electrolytes, blood glucose, liver and kidney function

22. Poisoning

 Give supportive care and correct fluid and electrolyte balance as required

If within 24 hours of overdose with over 10g of paracetamol:

- Give the specific antidote N-acetylcystine as an IV infusion in Glucose 5%
 - ➤ Initially give 150mg/kg in 200ml over 15 minutes
 - Then give 50 mg/kg in 500 ml over 4 hours
- ➤ Then give 100mg/kg in 1 over 16 hours If a serious reaction occurs:
 - Stop the infusion
 - Treat the reaction (see Section 5.1.1)
 - Restart the infusion

23.0 Nutritional disorders

23.1 Malnutrition

23.1.1 Definition Severe Malnutrition

- Children 6-59 months
 - < -3Z score of expected weight for hieght
 - > <11.5cm MUAC
- Children 5-9 years
 - ➤ MUAC <13.0cm
- Children 10-12 years
 - ➤ MUAC <16.0cm
 - <-3Z score WFH</p>
 - \triangleright Oedema +1 or +2
- Children 12–14 years
 - Bilateral pitting oedema (to be assessed by a clinician for medical causes)
 - ➤ Weight for Height (WHZ) < -3
 - Mid-Upper Arm Circumference (MUAC) < 16.0 cm</p>
- Children 15-18 years
 - Bilateral pitting oedema (to be assessed by a clinician for medical causes
 - ➤ MUAC < 18.5 cm</p>
 - ➤ Body-Mass Index (BMI) < 16

- Adults (non-pregnant/ post-partum)
 - Bilateral pitting oedema (to be assessed by a clinician for medical causes)
 - > MUAC < 18.5 cm
 - ➤ BMI < 16
- Pregnant/ post-partum women
 - ➤ MUAC < 19.0 cm

23.1.2 Definition Moderate Malnutrition

- Children 6-59 months
 - -3Z to <-2Z score of expected weight for Height
 - MUAC 11.5 to 12.5cm
- Children 5-12 years
 - ➢ BMI for age -3 to <-2Z score</p>
- Children 12–14 years
 - \triangleright WHZ ≥ -3 and < -2
 - MUAC ≥ 16.0 to < 18.5 cm</p>
- Children 15-18 years
 - ➤ MUAC ≥ 18.5 to < 22.0 cm
 - \triangleright BMI ≥ 16.0 to < 17.0cm
- Adults (non-pregnant/ post-partum)
 - ➤ MUAC ≥ 18.5 to < 22.0 cm
 - \triangleright BMI ≥ 16.0 to < 17.0cm
- Pregnant/ lactacting women
 - MUAC ≥ 19.0 to < 23.0 cm</p>

23.1.3 Malnutrition Investigations and Signs and Symptoms

Investigations

- Laboratory examinations identify nutrient deficiencies:
 - For blood (Hb, haematocrit)
 - Protein (serum albumin)
 - Micronutrient (vitamin B12, iron, zinc, and folate)
 - Blood glucose
 - Lipid (cholesterol and triglycerides)

Signs and Symptoms

- Symptoms that affect food intake
 - Diarrhoea
 - Nausea
 - Vomiting
 - Anorexia
 - Mouth and throat sores
 - Oral thrush
- Signs of clinical malnutrition
 - Wasting and weight loss
 - > Skin changes
 - Oedema
 - Apathy
 - > Hair changes
- Signs of anaemia
 - Pale conjunctiva, gums, nails, and skin
 - Breathlessness
 - Rapid pulse
 - Oedema

Dietary assessment

 Information about the types and amounts of foods eaten, appetite, food habits and eating behaviours to identify factors that affect food intake such as side effects of medications

23.1.4 Management of Malnutrition in Adolescents and Adults

- Refer to The National Guidelines for the Management of Acute Malnutrition in Adolescents and Adults
- All ambulatory adults and patients should be treated as outpatients
- The ready to use therapeutic food (RUTF)
 OR the CSB should be used in combination with a normal diet

23.1.5 Treatment of Severe Malnutrition

2 pots of Ready-To-Use Therapeutic Food
 (RUTF) 260g, 2700 kcal per day OR 6 sachets of
 RUTF 92g, 3000 kcal per day depending on
 what is available

 If patients achieves BMI of 17 or MUAC of 22cm (in adolescents WHZ <-2) then treat as moderate malnutrition (see Section 23.2.2 below)

23. 1.6 Treatment of Moderate Malnutrition

Give 4.5kg of Likuni Phala (containing 10% sugar) and one litre of Vegetable Oil per month (1500 kcal per day) OR 3kg of Corn Soya
 Blend++ (Super cereal)

23.1.7 Outpatient Follow Up

- Follow up patients monthly
- Plot their weight gain
- All patients not responding after three months should be reviewed by a clinician
- If not medically indicated, treatment can continue for up to six months after initiation, or if HIV infected up to three months after commencing ART

23.1.8 Use of Milk Based Formulations F75 and F100

- Patients requiring Naso Gastric (NG) feeding cannot tolerate RUTF or are severely ill require milk based formulations F75 and F100 feeds
- Patients should be weighed twice weekly
- Milk based formulations are given in two phases

23.1.8.1 Phase 1

- Use F75 to stabilize patient
- Treat infections and other urgent medical problems
- Provide sufficient energy and nutrients to stop further loss of muscle and fat tissue
- Correct fluid and electrolyte imbalance
- Give at least 5-6 feeds per day
- Night feeds may be helpful, particularly with NG tube feeding

How to prepare F75

- Mix one packet of F75 with 2 litres of cooled boiled water to make 2400ml of formula
- Give the amounts as in the Table 29 below unless patient is receiving IV fluids in which case amounts should be reviewed
- Intravenous fluids are discouraged in severe malnutrition as they have little nutritional value, and can cause fluid overload

Amounts of F75 given to patients in Phase 1

Class of weight (kg) 8 feeds per day 6 feeds per day 5 feeds per day ml for each feed

| 15.0-19.9 | 260 | 300 | 400 |
|-----------|-----|-----|-----|
| 20.0-29.9 | 300 | 350 | 450 |
| 30-60 | 350 | 400 | 500 |

- Patients should not eat any other foods or fluids, during Phase 1 unless they have diarrhoea
- Patients with diarrhea should be given ORS (Resomal)

23.1.8.1.1 Transition Phase

- Change to **F100** as soon as the patients appetite returns
- Control the amount of F100 during the transition phase to avoid the risk of heart failure
- RUTF may be introduced at this stage in addition to F100 so patients are familiar with it when they reach Phase 2
- Give 1 pot (or 3 sachets) of RUTF over the 2 day transition period as a taste dose
- The patient is not required to finish the RUTF
- The number of feeds, their timing and the volumes given remains exactly the same: F100 (100ml = 100kcal) is used in the transition phase
- Patients should normally move to Phase 2 after
 2 days on Transition phase
- Patients with NG feeding tubes should remain on transition phase quantities but these should be reviewed if NG feeding continues for more than two weeks

How to prepare F100

- Mix one packet of F100 with 2 litres of cooled boiled water to make 2400ml of formula
- Give the amounts as in the Table 30 below to each patient unless receiving IV fluids in which case amounts should be reviewed

23.1.8.2 Phase 2

- Aim of phase 2 is to achieve rapid weight gain and rebuild lost tissues and this requires more energy, protein and micronutrients than were needed for Phase 1
- **F100** without iron is given during this phase
- Give the amounts as in Table 31 below to each patient

Amounts of F100 given in Phase 2

F100 (ml) per feed phase 2 for 6 & 5 feeds

| Class of weight (kg) | ass of weight (kg) MI per feed 6 feeds per 24 hours | |
|----------------------|--|-----|
| 15.0-19.9 | 550 | 650 |
| >20 | 750 | 900 |

- Give one tablet of Fefol 200mg or Ferrous sulphate 200mg per day in phase 2 if clinically indicated
- If inpatient is well and can tolerate solid food,
 RUTF should be used in Phase 2 instead of
 F100

- RUFT treatment is the same as per for outpatients and should be used alongside a normal diet
- Give the amounts below until patients BMI reaches 17
 - 2 pots of RUTF 260g, 2700 kcal per day OR
 - 6 sachets of RUTF 92g, 3000 kcal per day
- Once patient achieves a BMI of 17 or MUAC 22cm (if BMI can not be taken), Or Weight/Height>80%, they should be transferred to treatment for moderate malnutrition

23.1.9 Discharge Criteria (when to stop treatment of malnutrition)

Adults

 BMI of 18.5 and bilateral oedema has gone for 10 consecutive days or MUAC 23cm (to be used only if BMI can not be taken)

Pregnant and lactating women up to 6 months after delivery

MUAC 23cm

Adolescents 12-18 years

 Weight/Height >85% and bilateral oedema has gone for 10 consecutive days

23.2 Malnutrition (protein-energy) in Children

23.2.1 Mild malnutrition

- Do a full assessment, especially looking for underlying disease such as TB, HIV infection, etc.
- Ensure weekly attendance at nutrition clinic
- Give food supplements
- Give Vitamin A 100,000 units as a single dose

23.3 Acute Malnutrition in Children

23.3.1 Moderate Acute Malnutrition

Signs and Symptoms

- Children 6-59 months
 - MUAC 11.5 to <12.5cm or</p>
 - ➤ WFH -3z to -2z score and
 - No Oedema
 - Appetite
 - Clinically well
- Children 5-12 years
 - BMI for age -3z to <-2z score and</p>
 - No oedema
 - All discharges from NRU or OTP

Investigations

- Look for visible signs of wasting
- Check whether the child has got good appetite

- Check whether the patient has medical condition that will impair nutrition status e.g TB or HIV
- Determine severity of malnutrition using MUAC, Oedema, WFH and BMI for age

Management

- Refer to Supplementary Feeding Programme
- Do a full assessment, especially looking for underlying disease such as TB, HIV infection, etc.
- Refer all children who have not been immunized to the nearest MCH clinic
- Give Vitamin A
 - Child 12-59 months: 200,000IU
 - > Child 6-11 months: 100,000IU
- Give Iron +Folic Acid Supplementation
 - <10kg: 1 tablet every two weeks</p>
 - > >10kg: 2 tablets every two weeks

 Give Albendazole or Mebendazole as follows:

| AGE | ALBENDAZOLE | MEBENDAZOLE |
|----------|--------------|------------------|
| <12 | None | None |
| months | | |
| 12 to 23 | 200mg single | 100mg twice |
| months | dose | daily for 3 days |
| | | |
| 24 to 59 | 400mg single | 100mg twice |
| months | dose | daily for 3 days |
| | | |
| 1 | | l . |

Provide 200g of CSB++ per day (3kg for 14 days)

Follow Up

- Follow up visits to SFP occur every 2 weeks
- At every visit assess nutrition status of the patient
 - Take weight and MUAC measurement at each follow up visit
 - Check for oedema at each follow up visit
 - Calculate WFH Z score or BMI for age at each follow up visit
 - Assess nutrition status for possible discharge or referral

Referral to OTP or NRU

 If the child 6 months to 12 years in SFP deteriorates and meets OTP admission criteria, refer to OTP If the child 6 months to 12 years deterrirates in SFP, cannot be treated in OTP and now meets the admission criteria for NRU refer to NRU

23.3.2 Severe Acute Malnutrition without Complications

Signs and Symptoms

- Children 6-59 months
 - ➤ MUAC <11.5cm or</p>
 - ➤ WFH <-3Z scores or
 - Oedema +1 or +2
 - With
- ✓ Appetite for RUTF
- ✓ Clinically well
- If the child is HIV + admit to OTP with MUAC <12.5cm</p>
- Children 5-12 years
 - MUAC <13.0cm (5-9 years)</p>
 - MUAC <16.0cm (10-12 years) or</p>
 - BMI for Age <-3Z score or Oedema +1 or +2
 - With
- ✓ Appetite
- ✓ Clinically well

Management

- Triage urgent cases
- Conduct Anthropometric Assessment
- Conduct Appetite test
- Conduct medical assessment
- Admit the child in the Outpatient Therapeutic Programme

Routine Medical Treatment

- Give Vitamin A Supplementation
 - 6-11 months (100, 000IU)
 - > 12 months-12 years: 200, 000IU
 - ➤ Give **Vitamin A** on admission
 - ✓ Do not give to children with oedema
 - ✓ Do not repeat the dosage of Vitamin A if the child is readmitted or has already received curative dose of Vitamin A during the last 6 months

• Give Amoxicillin as follows:

| Weight of the Child (KG) | Syrup | Syrup | Tablets |
|--------------------------|---------------------------------|----------------------------------|--|
| | 125mg/5ml | 250mg/ml | 250mg |
| | Dosage-give | Dosage-give tds | |
| ≤ 2.0 | 62.5mg (2.5ml) 3X per day | 62.5mg (1.25ml) 3X per day | 62.5mg (½4 tablet) 3X per day |
| 2.0-9.9 | 125mg | 125mg | 125 (½ |
| | (5ml) 3X | (2.5ml) 3X | tablet) 3X |
| | per day | per day | per day |
| 10.0-30.0 | 250mg | 250mg | 250mg (1 |
| | (10ml) 3X | (5ml) 3X | tablet) 3X |
| | per day | per day | per day |
| >30.0 | Give tablets | Give tablets | 500mg (2 tablets) 3X per day |

- Give on admission except for children under 2kg
- If Amoxicillin is not available, use
 Cotrimoxazole according to IMCI protocol
- Give Folic Acid
 - Give on first day
 - Give to all beneficiaries
 - ➤ Give 5mg single dose

Antimalarials

 LA should only be prescribed if there is a positive diagnostic test

| WEIGHT (KG) | DOSE |
|-------------|---------------------------|
| 5-14.9 | 1 TABLET (2x/day/3 days) |
| 15-24.9 | 2 tablets (2x/day/3 days) |
| 25-35 | 3 tablets (2x/day/3 days) |
| >35 | 4 tablets (2x/day/3 days) |

• Give Albendazole or mebendazole

| AGE | ALBENDAZOLE | MEBENDAZOLE |
|-----------------|--------------|------------------|
| <12 months | None | None |
| 12 to 23 months | 200mg single | 100mg twice |
| | dose | daily for 3 days |
| 24 to 59 months | 400mg single | 100mg twice |
| | dose | daily for 3 days |

 If the child is transferred between OTP and NRU ensure that this dose is not repeated

Nutritional management

 Ready to Use Therapeutic Food (RUTF) is used to treat patients with SAM in OTP

- The amount of RUTF is based on weight of the patient (Refer to Table 9; RUTF Ration Table in the Guidelines for Community management of Acute malnutrition)
- Give the required RUTF to the carer
- Counsel the carer on how to give the RUTF to the child

Follow Up

- The patient should come every week for nutrition assessment, medical checkup and to receive weekly supply of RUTF
 - Key activities should be done to assess the patient
 - ✓ Take weight and MUAC measurements
 - ✓ Calculate nutritional status
 - ✓ Review target weight for discharge
 - ✓ Assess for any illness since the previous visit
 - ✓ Conduct physical assessment for any illness
 - ✓ Conduct appetite test for RUTF
 - ✓ Make decision if there is need for any additional management
 - ✓ Reinforce key health and nutritional key messages

Referral

- Children should be transferred to NRU if:
 - Medical condition deteriorates or any condition develops
 - ✓ The child refuses RUTF
 - ✓ There is an increase in bilateral pitting oedema (or Oedema appears when it was previously absent)
 - ✓ There is weight loss for 3 consective OTP sessions
 - ✓ There is static weight (no weight gain) after five consecutive OTP sessions
 - ✓ Discharge criteria have not been reched after 3 months in OTP

23.3.3 Severe Acute malnutrition with Complication

Signs and Symptoms

- Bilateral Pitting Oedema +3
- Marasmic Kwashiorkor
 - MUAC <11.5cm (6-59 months)</p>
 - MUAC <13.0cm (5-9 years)</p>
 - MUAC <16.0cm (10-12 years) or</p>
 - ➤ WFH <-3Z score (6-59 months)
 - ➢ BMI for Age <-3Z score (5-<12 years)</p>

Together with

- ➤ Any grade of Oedema +/++
- MUAC <11.5 or WFH <-3Z score (6-59 months)</p>

- MUAC <13.0cm (5-9 years, MUAC <16.0cm (10-12 years or BMI for Age <- 3Z score (5-12 Years) or</p>
- Bilateral Pitting Oedema + and ++

With any of the following medical complications

- > Anorexia, poor appetite
- High Fever
- Hypothermia
- Vomiting
- Severe dehydration
- Severe anaemia
- Not alert, very weak, apathetic, unconscious, convulsions
- Moderate to severe skin lesions
- Difficult or fast breathing

Special Cases

- All infants >6 months and weighing <3kg or
- Infants <6 months who meet any of admission for NRU as follows
 - Visible severe wasting
 - Bilateral Pitting Oedema
 - WFH <-3Z score (for children ≥45cm</p>
 - Static weight or not gaining weight or losing weight for 3 consecutive weighing in GMP but not severely malnourished
 - Mother reports that she does not produce enough milk and infant is not gaining weight on breastmilk despite counseling on appropriate breastfeeding technique and attachment

- Infant if sick, feeble and not suckling well but not severely malnourished
- Referrals from OTP

Management

- Must be admitted to hospital
- Do a full assessment, especially looking for underlying disease such TB, HIV infection, etc.

Assess and treat common complications and other medical condition (as indicated in the CMAM Guidelines; 2012 Edition) such as:

- Hypoglycaemia
- Hypothermia
- Diarrhoea
- Dehydration
- Severe anaemia
- Skin lesions
- > Eye problems
- Oral Candidiasis

Nutrition Management

- Phase 1
 - F75 is used to stabilize
 malnourished children in Phase 1
 of treatment
 - Give 8 feeds of F75 per day (every 3 hours). This will assist to prevent hypoglycaemia and Hypothermia

- Give F75 to all children except infants <6 months where breastfeeding is being established. For these infants use F100-D except if Oedema is present
- Continue breastfeeding on demand

Preparation of F75

- Mix one packet of F75 with 500ml of cooled boiled water (see annex 5-3 in the CMAM Guidelines 2012 edition)
- If pre-packed F75 is not available use one of the recipes (Annex 5-12 in the CMAM Guidelines 2012 Edition)

Amounts to Give in Phase 1

- Amounts to give depends on the admission weight
- Keep the amount the same for all of Phase 1 even if the child is gaining or loosing weight

Transitional Phase

- Criteria to progress from Phase 1 to Transition Phase
 - The child has good appetite (easily finishes feeds)
 - Bilateral Pitting oedema is subsidizing
 - No serious medical complications

- Nutrition Treatment in Transition Phase
 - RUTF is introduced alongside F75 in preparation for Phase 2
 - Last for 2-3 days
 - The number of feeds, their timing and the volume remains the same as in Phase 1
 - Give the caregiver the daily ration amount of RUTF. The caregiver giver should give the child RUTF throughout the day
 - After every 24-hour period, note the quantity of RUTF consumed
 - Give the child safe drinking water with RUTF in between feeds
 - Every three hours provide Phase1 ration of F75
 - If the child is breastfed, breastfeeding should continue throughout the day and night
 - When the child finishes 50% of RUTF, reduce the volume to 50 % of the normal F75 ration
 - Stop F75 when the child is able to finish 75-100% of the daily RUTF ration
 - Give RUTF and water only. Observe for at least 24 hours eating RUTF to ensure he/she does not develop complications
 - If the child develops complications return him/her to Phase 11 and provide appropriate medical care.

Discharge Criteria from NRU to OTP

- Appetite returned
- Medical condition resolved or stabilized (for chronic conditions).
- Bilateral pitting oedema subsidizing

Transition Phase in Classical Approach

- Where RUTF is not available or cannot be used Classical or Traditional Approach should be used in transitional phase
- F100 is used in the classical transitional phase.
- The number of feeds, timing and volume given remains exactly the same as in Phase 1 (see annex 5.4 of the CMAM Guidelines 2012 Edition)
- Breastfeeding should continue on demand

Preparation of F100

- Mix one packet of F100 with 500ml of cooled boiled water (see annex 5.3 in the CMAM Guidelines 2012 edition)
- If pre-packed F75 is not available use one of the recipes (Annex 5.12 in the CMAM Guidelines 2012 Edition

Criteria for change to Phase 2

- Transfer after 2 days in transition phase (if stable and no Oedema)
- If Oedema is present the child should remain in transition phase until all oedema has resoilved

Criteria of moving back to Phase 1

- If there is any sign of fluid overload
- If tense abdominal distension develops
- If the patient gets significant re-feeding diarrhea so that there is weight loss
- If a complication arises that necessitates an intravenous infusion
- If the child loses his/her appetite for therapeutic milk and requires NG feeding

Phase 2 (Rehabilitation Phase)

- F100 (High energy milk) is used in this phase
- Children should be given unlimited intake of F100
- Give 6 feeds of **F100** per day
- Offer 2 feeds of CSB++
- Breastfeeding should continue on demand as usual
- Give F100 milk according to weight (see annex
 5.6 of CMAM Guidelines 2012 Edition)
- Encourage the child to eat until full

Routine Medical Treatment and Prophylaxis

Vitamin A: (on day 1, Day 2) as follows:

| AGE | Vitamin A orally on day 1 and 2 (and |
|-----------|--------------------------------------|
| | day 14 if the child stays in NRU) |
| <6 months | 50, 000 IU (2 drops or one third red |
| | cap) |
| 6 to 12 | 100, 000 IU |
| months | |
| >12 | 200,000 IU |
| months | |

Antibiotic Regime in NRU

| Antibiotic Regime III 1410 | | | |
|----------------------------|----------------------------------|--|--|
| On admission if | Give | | |
| No complications | Amoxicillin orally, 15mg/kg tds | | |
| | for all of Phase 1 and four days | | |
| | extra | | |
| Complications | Give Benzyl penicillin | | |
| | 50,000iu/kg q6h IV/IM for 48 | | |
| | hours then oral amoxicillin | | |
| | 15mg/kg tds for 5 days AND | | |
| | If the child fails to improve | | |
| | within 48 hours add | | |
| | Gentamycin 7.5mg/kg q24h | | |
| | IV/IM for 7 days or | | |
| | Chloramphenicol 25mg/kg | | |
| | IM/IV q8h for 5 days | | |
| | Other antibiotics based on | | |
| | specific diagnosis | | |
| If child is HIV | Cotrimoxazole: | | |
| infected or | <6 months-120mg/day | | |
| exposed give | >6 months-5 years 240mg od | | |
| cotrimoxazole | >5 years-480mg od | | |
| preventive | | | |
| therapy (CPT) | | | |

- Assess dehydration (see Section 7.5.1.4)
 - Skin turgor is not a reliable sign in these children
 - Look for tears, sunken eyes or fontanelle
 - Assess urine output
 - Prevent and/or correct dehydration (see Section 6.1)

- The fluid tolerance of these children is limited. ReSoMal and especially IV solutions may cause fluid overload and heart failure
 - ✓ Avoid fluid overload in severe malnutrition
- Non-dehydrated children with mild diarrhea should continue milk feeds to prevent dehydration – they do not need ReSoMal
- Whenever possible, rehydrate these children orally using ReSoMal (to prepare this see above)
- If IV rehydration is necessary, limit this to a few hours.
 Continue feeding and rehydrate orally as soon as possible see above
- Start an intensive feeding regime with high energy milk:
 - ✓ Administration via NGT is necessary in many children with poor appetite
 - ✓ Start with 2-hourly feeds then reduce to 3-hourly

- ✓ Start with **F-75** refer to severe malnutrition manual
- ✓ Move to F-100 when the child is stable refer to severe malnutrition manual

Note: Frequent feeds spaced throughout the whole 24 hours are essential to prevent hypoglycaemia

- Provide antibiotic cover for 5 days
- ➤ Give **Cotrimoxazole** 24 mg/kg bd Alternatively if sepsis suspected or child very ill:
 - Give Benzylpenicillin 50,000 units/kg IM or IV q6h, and
 - Give Gentamycin 6 mg/kg IM or IV q24h

Alternatively to Gentamycin:

Give Chloramphenicol 25 mg/kg IM or IV q8h

If the child improves, then switch to oral **Ciprofloxacillin** for 5 days

Give supplements:

- Give Vitamin A (Retinol) 200,000 units on days
 1,2 and 8
 - Children < 1 yr: 100,000 units (½ capsule)</p>
- Give Multivitamin syrup 5 mL daily for 1 week
- Give Potassium 1 mmol/kg qid for 2 weeks mixed with feeds
 - ➤ To prepare a stock solution: add 7.5g of **Potassium Chloride** to 1L of pure water. This gives 1 mmol potassium chloride/mL

One Potassium Chloride slow-release (Slow K ® tablet = 13 mmol of K+)

From day 7:

- Give Ferrous sulphate paediatric mixture 2.5 mL bd for 2 weeks plus
- Give Folic acid 5mg od for 5 days
- Give Albendazole 400 mg single dose, when recovering
 - Children<2 give 200 mg</p>
- Treat complications:
 - > Hypothermia:
 - ✓ Re-warm
 - ✓ Consider the possibility of sepsis or hypoglycaemia
 - Hypoglycaemia:
 - ✓ Give Dextrose 50%
 - ✓ See Section 5.4 for dilution, dose, administration
 - ✓ Then give F75 orally or via NGT as soon as possible and recheck the blood sugar after 1 hour

Note: Hypothermia and hypoglycaemia are frequently signs of sepsis. Consider sepsis treatment if present

- Cardiac failure:
 - ✓ Give Frusemide 1-2 mg/kg IV or IM
 - ✓ Digoxin is contraindicated in kwashiorkor

- Severe anemia:
 - ✓ Transfuse 10 mL/kg packed cells
- Mouth ulceration:
 - ✓ If not severe use **GV Paint**If severe like cancrum oris use:
 - ✓ Give Benzylpenicillin 25,000 units/kg per dose IM q6h and
 - ✓ Give Metronidazole 7.5 mg/kg every tds for 7 days
- > Skin ulcers:
 - ✓ Soak lesion with Potassium Permanganate 1% solution for 10-15 minutes then
 - ✓ Apply a paraffin Gauze dressing

23.4 Pellagra

 Usually multiple vitamin deficiency is present and other vitamins may therefore be necessary

Treatment

 Give Nicotinamide 50 mg every tds for 28 days

23.5 Vitamin A Deficiency, Xerophthalmia

Prevention

 Give Vitamin A is a prophylactic vitamin A supplementation to all children 6 months-5 years old (every 6 months) nursing mothers and to risk groups at every available opportunity **Note:** Xerophthalmia is a medical emergency

Treatment

Adults

 Give Vitamin A 200,000 units/ dose on days 1, 2, and 8

Children

• Give **Vitamin A** < 1:100

24.0 Palliative Care

Palliative care is an approach that improves the quality of life of patients and their families facing the problems associated with life threatening illness, through the prevention and relief of pain and suffering by means of early identification and impeccable assessment and treatment of pain and other problems – physical, psychosocial and spiritual

24.1 Pain Management in Palliative Care

24.1.1 Areas to Consider in Pain Management

- Effective pain control requires:
 - Holistic assessment including psychological, spiritual, and social aspects
 - Acurate diagnosis of the cause: defined as somatic and neuropathic
 - Appropriate analgesics: Non opiod, opioids and adjuvants
 - To consider:
 - ✓ Radiotherapy if available
 - ✓ Chemotherapy if available

- ✓ Surgical / orthopeadic procedures
- ✓ Neurolytic procedures if available

24.1.2 Principles of Analgesic Use

- By the clock
 - Regular analgesia is necessary as patients require prevention therapy towards their persistent pain
 - Medication given when required for chronic pain does not work
- By the mouth
 - Oral medication is the standard preferred in chronic pain
- By the patient
 - Dosage is determined on an individual basis
- By the ladder
 - Three step analgesic ladder with treatment moving up as pain increases should be used when prescribing analgesics

24.1.3 The WHO Analgesic Ladder

 The choice of analgesic drugs should be based on severity (intensity), type (neuropathic or nociceptive) and cause of pain



Mild Pain - Step 1

- Give Paracetamol 500mg 1g every 4-6 hours (max 4g daily)
- Give NSAIDS anti inflammatory effects are good for metastatic bone and soft tissue pains
- Give **Aspirin** 300-600mg every 4-6 hours
- Give Ibuprofen 400mg P.O every 6-8 hourly (max 1.2 g daily), children>7kg 20mg/kg (max 40mg/kg/day)
- Give Diclofenac (NSAID) 50 mg, PO 8hourly (max 150mg)

Notes:

- Do not give Aspirin to children under 16 years because of the risk of Reye's syndrome
- Aspirin causes gastric irritation and ulceration, therefore administer with food and milk
- Do not use Aspirin or other NSAIDs (e.g. Ibuprofen, Indomethacin, Diclofenac) in patients with symptoms suggesting gastritis or peptic ulcer disease, pregnancy or bleeding disorders or in asthmatic patients
 - ✓ Do not use two **NSAIDS** at the same time

Moderate Pain - Step 2

- Codeine phosphate 30-60mg every 4 hours (max 240mg daily), children >1 year 3mg/kg in divided doses every 4 hours
- Can be combined with non-opiates and/or adjuvants, but <u>not</u> with morphine
- Give **Tramadol** 50-100 mg every 8 hours oral
- Always prescribe codeine with a laxative (not tramadol) e.g. Give Bisacodyl 10mg at night, unless the patient has diarrhoea

Severe Pain - Step 3 (strong opiod)

- Give Morphine a drug of choice for severe pain
- Can be sedalong with non-opiods for severe pain

- Oral Morphine (1 mg/ml) solution starting dose 2.5ml -5mls, 4 hourly and double dose at night (10pm) titrating upwards every 12 hours to achieve pain control
- Patients who were previously on Codeine should start on morphine 10mgs po 4 hourly
- For children, 0.2- 0.4mg/kg po 4 hourly (max 3mg)
- Cachexic patients start Morphine with a low dose 2.5mg 4 hourly
- Elderly / frail patients start
 Morphine at 2.5mg po 6-8 hourly
 due to potential risk of renal
 imparment
- Where Oral Morphine Tablets (MST) is available start with 10mg tablets bd and titrate upwards (take 24-28 hours to reach steady state level)
- May be given together with step 1 drugs (nonopioids) and/or adjuvants
- Always prescribe a laxative e.g. Bisacodyl
 10mg at night unless the patient has diarrhoea
- Nausea and vomiting are common side effects and may require medical management if persist after 24 - 48 hours
- Morphine does not have a ceiling effect and must be titrated up to gain effective control of pain

- Increase Morphine dosage by 30-50% of the normal dose, for maximum pain control
- For breakthrough pain, give the regular dose of Morphine as a "rescue" dose as often as required and keep record
- Initiate Weak Liquid Morphine (1mg/ml) and if the required dose is very high can be changed to Strong Morphine (10mg/ml) 4 hourly
- If patient cannot swallow, Liquid Morphine can be administered through bucal and rectal
- Use fentanyl patches if available as soon as pain is under control for maintainance

Note:

- Never prescribe more than one modified release opioid at a time
- Explain common opiod side effects to patients (constipation, drowsness, nausea)
- If pain has been controlled reduce the dose gradually to avoid withdraw symptoms (sweating nausea, agitation)
- For painiful procedures (dressing, fructure reduction), give **Morphine** 5mg start (adult) or 0.2- 0.4mg/kg (children) 3- 40 minutes before procedure to reduce pain and distress. If the procedure is repeated you can titrate the dose to the best effect.

- For regional anaesthesia (epidural, caudal, or specific LA blocks) and chronic pain relief can give bupivacaine 0.1% 0.5% (preservative free), ropivacaine (preservative free) 0.1%. For management of local anaesthetic toxicity refer dental section
- Pethidine is not recommended for use in chronic pain due to its short duration of action and its side effects

Adjuvant Analgesics

- Drugs useful in pain that is only partially sensitive to opioids, such as nueropathic and bone pain, smooth or skeletal muscle spasms, or pain related to anxiety
- Can be used alone, or in conjuction with step 1,
 2 and 3 analgesics

Corticosteroids

- Can be helpful in reducing tumour related
 Oedema e.g. liver capsule pain, nerve
 compression
- Combination of a steroid and opioid is usually effective
 - ➤ Give **Dexamethasone** 4 8mg in divided doses daily for a minimum of 10 days
 - For increased intracranial pressure start 24mg daily and reduce by 2mg daily to the lowest efffective dose

- For nerve compression 8mg daily
- ➤ For spinal cord compression 16 mgs

 If dexamethasone not available
 - Give Prednisolone 30 50mg od for 10 days
 - Then reduce gradually to lowest
 - effective dose, depending on prognosis
 - In advanced disease, corticosteroids can improve appetite, decrease nausea and improve quality of life

Monitor side effects such as:

- Neuropsychiatric syndrome
- GIT disturbance
- Immuno suppression

Anti-depressants

 Helpful for neuropathic pain, which may present as burning, pricking, allodynia, paraesthesia or sharp, shooting pain

For adults

• Give **Amitriptiline** 25mg at night. Dose can be increased slowly up to 75mgs

For children

- 2-12 years: 0.2 0.5mg/kg po at night
- 12 -18 years: 10-25mg/kg po at night

14. Hiccup 1.1

This is a relatively common symptom in advanced disease which causes distress to the patient

| | 1.2 Common Causes | 1.3 | Management |
|--|--|---------------------------|---|
| | | | |
| | Irritation of the phrenic nerve by tumour | Immediate: | 22 |
| | involvement at the hilum of the lung. | | |
| | Direct irritation of the diaphragm (infection, | Phary | Pharyngeal stimulation e.g. by swallowing dry bread or two |
| | tumour). | teaspo | teaspoons of sugar. |
| | Uraemia. | Correct | Correct uraemia if possible. |
| | Dyspepsia (especially with hiatus hernia). | ■ Simple | Simple re-breathing from a paper bag to elevate pCO2 level. |
| | Elevation of diaphragm (from enlargement of | Drugs: | |
| | the liver or ascites). | (i) | (i) Haloperidol 0.5mg bd orally or 1.5mg intramuscularly |
| | CNS tumour. | dur | during attacks. |

(iii) Metoclopramide 10mg qid or domperidone 10-20mg (iv) Muscle relaxants e.g. baclofen 5-10mgs TDS are (although its sedative effect may distress the patient). qid may be quite effective if due to gastric distension.

(ii) Chlorpromazine 12.5- 25mg bd. or prn during attacks

Anti – convulsants

Adults

- Give Phenytoin 100mg bd, maybe increased slowly to 100mgs tds
- Give Carbamazepine 100 200 mg tds

Children

2-12 years 10/kg day 1, then bd day 2, tds day
3

24.5.3 Antispasmodics

- Helpful in relieving visceral distension pain and colic
- Give Hyoscine butylbromide 10-20mg every 8 hours tds orally

Children

- 1 month 2 years give 0.5mg/kg po 8 hourly
- 2-5 years give 5mg po 8 hourly
- 6-12 years give 10mg po 8 hourly

24.5.4 Muscle Relaxants / Anxiolytics

 Helpful in painful muscle spasms (cramps), myofascial pain and anxienty related pain

Adults

• Give **Diazepam** 2.5mg every 8 hours

Children

• 6 - 14 years give 2-10mg/day in 2 or 3 divided doses

24. 5 Management of Common Symptoms in Palliative Care

General Approach

- Assess cause and severity
- Treat reversible causes
- Initiate disease specific palliative care (drug and non-drug measures)
- Involve and explain the plan to patient and family
- Review

1.1 16. Halitosis - an unpleasant or bad breath

Gingivitis (gum infection), sepsis in the mouth, pharynx, nose, nasal sinuses, lungs tonsillar disease, lung abscess/cancer Common causes: Poor dental and oral hygiene, Oral thrush, Dry mouth – loss of saliva, mouth breathing, dehydration, and necrotic ulcers

1.2 Management

1.3 General Measures

Good oral hygiene

- The patient may be unable to do this for himself,
- Cleaning with soft tooth brush
- Rinse the mouth with salty water or diluted lemon juice
- Encourage oral fluids, chew or suck small pieces of fruit (pine apple or orange)
- Mouth wash and paste- crushed flagyl + sobo+crushed steroids (predinisolone) to reduce odour and pain

L 23. Squashed Stomach Syndrome

This is a reduction in stomach capacity leading to a feeling of fullness after taking only a small amount of food.

- .2 Clinical Features
- Early satiety / constant feeling of fullness
- Oesophageal regurgitation
- Indigestion

Common causes

Liver enlargement, gross ascites, pelvic tumours, e.g. ovarian, bowel

Management

- Small frequent meals
- Antacids to remove gas, thus increasing stomach capacity
- Antiemetics, e.g. Metoclopramide 10mg tds, taken 30 mins before meals, to control nausea and speed gastric emptying
- Steroids to reduce tumour size

| 1.1 18. Anorexia - loss of appetite | |
|---|--|
| 1.2 Common Causes | 1.3 Management |
| Cancer | Explain to the patient and family that anorexia is normal as disease |
| Opportunistic infections | progresses |
| Pain | Identify and treat underlying causes, if possible |
| Weakness / fatigue | Encourage small, frequent meals of the type of food which the |
| Treatment side-effects – antibiotics, chemotherapy | patient likes |
| Gastrointestinal problems – sore mouth, dysphagia | Present food attractively |
| Anaemia | Cook food away from where the patient is, to reduce cooking smells |
| Changes in taste sensation – common in cancer and HIV | Don't force feed |
| Psychological causes – depression, anxiety, fear | Encourage the patient to eat with other family members – appetite is |
| Disease process – anorexia increases as disease | usually better when meals are eaten in company |
| progresses | Encourage the patient to sit in an upright position when eating |
| Unappetizing food, poorly presented | Encourage gentle exercise, to promote appetite |

Appetite stimulants – eg. Prednisolone 10mg tds, before meals; small

Drugs, e.g. antiemetics to reduce nausea, Metoclopramide to speed

Generally contraindicated in advanced cancer.

up gastric emptying.

amounts of alcohol

Strong food smells during cooking – may cause nausea

Management of Nausea and Vomiting in Palliative Care in Adults

Treat

- Oral/oesophageal candidiasis
- Constipation
- Infections malaria, gastroenteritis, urinary tract infection etc.,
- Indigestion/heartburn

Care

- Review new medication that may have caused nausea/vomiting
- Encourage clear fluids small sips better absorbed
- If the patient is dehydrated give ORS
- Ginger chewed or boiled as a drink may help

Prescribe

- Nausea and vomiting can arise from many different causes and respond to particular drugs – see table below
- If vomiting is severe, injectable anti-emetics are best until the vomiting is controlled

| Pattern of nausea and vomiting | Causes | înS | Suggested drugs |
|--|--|---------------|-----------------------------------|
| Poor stomach emptying: | • Opioids | • Me | Metoclopramide |
| Main symptom is vomiting | Constipation | 10- | 10-20mgs tds before meals |
| Vomiting relieves nausea | Stomach & bowel | or | |
| Patient feels full quickly | conditions | • Dor | Domperidone 10-20mg bd |
| May have reflux | | | |
| Blood chemistry disturbances | • Drugs | • Hal | Haloperidol 1.5mg nocte or |
| Nausea is the main symptom | Renal failure | • Pro | Prochlorperazine 5-10mg |
| Vomiting does not relieve | Hypercalcaemia | tds | |
| nausea | | | |
| Inflammation or swelling in the head | Ear infections | • Pro | Prochlorperazine 5-10mg |
| May be worse on movement | Brain tumours | tds or | or |
| Vomiting does not relieve | Meningitis | • Cyc | Cyclizine 25-50mg tds |
| nausea | Malaria | | |
| May be worse in the morning | | | |
| Vomiting with diarrhoea | Infectious diarrhoea | • Pro | Promethazine 25mg tds or |
| Exclude constipation with | | • Cyc | Cyclizine 25-50mg tds |
| overflow | | | |
| Partial bowel obstruction | Constipation | • Me | Metoclopramide 10-20mg |
| Large volume vomiting | Abdominal or pelvic | ⊠ | IM. STOP if increasing |
| Patient still passing occasional | tumour | apo | abdominal pain and |
| flatus/stools | | pre | prescribe as below |
| Complete bowel obstruction | Abdominal or pelvic | • Pro | Promethazine 25mg s/c tds |
| Large volume vomiting | tumour | or | |
| Patient not passing flatus or | | • Cyc | Cyclizine 50mg s/c tds |
| faeces | | | |

Chapter 25: Psychiatric Conditions

These are the common conditions covered in this chapter:

- Organic mental disorders delirium and dementia
- Alcohol and drug use related disorders
- Psychotic disorders schizophrenia
- Mood disorders depression and bipolar affective disorder
- Anxiety disorders
- Mental disorders during pregnancy and postpartum period

25.1 General Guidelines on Psychotropics Use

- Identify complicating factors i.e. psychological, medical, social, etc.
- Review past successful medication and other interventions
- Weigh the risks and benefits of any intervention
- Involve the patients and their families in all decisions around treatment
- Target symptoms e.g. use a more sedating agent if insomnia is a problem
- Start with low doses and go slow
- Avoid poly-pharmacy

- Never stop medication suddenly. Wean off slowly
- Take caution in special circumstances such as:
 - Pregnancy and breast feeding mothers
 - Elderly patients
 - > HIV Infection
 - Renal or hepatic impairment
 - Medicine interaction
- Psychosocial interventions are imperative

25.2 Organic Mental Disorders

25.2.1 Delirium

 Sudden onset of confusion often accompanied by impairment of consciousness and agitated behaviour often caused by underlying organic conditions (e.g. infections) and usually reversible

Signs and Symptoms

- Confusion
- Agitation
- Perceptual disturbances
- Possible physical complaints such as headache or fever
- Fluctuating level of consciousness
- Disorientation
- Raised temperature
- Sweating
- Hallucinations and delusions

Investigations

- Look for possible causes:
 - Systemic and CNS infections
 - Hypoxia
 - Hypo- or hyperglycaemia
 - Drugs
 - > Alcohol intoxication or withdrawal
 - Post-convulsion phase in epilepsy
 - > Head trauma
 - Subdural hematoma
 - Stroke etc.

Full physical examination including vital signs

- Full blood count
- Urine/ Blood glucose test
- Blood film or rapid diagnostic test for malaria
- Consider LP for CSF analysis if suspect meningitis or encephalitis
- Advisable to determine HIV serostatus

Treatment

- The aim of the treatment is to control the behavioural disturbances in order to allow identification and management of the underlying cause (e.g. infections, alcohol withdrawal, hypoglycaemia etc)
- For behavioural disturbances or aggression give short course of low dose of antipsychotic medications, preferably orally

- Ist line: Chlorpromazine 50 100mg bd or Haloperidol 2 - 5mg bd for 7 days or until agitation / confusion resolves
- ➢ If the patient is refusing oral medications, use Chlorpromazine 50 − 100mg IM bd or Haloperidol 2.5 − 5mg IM bd until sufficiently improved to accept medications orally
- ▶ If agitation is severe consider adding Diazepam 5 – 10mg po or slow IV push for short time or until less agitated (3-5 days)
- Nurse the patient in calm, quiet and well-lit environment with frequent reassurance and orientation

Red Flags

- For referral
 - Decreasing level of consciousness
 - Worsening physical health
 - Investigations required not available at health facility
- For admission
 - If underlying condition and behavioural disturbance cannot be managed at home or health facility
 Note: Intravenous or high dose
 Diazepam can cause respiratory depression/ distress

Follow-up

If agitation is severe review the patient regularly

- As an outpatient review at least every 24-48 hours until stable
- If admitted review every 8 hours until stable
- Once symptoms have resolved, stop antipsychotics and arrange review in 5-7 days' time to ensure patient remained stable

25.2.2 Dementia

Signs and Symptoms

- Forgetfulness
- Deterioration in personal hygiene and social interaction
- In later stages:
 - Wandering away from home
 - Poor memory of recent events
 - Disturbances of orientation
 - Personality changes such as irritability, and suspiciousness (sometimes persecutory delusion)

Investigations

- Formal memory tests can also be used; for example, ask patient to:
 - ➤ Give the names of 3 common objects and repeat them after about 3 minutes
 - Accurately identify the day of the week, month, year etc (as appropriate)
 - Give their full name and where they live

Any impairment symbolises possibility of dementia, then

- Do physical examination to look for possible causes such HIV, high BP, alcohol, vitamin deficiencies etc.
- Assess for depression which can cause confusion in elderly

Treatment

- The aim of treatment is supportive
- Maintain patient's physical health good diet exercise, personal hygiene
- Treat any physical illness promptly
- Encourage patient to use remaining abilities a much as possible
- Use reminders and prompts to help memory
- If depressed, refer to depression section for treatment details
- If the psychotic symptoms are prominent, use low dose antipsychotic medications
 - ➤ 1st line: **Haloperidol** 1.25-2.5mg po od/bd until the symptoms resolve or
 - 2nd line: Risperidone 0.5 1mg po od/bd until the symptoms resolve
 - Consider long term antipsychotic medication use if the psychotic symptoms persist after stopping the medication.
 - If possible, avoid use of medication especially sleeping tablets (e.g. Diazepam) as they make confusion worse. Aspirin in low doses may slow down vascular dementia

Red Flags

- Admission
 - If there is sudden increase in confusion in known cases of dementia which may be due to acute infection, toxic reaction to medication, acute psychosis, misuse of alcohol or drugs (rule out Delirium or Acute Psychosis)
 - If the possible cause is treatable or manageable diseases such as HIV, High blood pressure

Follow-up

- Consider reviewing the patient once a week until stable
- Once the condition is stable, review once a month
- Always do a physical check up to rule out comorbid illnesses which can complicate the dementia
- Assess risk for committing suicide
- Consider stopping antipsychotics if symptoms resolve
- Consider continuing with low dose antipsychotic medications if the psychotic symptoms persist
- Assess for and treat moderate severe depression in the carers (guardians)

25.3 Alcohol Related Disorders

25.3.1 Alcohol intoxication

 Behavioural change with disinhibition, potentially agitated and aggressive behaviour after recent ingestion of alcohol

Signs and Symptoms

- Argumentativeness
- Lability of mood
- Impaired attention and judgment, and interference with personal functioning
- Unsteady gait
- Difficulty standing
- Slurred speech
- Decreased conscious level
- Flushed face
- Conjunctival injection

Investigations

Blood or breath alcohol (if available)

Treatment

- Management is primarily supportive until the effects of the alcohol have worn off
- If very aggressive then follow treatment guidelines for Violence and Aggression management

Note: Diazepam should be avoided as there is increased risk of respiratory depression

Red Flags

- Admission
 - If underlying condition and behavioural disturbance cannot be managed at home or health facility

Follow-up

 Give advice about safe levels of alcohol intake and screen for alcohol use disorders (CAGE questionnaire)

25.3.2 Alcohol Withdrawal Syndrome

- Acute confusional state that occurs within hours to days of cessation, or reduction of alcohol intake after prolonged (weeks to months) of heavy consumption
- The peak onset is at 24-48 hours post last ingestion and it can last for 7 – 10 days if untreated

Signs and Symptoms

- Headache
- Nausea
- Anxiety
- Fevers
- Shaking
- Tremor
- Sweating
- Vomiting
- Increased pulse and blood pressure
- Agitation

 Can present as Delirium Tremens with: confusion, marked agitation, aggression, hallucinations (frequently visual) and delusions, seizure

Investigations

- Full physical examination including vital signs (to exclude other causes of delirium if present)
- Consider FBC LFT
- Blood/ Urine glucose

Treatment

- The aim of treatment is to reduce the symptoms associated with alcohol withdrawal, which can result in seizures and potentially be fatal
- Primary treatment
 - A short course of oral **Diazepam** should be given at least four times a day, reducing in dose over a week to ten days, titrated according to symptom resolution; for example:
 - ✓ Diazepam 20mg qid for 2 days
 - ✓ Diazepam 10mg qid for 2 days
 - ✓ Diazepam 5mg qid for 2 days
 - ✓ Diazepam 5mg bd for 2 days

- It should be accompanied by oral
 Thiamine supplements
 - ✓ Thiamine 300mg OD orally for 1 month
- Secondary /alternative treatment
 - If markedly agitated and unable to comply with oral medication IV
 Diazepam 5-10mg can be used up to 4 times per day until able to comply with oral treatment
 - ✓ High dose IM / IV thiamine can be given if available
 - ✓ IV fluids may be required if evidence of dehydration (low BP, tachycardic)

Red Flags

- For admission:
 - Delirium tremens should be treated as an inpatient (medical emergency)
 - People with a high risk of seizures (previous seizures, known epilepsy, prolonged heavy alcohol use)
 - People with co-morbid physical illnesses (HIV, jaundice)

Note: The dose of **Diazepam** should be reduced in the physically frail of those with liver impairment

Follow-up

- If agitation is severe review the patient regularly
- As an outpatient review at least every
 24-48 hours until stable
- If admitted review every 8 hours until stable
- Once detoxification complete offer advice regarding safe levels of alcohol intake and counseling support if planning to stop drinking

25.3.3 Wernicke -Korsakoff Syndrome

- Is characterized by confusion and drowsiness, ataxia and ocular disturbances (usually due to weakness or paralysis of 6th cranial nerve) including nystagmus
- May have acute onset or develop slowly over 1 week or so
- Korsakoff's psychosis is a state of amnesia that usually follows wernickes' syndrome
- This is due to thiamine (vitamin B1) deficiency in alcoholics and malnourished non-alcoholics

- Mostly anterograde amnesia (inability to retain new memories) and possibly retrograde amnesia (inability to recall the past),
- Fabricating answers or confabulating to cover their memory problems, and
- Oculomotor disturbances nystagmus.
- Patient is alert but can be confused if having wernicke's disease, responsive and normal

Investigations

- Substance use history
- String test to diagnose Korsakoff's psychosis (clinician asks the patient to take an imaginary string in his or her hands, and the patient complies, as though the string were real)
- Do a physical examination to rule out medica complications or comorbid illnesses
- Laboratory tests include FBC, offer PITC for HIV, LFTs, Urine/ Blood glucose test

Treatment

 The aim is to rapidly treat Wernicke's syndrome in order to prevent onset of Korsakoff's psychosis

- Immediately give Thiamine (Vitamin B1) 100 mg IV or IM followed by 100 mg IM or orally od for the next 2 days for complete reversal of ocular abnormalities and ataxia
- Give Glucose (5-10%) IV with multivitamins orally and/or Vitamin B Complex can be given
- Glucose without Vitamin B1 can worsen
 Wernicke's encephalopathy
- Advise the patient to abstain from alcohol use or consider alcohol detoxification to prevent alcohol withdrawal
- Always treat medical complications of chronic alcohol use such as GIT, neurologic, cardiovascular, pulmonary, hematologic and endocrine.
- To treat delirium tremens: see alcohol withdrawal section

Red Flags

- For Admission
 - If delirium tremens or alcohol withdrawal is suspected
 - Severe medical complications of chronic alcohol use such as vomiting blood
 - Continued high risk alcohol use or multiple substance use

Follow up

- Review the patient once a week until the confusion resolves
- Assess for continued alcohol use and offer brief intervention to promote change in alcohol use on each visit
- Assess for and treat the medical complications

25.3.4 Alcoholic Hallucinosis

 Is a substance-induced psychotic illness which is a rare complication of prolonged heavy alcohol use, and often starts during a phase of abstinence.

Sign and Symptoms

- Perceptual disturbances
- Signs: auditory hallucinations in clear consciousness and while sober
- If insight is not present, may form basis for delusional thinking

Investigations

 Comprehensive alcohol use history from patient as well as guardians

Treatment

- The aims of treatment are abstinence and supportive care
- It is a self-limiting condition, symptoms resolve rapidly after ceasing alcohol consumption after a few months
- If symptoms persist for more than 6 months, consider giving antipsychotics (refer to the substance induced psychosis section)

Red Flags

- For admission
 - Persisting symptoms for more than 4-6 weeks of antipsychotic medication trial
 - Continued risky alcohol use
 - Severe medical complications
 - Emergence of more typical schizophrenia symptoms

Follow-up

- Assess for emergency of schizophrenia symptoms on each visit once a month
- Assess for continued alcohol use and offer brief intervention
- Assess for and treat medical complication for chronic alcohol use

25.3.5 Management of Physical and Neurological Complications of Alcohol Dependence

- Counsel the patient
- Abstinence may be essential
- Refer to Alcoholic Anonymous groups if available or link with other agencies such as religious organizations, social welfare services, etc
- Encourage a healthy diet with high protein and vitamin content (give Vitamin B Complex if necessary)
- Treat specific disorders symptomatically (e.g. gastro-intestinal disorders, cirrhosis, neuropathy) as per guidelines

25.3.6 Treatment of Psychological and Social Complications

- Counsel the patient using problem solving technique
 - Identify the problems
 - Prioritize the problems
 - Select the problem to be addressed
 - Think about all possible solution to the problem
 - Select the most appropriate solution
 - > Implement the solution
- Educate and support the family

25.4 Anxiety disorders

- May be a perfectly normal response to stressful life events or circumstances
- Only illness if no obvious stress or threat, or worse than the situation warrants
- Types:
 - Panic disorder: characterized by spontaneous panic attacks
 - Generalized anxiety disorder: excessive worry about actual circumstances/ events

- Phobias: irrational fear of objects/ public situations
- Post traumatic stress disorder: anxiety produced by extraordinary stressful events, re-occurs as flashbacks.
- Obsessive compulsive disorder: recurrent intrusive ideas, images thoughts or repetitive ritualized patterns of behaviour

Signs and Symptoms

- Feeling tense and anxious and worry a lot about things
- Associated physical health symptoms, heart palpitations
- Sweating, chest pains or difficulty breathing, feeling dizzy, lightheadedness or faint

Investigations

- Perform a physical examination to rule out other causes of anxiety such as thyrotoxicosis, asthma, etc.
- Elicit psychosocial stressors including alcohol and drug use from patients as well as guardians (carers)

Treatment

- Explain anxiety to patient especially the link between physical and psychological symptoms
- Give good health advice such as reducing or stopping substance use, good diet, etc.
- Educate on relaxation methods such as breathing exercise during panic attack

- If symptoms are severe prescribe
 - 1st line: Give **Diazepam** 2-10mg 2-4 times daily po for 5-7 days
 - 2nd line: Give **Amitriptyline** 50-150mg nocte po for 4-6 weeks
 - 3rd line: Give **Fluoxetine** 20-60mg od (morning) po for 4-6 weeks
- Psychotherapy such as cognitive behavioural therapy (CBT) if available

Red flags

- For referral/admission
 - If there is significant comorbid physical illness
 - The symptoms are interfering with activities of daily living
 - The patient is suicidal with no significant psychosocial support at home

Follow up

- Review patient once a week until symptoms resolve
- Once symptoms resolve, continue treatment for
 6 months and consider stopping thereafter
- Assess for substance use and other psychosocial stressors

25.8 Psychiatric Emergencies

25.8.1 Self-harm and Attempted suicide

- An act of self-harm without suicidal intent is deliberately harming oneself, e.g. cutting oneself. A suicide attempt is an act of selfharm with suicidal intent but not resulting in death
- Both are to be taken very seriously because they are high risk factors for completed suicide in the future
- Risk of future attempt is raised if:
 - Underlying psychiatric disorder (depression, bipolar affective disorder, schizophrenia/ drug or alcohol misuse)
 - Ongoing suicidal thoughts or plans
 - Regret at having survived the attempt
 - Access to dangerous means (firearms/ agro-chemicals/ medicines etc)
 - Evidence of hopelessness, marked emotional distress
 - Previous self harm/ suicide attempts
 - > Family history of completed suicide
 - Male gender, older age, lack of social support

Sign and Symptoms

 Evidence of injuries from or history of actual suicide attempt method e.g. hanging, poisoning, drowning, etc.

General Management

- Asking about suicidal thoughts in a routine assessment
 - First establish a good relationship
 - Ask how they feel about their life at the moment. How do they see the future for themselves? Do they ever think that life is not worth living?
 - Do they have thoughts about trying to harm them-self or end their life? If yes, have they made any plans harm themselves or end their life?
 - What has prevented them from acting on these thoughts? What protective factors are in their life? (family/ religious faith/ hope that things will get better)
 - Have they made at attempt to harm them-self end their life in the past? If yes, what happened?

Treatment

- If assessing someone presenting with a medically serious attempt at self-harm, first treat the underlying condition whilst ensuring that they are in a safe environment e.g. suture any wounds and manage bleeding, monitor appropriately if ingested poisons such as rat poison/fertiliser/ overdose of prescribed or over-the-counter medication
- All people with a self-harm/ suicide attempt should be referred for assessment by the district mental health team or local psychiatry services before leaving the health facility
- If there is current active suicidal ideation with a plan and access to dangerous methods, the person should be referred for assessment and possible admission by the psychiatry team urgently
- If suicidal thoughts are present but there is no plan and protective factors are in place, treat any underlying psychiatric disorder, give supportive counselling and monitor the suicidal thoughts at follow up visits. If the suicidal ideation does not improve, refer for assessment by the psychiatry team

25.8.2 Acutely Disturbed or Violent Behaviour Management

- The vast majority of people with psychiatric disorders are never aggressive or violent
- However some factors do make it more likely that some people may become aggressive when unwell (active psychotic symptoms, agitation and over activity associated with mania, auditory hallucination, confusion and disorientation, alcohol or drug use)
- Other people attending health care facilities can also be aggressive or violent for a variety of reasons, so it is important to know how to manage this situation in a way that is safe for the patient, their carers and the heath care staff
- Factors associated with violence or aggression
 - Young male
 - Previous or recent history of aggression
 - Drug or alcohol use
 - Increased impulsivity eg delirium, brain injury, learning disability, dementia
 - Psychiatric disorders e.g. schizophrenia with current active psychotic symptoms especially command hallucination or paranoid persecutory delusion, mania.

Signs and Symptoms

- Making verbal threats or shouting
- Agitation or irritability
- Suspiciousness/ anxious look
- Pacing up and down
- Actual physical aggression towards people or property

Investigations

Assess for biopsychosocial causes of the acute disturbed or violent behaviour

Treatment

- The aim is to alleviate suffering and to prevent harm/ injury to the patient and the health care staff
- Also to allow investigation and management of the underlying cause of the aggression e.g. delirium, psychosis, mania

General Measures

- De-escalation techniques and skills such as
 - First ensure your own safety Move towards a safer place – avoid being trapped in a corner; have other staff or guardians with you;
 - > Give clear, brief, assertive instructions
 - > Explain your purpose or intention
 - Negotiate options and try to understand the reason for their distres
 - > Avoid verbal and non-verbal threats
 - Allow greater body space than normal

25. Psychatric Conditions

- Pharmacological management (rapid tranquilization) may be optional if deescalation fails
- Have at least four additional people to handle patient if rapid tranquilization is needed
- If patient comes while tied do not immediately remove physical restraints until safe to do so

Steps of Rapid Tranquilization (RT)

| Step | Intervention | Dosages of | Other/ Adjuvant |
|------|---------------------|----------------------|-------------------------|
| | | Medication | treatment |
| 1 | De-escalation | | |
| 2 | Offer oral | Haloperidol 2.5- | With or without |
| | treatment | 5mg or | oral Diazepam 5- |
| | Repeat this up to | Chlorpromazine | 20mg or |
| | 2 more time at | 100-200mg | Promethazine |
| | 30-minute | | 50mg |
| | intervals if person | | |
| | remains agitated | | |
| 3 | Consider IM | Haloperidol 5mg | Promethazine |
| | treatment (if the | or | 50mg IM is an |
| | person doesn't | Chlorpromazine | alternative in |
| | accept oral | 50-200mg or | benzodiazepine- |
| | medication or is | Lorazepam 1-2mg | tolerant patients |
| | not effective. | if available | (people with |
| | These | | alcohol |
| | medications can | | dependence) |
| | be combined in | | |
| | one syringe & | | |
| | given into large | | |
| | muscle e.g. | | |
| | gluteus or deltoid | | |
| 4 | Consider IV | Diazepam 10mg | Note: if giving IV |
| | treatment using | slow push over at | Diazepam , care |
| | large vein | least 5 min. | should be taken |
| | | repeat after 5-10 | as it causes |
| | | min if insufficient | respiratory |
| | | effect (up to | depression: |
| | | three times) | patient could stop |
| | | , | breathing |
| 5 | Seek expert | | |
| | advice | | |

General monitoring after RT

- Pulse
- BP
- Respiratory rate
- Temp
- If available, oxygen saturation should be monitored every 15 minutes for the first hours and then every 30 minutes until the patient is awake and alert
- Full physical examination including vital signs to determine if any physical cause for the aggression (e.g. delirium, drug or alcohol withdrawal)

25.9 Psychotic Disorders

- This is an umbrella term for a group of conditions where the person thinking, emotions and behaviour are affected
- They experience hallucinations, normally auditory and have unusual beliefs, which are often paranoid about other people
- They can behave in unusual ways and lack insight into the fact that they are ill

25.9.1 Schizophrenia

Signa and Symptoms

- To diagnose schizophrenia there must be a one month history of symptoms listed below
 - Hallucinations (often auditory but can be visual etc) and delusions (fixed false belief) often paranoid/ suspicious in nature

25. Psychatric Conditions

- Speech can be irrelevant and incoherent
- Abnormal behaviour and at times agitation and aggression
- > Lack of insight is prominent

Investigations

- In a first episode of psychotic symptoms, physical causes for the symptoms must be excluded (delirium, drug and alcohol use/ withdrawal etc)
- Full physical examination
- FBC
- VDRL
- Urine drug screen if available
- Consider HIV serotesting.

Treatment

- Treatment is both biological (medications) and psychological (supportive counselling about the illness, compliance with medication, education to the guardians
- The aim of treatment is to remove all symptoms if possible and to help the person to return to their previous level of functioning

- Primary treatment
 - First line commence an anti-psychotic medication which will need to be continued for at least one year if this is a first episode or several years if the person has had many previous episodes
 - Give Chlorpromazine 100-300mg nocte or Haloperidol 2.5-5mg nocte
 - All anti-psychotic medication have a delayed onset of action – advice the patient/ guardian it will take 1-2 weeks before improvement is noted
 - Advise about the side effects:
 - ✓ Chlorpromazine: sedation, postural hypotension, constipation, photosensitivity
 - ✓ Haloperidol: Extra-pyramidal side effects (EPSE) eg. tremor, stiffness of limbs/jaw, salivation
 - ✓ This dose can be increased at subsequent reviews if symptoms remain present
 - ✓ If EPSEs persist consider reducing the dose of antipsychotic or adding Benzhexol 5mg daily

- Secondary /alternative treatment
 - If symptoms have not improved on chlorpromazine or if the person has lots of side effects use a second generation anti-psychotic Risperidone 1-2mg nocte
 - Advice about side effects:
 - ✓ Risperidone: weight gain, sedation, impaired glucose tolerance OR
 - ✓ If compliance with medication is poor despite trying to reduce side effects and counselling on the importance of compliance with medication consider a long-acting depot antipsychotic Fluphenazine 12.5mg IM into a large muscle (eg gluteal/ deltoid) as a test dose
 - ✓ Caution: risk of Acute Dystonic reaction (painful spasm of head and neck muscles)
 - ✓ If occurs give **Benzhexol** po if able to swallow
 - ✓ Otherwise IV/IM Procyclidine or Benzhexol 5mg OR slow IV push Diazepam 5-10mg
 - ✓ Advice about side effects: EPSE
 - ✓ Maintenance dose
 Fluphenazine 25-50mg IM
 every 4 weeks

✓ If EPSE persist consider reducing the dose adding Benzhexol 5mg daily

Red Flags

- For referral
 - If symptoms persist/ worsen despite 6-8 weeks of anti-psychotic medication at an effective dose
 - If side effects are not manageable
 - ➤ If the person has Catatonic symptoms (mute, maintaining unusual postures for many hours, stiffness/ rigidity despite not being on medication)
- For admission
 - Marked agitation / aggression should be managed as an inpatient following the Violence and Aggression treatment guidelines
 - Evidence of dehydration and malnutrition due to prolonged poor self care
 - Evidence that the patient is a risk to themselves (self harm/ neglect/ vulnerable to exploitation) or a risk to others (agitation/ aggression)
 - If insight is lacking and there is no guardian to ensure compliance with medication at home

Note: Neuroleptic Malignant
Syndrome is a severe but rare
complication of anti-psychotics,
presenting with fever, rigidity,
fluctuating pulse and BP and reduced
conscious level. It is a medical
emergency. All anti-psychotics should
be stopped and the person referred for
medical admission.

Follow-up

- If inpatient review the person every 24 hours until stable
- If outpatient review in 1-2 weeks initially and then every 4-8 weeks once more stable
- Screen for ongoing symptoms and monitor for side effects at each review
- Adjust the dose of medication accordingly
- Ask about any drug or alcohol use and give advise about use
- Screen for low mood and suicidal ideation at each review and follow Depression Treatment Guideline if present
- Once symptoms are improving advice the patient to return to their usual daily activities, including work if employed

Treatment duration:

- First episode: Medication should be continued at an effective dose for one year <u>from when all</u> symptoms are resolved
- Recurrent schizophrenia: Medication should be continued for 2-5 years from complete resolution of symptoms. If many episodes of relapse or symptoms never fully resolves may need to remain on medication lifelong.
- When considering stopping medication, discuss carefully with the patient and guardian and start to reduce slowly over 4-8 weeks
- Advise about symptoms that would indicate relapse (difficulty sleeping, auditory hallucinations, suspicious thoughts) and inform to return to the clinic promptly. Continue to monitor until medication free for 2-3 months before discharging
- Advise them to return to the clinic for review if have any concerns in the future

25.9.2 Substance Induced Psychosis

Development of psychotic symptoms

 (abnormal behaviour, hallucination and delusions) related to prolonged use of psychoactive drugs (eg cannabis) within the last month

Signs and Symptoms

- Hallucinations (often auditory but can be visual etc) and delusions (fixed false belief) often paranoid/ suspicious in nature
- Speech can be irrelevant and incoherent
- Abnormal behaviour and at times agitation and aggression
- Lack of insight is prominent

Investigations

- Physical causes for the symptoms must be excluded (delirium, drug and alcohol use/ withdrawal. Etc.)
- Full physical examination
- FBC
- VDRL
- Urine drug screen if available
- Consider HIV serotesting

Treatment

- Aim of treatment is to reduce/ remove symptoms and to encourage abstinence from the psycho-active drug
- Treatment duration is shorter (6 months) than for schizophrenia
- Primary treatment
 - Advise to stop using the drug (eg cannabis)
 - In some cases the symptoms will resolve without any further treatment

- A short course of oral **Diazepam** may help with agitation, insomnia while symptoms resolve (**Diazepam** 10mg bd for 1 week)
- Secondary /alternative treatment
 - If symptoms persist despite cessation of substance use, or markedly agitated or aggressive behaviour, an antipsychotic can be used
 - Give Chlorpromazine 100-300mg nocte or Haloperidol 2.5-5mg nocte
 - All anti-psychotic medications have a delayed onset of action – advice the patient/ guardian it will take 1-2 weeks before improvement is noted
 - Advice about the side effects:
 - ✓ Chlorpromazine: sedation, postural hypotension, constipation, photosensitivity
 - ✓ Haloperidol: Extra-pyramidal side effects (EPSE) eg. tremor, stiffness of limbs/jaw, salivation
 - ✓ This dose can be increased at subsequent reviews if symptoms remain present
 - ✓ If EPSEs persist consider reducing the dose of antipsychotic or adding Benzhexol 5mg daily

Third Line

- If symptoms have not improved on Chlorpromazine or if the person has lots of side effects use a second generation anti-psychotic
- ➤ Give **Risperidone** 1-2mg nocte
- Advice about side effects:
 - ✓ Risperidone: weight gain, sedation, impaired glucose tolerance

OR

- ✓ If compliance with medication is poor despite trying to reduce side effects and counselling on the importance of compliance with medication consider a long-acting depot anti-psychotic
- ✓ Give **Fluphenazine** 12.5mg IM into a large muscle (eg gluteal/ deltoid) as a test dose
- ✓ Caution: risk of Acute Dystonic reaction (painful spasm of head and neck muscles)
- ✓ If occurs give Benzhexol po if able to swallow
- ✓ Otherwise IV/IM Procyclidine or Benzhexol 5mg OR slow IV push
 Diazepam 5-10mg
- ✓ Advise about side effects: EPSE
- ✓ Maintenance dose Fluphenazine25-50mg IM every 4 weeks

✓ If EPSE persist consider reducing the dose adding **Benzhexol** 5mg daily

Red Flags

For referral

- If symptoms persist/ worsen despite 6-8 weeks of anti-psychotic medication at an effective dose
- > If side effects are not manageable

For admission

- Marked agitation / aggression should be managed as an inpatient following the Violence and Aggression treatment guidelines.
- Evidence that the patient is a risk to themselves (self harm/ neglect/ vulnerable to exploitation) or a risk to others (agitation/ aggression)
- If insight is lacking and there is no guardian to ensure compliance with medication at home or if compliance is poor at home despite supervision.

Note: Neuroleptic Malignant Syndrome is a severe but rare complication of antipsychotics, presenting with fever, rigidity, fluctuating pulse and BP and reduced conscious level. It is a medical emergency. All anti-psychotics should be stopped and the person referred for medical admission.

Follow-up

- If inpatient, review the person every 24 hours until stable.
- If outpatient, review in 1-2 weeks initially and then every 4-8 weeks once more stable.
- Screen for ongoing symptoms and monitor for side effects at each review. Adjust the dose of medication accordingly.
- Ask about any other drug or alcohol use and give advise about use.
- Screen for low mood and suicidal ideation at each review and follow Depression Treatment Guideline if present.
- Once symptoms are improving advice the patient to return to their usual daily activities, including work if employed.

Treatment duration:

- Medication should be continued for 6 months from complete symptom resolution resolution
- When considering stopping medication, discuss carefully with the patient and guardian and start to reduce slowly over 2-4 weeks
- Advise about symptoms that would indicate relapse (difficulty sleeping, auditory hallucination, suspicious thoughts) and inform to return to the clinic promptly
- Continue to monitor until medication free for
 4- 6 weeks before discharging
- Advise them to return to the clinic for review if have any concerns in the future

25.9.3 Puerperal Psychosis

- A psychotic disorder affecting the mother, which can develop within a few hours and up to 4 weeks of childbirth, with peak incidence around 24-28 hours
- It can be particularly florid with marked agitation and aggression due to vivid hallucinations and delusions
- Care must be taken, as there can be significant risk to both mother and baby

Signs and Symptoms

- Confusion
- Insomnia
- Anxiety
- Agitation
- Possible aggression
- Hallucinations (in any modality) and delusions (particularly paranoid persecutory)

Investigations

- The puerperium is a time of increased risk of many physical conditions such as sepsis, postpartum haemorrhage, metabolic imbalance, eclampsia etc.
- These conditions can present with delirium and so care must be taken to exclude underlying physical causes

- Full physical examination including vital signs
- FBC U&E LFT
- Blood/ urine glucose
- MRDT
- Consider HIV serotesting

General Management

- The aim of treatment is to reduce/ alleviate symptoms to allow a return to usual functioning and to promote good bonding between mother and baby
- As with all psychotic disorders treatment duration is for at least 1 year
- First assess the risk of the mother to herself and her baby (some may have thoughts of harming their baby due to the psychotic symptoms) and the risk of aggressive behaviour
- If present follow Violence and Aggression
 Treatment Guideline. Ensure that the baby is in
 the care of the guardian and that all mother
 baby interactions are supervised until more
 stable
- Care should be taken in breastfeeding mothers as medication can be found in the breastmilk.
 Use low doses and increase slowly. Monitor the baby for evidence of sedation
- Primary treatment
 - ➤ Give Chlorpromazine 100-200mg nocte or Haloperidol 1.25-2.5mg nocte

- All anti-psychotic medication have a delayed onset of action – advise the patient/ guardian it may take 1-2 weeks before significant improvement is noted
- Advise about the side effects:
- Chlorpromazine: sedation, postural hypotension, constipation, photosensitivity
- Haloperidol: Extra-pyramidal side effects (EPSE) eg. tremor, stiffness of limbs/jaw, salivation
- This dose can be increased at subsequent reviews if symptoms remain present
- If EPSEs persist consider reducing the dose of anti-psychotic or adding Benzhexol 5mg daily
- Secondary /alternative treatment
 - If no improvement after 4-6 weeks on adequate dose of Chlorpromazine/
 Haloperidol or unacceptable side effects persist, consider a second generation anti-psychotic
 - ➤ Give **Risperidone** 0.5-1mg nocte
 - Advice about side effects:
 - ✓ Risperidone: weight gain, sedation, impaired glucose tolerance

Red Flags

- For referral
 - If symptoms persist or worsen, despite adequate doses of anti-psychotic for 6-8 weeks
 - ➤ If there is evidence that the baby is failing to thrive (dehydration, weight loss, inadequate care etc)

For admission

- Marked agitation / aggression should be managed as an inpatient following the Violence and Aggression treatment guidelines
- Evidence that the patient is a risk to themselves (self harm/ neglect/ vulnerable to exploitation) or a risk to others (agitation/ aggression)
- If there is no guardian to take care of the baby and to ensure compliance with medication at home or if compliance is poor at home despite supervision

Follow-up

- If an inpatient review every 8 hours until the symptoms improve.
- If an outpatient review twice weekly until symptoms improve. Then review every 2-8 weeks.
- Screen for ongoing symptoms and monitor for side effects at each review. Adjust the dose of medication accordingly.

- Ask about any drug or alcohol use and give advise about use.
- Once symptoms are improving advise the patient to return to their usual daily activities, including work if employed
- Treatment duration
 - If first episode: Medication is continued for 1 year from complete resolution of symptoms
 - ➤ If previous episodes of psychotic illness (e.g. schizophrenia/ Bipolar affective disorder) continue medication for 2-5 years from complete resolution of symptoms
 - When considering stopping medication, discuss carefully with the patient and guardian and start to reduce slowly over 4-8 weeks
 - Advise about symptoms that would indicate relapse (difficulty sleeping, auditory hallucination, suspicious thoughts) and inform to return to the clinic promptly
 - Continue to monitor until medication free for 2-3 months before discharging
 - Advise them to return to the clinic for review if have any concerns in the future or when pregnant again (high risk of recurrence in future pregnancies)

25.10 Mood Disorders

 These disorders mainly present with a disturbance of mood, which can either be elevated or depressed, with associated changes in activity levels and behaviour

25.10.1 Depression

- The person experiences persistent sadness with decreased energy and enjoyment of usual activities, everyday for at least 2 weeks
- If it is moderate to severe there is impairment of the usual occupational and functional activities such as going to work, housework and self care
- Suicidal ideation should be asked about at every assessment

Signs and Symptoms

- Low mood or irritability
- Reduced energy and enjoyment
- Poor sleep
- Reduced appetite and weight loss
- Poor concentration
- Feelings of guilt and worthlessness
- Possible suicidal ideation
- Looks sad
- Poor eye contact
- Reduced speech at low volume
- Evidence of weight loss and poor self care
- Possible agitation
- If severely unwell can develop psychotic symptoms (delusions and hallucinations)

Investigations

- Some physical illnesses can present with depressive symptoms (anaemia, hypothyroidism) so these must be excluded and the person assessed for dehydration and malnutrition
- Full physical examination including vital signs.
- FBC ,U&E, glucose
- If evidence of hypothyroidism on examination (weight gain, dry skin, goitre) check TFTs if possible
- VDRL
- Consider testing HIV serostatus
- Urine drug screen if available

General Management

- The aim of management is to completely resolve symptoms, if possible, to allow the person to return to their previous activities and occupation
- Duration of treatment varies from 6 months to 2 years depending on whether it is a first episode of a recurrent illness

Primary treatment

- If mild to moderate depression (symptoms of depression present but still managing to undertake most of daily activities), first line treatment should be supportive counselling (listen to and try to understand their problems, help them to think about possible solutions, encourage enjoyable activities)
- Consider referring to local supportive groups (church, womens groups, etc.)
- If moderate to severe, or if mild depression persists despite counselling commence anti-depressant treatment
- ➤ Give **Amitriptyline** 50-75mg nocte
- Increase after 2 weeks by 25mg if symptoms persist up to a maximum of 200mg
- Monitor for side effects (sedation, hypotension, dry mouth, constipation, sexual dysfunction)
- Explain that antidepressants have delayed onset of action and it may take 1-2 weeks before improvements are noted
- If suicidal ideation is present, give medication to the guardian to administer (Amitriptyline can be fatal in overdose due to cardiac effects). If no guardian, admit the patient until suicidal ideation is resolved.

- If person is elderly or physically frail (eg HIV) reduce the starting dose to Amitriptyline 25mg nocte and increase the dose slowly.
- Secondary /alternative treatment
 - ➢ If not improved on an adequate dose of amitriptyline for 4-6 weeks or unable to tolerate due to side effects change to SSRI
 - ➤ Give **Fluoxetine** 20mg daily
 - Increase after 2 weeks by 20mg upto a maximum of 60mg daily (rarely needed)
 - Monitor for side effects (agitation, increase in anxiety initially, insomnia, Gl upset)
 - If psychotic symptoms (hallucinations or delusions) present, treat with an anti-psychotic and consider referral to district psychiatry team
 - ✓ Give Chlorpromazine 75 150mg nocte or Haloperidol
 1.25-2.5mg nocte
 - ✓ Increase dose every 2 weeks until psychotic symptoms have resolved

Red Flags

- For referral
 - Ongoing or worsening symptoms despite adequate anti-depressant treatment for 2- weeks
 - Depression with psychotic symptoms

- Persistent or increasing suicidal ideation or if the person has developed a suicide plan
- Evidence of self neglect, dehydration or malnutrition

For admission

- Evidence that the patient is a risk to themselves (self harm/ neglect/ vulnerable to exploitation) or a risk to others (agitation/ aggression)
- ➢ If presents with severe depression with psychotic symptoms and evidence of psychomotor retardation, this is an emergency and should be referred for admission for consideration of Electro Convulsive Therapy (ECT)
- ➤ If has active plans of suicide or a recent suicidal attempt.
- If has thoughts of harming others.
- Evidence of self neglect including not eating and drinking properly.

Follow-up

- If inpatient review daily until more stable, monitoring vital signs if food and fluid intake is poor
- IV fluids may be required
- If outpatient and suicidal ideation or psychotic symptoms present review weekly, until symptoms improved. Thereafter review every 2-4 weeks.
- At each review, ask about symptoms, suicidal ideation and side effects from medication.

- Monitor for emergence of manic symptoms (see below) as some people can develop manic symptoms when treated with anti-depressants.
- Give supportive counselling to patient (see mild –moderate depression above).
- Advise to increase physical activity gradually once improved and to return to usual activities when possible, including work if employed.
- Once stable review every 8 weeks.
- Duration of treatment
 - First episode of depression: continue medication for 6 months from complete resolution of symptoms.
 - Recurrent depressive illness: continue medication for 2 years after complete resolution of symptoms
 - When considering stopping medication, discuss carefully with the patient and guardian and start to reduce slowly over 4-8 weeks. Advise about symptoms that would indicate relapse (difficulty sleeping, low mood, reduced energy, increase worries) and inform to return to the clinic promptly. Continue to monitor until medication free for 1-2 months before discharging

25.10.1.1 Post Partum Depression

within 6 months of childbirth.

- Assess risk: of self harm, of harm from mother to baby and of neglect of the baby.
- If any concerns admit or refer for review.

Treatment

- Caution should be used when prescribing medication to pregnant and breastfeeding women.
- If mild to moderate depression, treat with supportive counseling.
- If medication is required:
 - Give Amitriptyline: start at 25-50mg nocte and increase slowly. Avoid during first trimester of pregnancy. In breastfeeding mothers it is the treatmet of choice. Monitor the baby for evidence of sedation as small amounts can be found in the breast milk.
 - Give Fluoxetine: Generally safe in pregnancy. Avoid in breastfeeding mothers.
 - Otherwise follow Depression Treatment Guidelines as above.

25.11 Mania

 Mania presents with elevated or irritable mood with associated increase in energy and activity and possible aggression and psychotic symptoms present for at least a week or resulting in complete disruption of usual functioning.

Signs and Symptoms

- Elevated or irritable mood
- Reduced need for sleep
- Increased energy and activity
- Talkativeness
- Over spending
- Increased libido
- Agitation and over activity
- Possible aggression
- Rapid speech
- Reduced attention and concentration
- Possible psychotic symptoms (hallucinations and delusions which are grandiose in nature – belief that special/important/ very rich person etc)

Investigations

- Full physical examination to exclude underlying organic causes (delirium)
- Vital signs: If severely unwell, dehydration/ exhaustion can be potentially fatal
- FBC, U&E, glucose
- If evidence of hyperthyroidism on examination (weight loss, tremor, exopthalmos, goitre) consider TFTs if available
- VDRL
- Consider HIV serotesting (HIV can present with secondary mania)

General Management

- The aim of management is to reduce/ alleviate symptoms, to allow the person to return to their previous daily activities and occupation if employed and to prevent any further relapses in the future. It is continued for at least 2 years.
- If marked agitation / aggression follow the Violence and Aggression Treatment Guideline.
- Always assess risk of suicide, harm to others and of self neglect
- Primary treatment
 - Commence either a mood stabiliser or an anti-psychotic
 - Give Sodium Valproate 400mg nocte increasing every few days by 200mg until symptoms improved. Usual effective dose 600- 1000mg.
 - Caution: Sodium Valproate should be avoided in women of child-bearing age due to teratogenic effects. If no available alternative advise about contraception.
 - Side effects: sedation, weight gain, liver impairment
 - Give Carbamazepine 200mg nocte increasing by 200mg every few days until on 400-600mg. Usual dose 400-600mg daily.

- Caution: **Carbamazepine** should be avoided in women of childbearing age due to teratogenic effects. If no available alternative advise about contraception.
- Side effects: sedation, rash, incoordination, RARELY Steven Johnsons syndrome
- If psychotic symptoms present also give an anti-psychotic medication
- Give Chlorpromazine 100-200mg nocte or haloperidol 2.5mg nocte
- Alternative treatment
 - ➢ Give Chlorpromazine 100-200mg nocte increasing every week by 100mg until symptoms improve. Usual dose 100 − 500mg daily
 - Side effects: sedation, postural hypotension, constipation, photosensitivity
 - Give Haloperidol 2.5mg nocte increasing by 1.25mg every week until symptoms resolves. Usual dose 1.25-5mg daily

Red Flags

- For referral
 - Ongoing or worsening symptoms despite adequate treatment for 2-4 weeks

For Admission

- Evidence of exhaustion, dehydration due to over-activity. IV fluid may be necessary
- Evidence that the patient is a risk to themselves (self harm/ neglect/ vulnerable to exploitation) or a risk to others (agitation/ aggression)
- No guardian available to monitor adherence to medication

Follow-up

- If inpatient review every 24 hours until stable.
- If outpatient and suicidal ideation/ psychotic symptoms present review twice a week until more stable.
- Otherwise review every 1-2 weeks until symptoms improve.
- Once stable review every 4-8 weeks.
- At every review, ask about symptoms, suicidal ideation and medication adherence and side effects.
- Provide education about illness to patient and guardians. Give supportive counselling and encourage return to previous activities and occupation as soon as able.
- Duration of treatment
 - First episode treatment should be continued for 2 years from complete resolution of symptoms.

- ➤ Multiple episodes: treatment should be for at least 5 years from complete resolution of symptoms and some may need lifelong medication.
- When considering stopping medication, discuss carefully with the patient and guardian. There is significantly increased risk of relapse if medication is stopped suddenly. Start to reduce slowly over 2-3 months. Advise about symptoms that would indicate relapse (difficulty sleeping, low/elevated mood, reduced or increased energy, increased worries) and inform to return to the clinic promptly. Continue to monitor until medication free for 1-2 months before discharging

Index

Abnormal Presentation of the

Fetus, 294

Abnormal Vaginal Bleeding, 282 Abnormal vaginal discharge, 398 Abortion and Its Complications, 283

Acute Carditis, 253 Acute Diarrhoea, 139 Acute eczema, 428

Acute Haemorrhage with Shock, 7

Acute Heaemorrhage, 9
Acute Heart Failure, 47
Acute Kidney Injury, 353
Acute Leukemias, 243, 244
Acute Liver Failure 161
Acute Myelopathy, 91
Acute Osteomyelitis, 251
Acute Pulmonary Oedema, 353

Acute Renal Failure, 357
Acute renal failure, 461
Acute Respiratory Distress

Syndrome, 353

Acute Respiratory Infections

(ARI) in Children, 371 Acute scrotal pain, 405 Acute scrotal swelling, 405 Acute Vaginal Bleeding, 282

AIDS Dementia, 179
Alcohol hallucinosis, 534
Alcohol intoxication, 527
Alcohol related disorders, 527

Alcohol withdrawal syndrome, 528

Allergic Conjuctivitis, 304
Allergic sinusitis, 103
Altered Consciousness and

/or Convulsions, 351 Amoebiasis, 134 Anaemia, 17

Ancylostoma Infestation, 368 Anemia in Pregnancy, 271

Animal bite, 439 Antenatal Care, 257

Antepartum Haemorrhage, 269

Antispasmodics, 513 Anxiolytics, 513 Aplastic Anaemia, 21

Arthritis (Non-infective), 246

Arthritis (Septic), 246

Ascaris Infestation, 368 Asthma (Recurrent Wheezing), 380 Asthma, 381

Attempted suicide, 539
Atypical Pneumonia, 389
Bacillary Dysentery, 135
Bacterial Conjunctivitis, 305
Bacterial Endocarditis, 255
Bacterial Keratitis, 307
Bacterial Meningitis, 195
Bacterial sinusitis, 102

Bacterial skin infections, 417 Balanitis, 408 Balapostitis, 408 BCG vaccination, 437 Bee stings, 450 Bells' palsy, 89 Bite, 439

Bleeding Disorders, 33 Blood Conditions, 1 Brain abscess, 81 Brain oedema, 93 Breast Abscess, 288 Breast Cancer, 234 Bronchiectasis, 390 Bronchitis (Acute), 379

Bronchitis, 379 Bubo, 406 Bullae, 420

Bullous Keratopathy, 307

Burns, 443 Buruli ulcer, 436 Cachexia, 225 Cancer, 225 Candidiasis/

Oroesophageal, 320 Carbuncles, 418 Cellulitis and Tropical Pyomyositis, 256 Cellulitis, 421

Central Nervous System

Disorders, 177

Cerebral Malaria, 264 Cerebral Oedema, 92

Cerebral Toxoplasmosis, 178

Cervical Adenitis, 379

Cervical Cancer, 238 Chancroid, 395 Chemical Injury, 317 Chicken pox, 431 Cholera, 136 Chorea, 254 Chorea, 87

Chorioamnionitis, 268 Chronic Blepharitis, 307 Chronic Diarrhoea, 154 Chronic Disorder

Anaemia, 22

Chronic eczema, 429 Chronic Leukemias, 244 Chronic Lung Disease

/ COPD, 391

Chronic Lymphocytic Leukemia, 244 Chronic Myeloid Leukemia, 244

Chronic Osteomyelitis, 252 Cluster Headaches, 76 Coagulation Defects, 40 Coagulopathy, 352 Common Cold

(Nasopharyngitis), 374 Common Infections, 353 Community Acquired Pneumonia (CAP), 388

Congenital Heart Disease, 50

Congenital syphilis, 414 Congestive Heart

Failure, 48

Conjuctivitis, 304
Constipation, 137
Convulsions , 264, 298
Convulsions in Children, 114
Corneal Anaestasia, 307
Cryptococcal Meningitis, 197

Cyanotic congenital Heart Disease, 51 Cytomegalovirus (CMV) Retinitis, 313 Cytopenias, 241 Cytoses, 241

Delirium tremens, 521 Delirium tremens, 529 Delirium, 521
Dementia, 524
Dental Abscess, 324
Depression, 563
Depression, 563

Diabetes in Pregnancy, 272 Diabetes Mellitus, 121 Diabetic Ketoacidosis, 117

Diarrhoea, 139

Disciform Keratitis, 309 Disseminated Intravascular

Coagulation, 41
Dysfunctional Labour
Syndrome, 275
Dysfunctional Uterine

Bleeding, 283

Dysmenorrhoea, 287 Dyspepsia, 158 Dysphagia, 237 Ear, nose and Throat

(ENT), 95

Early syphilis, 413 Eclampsia, 263 Ecthyma, 419

Ectopic Pregnancy, 286

Eczema, 427

Emergency Hypertension, 57

Empyema, 390 Encephilitis, 264

End Organ Dysfunction, 164
Endogenous Bacterial
Endophthalmitis, 314
Endophthalmitis, 314
Enlarged inguinal node, 406
ENT Emergencies, 105
Enterobius Infestation, 368

Envenomation, 450 Epilepsy, 264 Epilepsy, 66 Epistaxis, 100

Epithelial Keratitis, 309

Erysipelas, 421

Erythema multiforme, 423 Essential Tremor, 86 Exposure Keratopathy, 307

Extra Pulmonary Tuberculosis, 215 Failure to Thrive, 182 Febrile Neutropenia, 30

Fever, 174
Fixed medicine
eruption, 425
Foetal Well-being
or Maturity, 263
Folluculitis, 418

Foreign Body Inhalation, 105 Fresh Frozen Plasma, 8 Fungal skin infections, 430

Furunculosis, 418 Gastric lavage, 462 Gastritis, 159 Gastrointestinal Conditions, 134

Genital ulcer disease, 395

Genital warts, 410 Gingivitis, 325 Glaucoma, 316 Gout, 249 Haematological Conditions, 1 Haematological

Malignencies, 240, 241 Haemolytic Anaemia, 23

Halitosis, 515

Head and Neck Cancers, 233 Headaches and Facial Pain

Syndrome, 74
Hemophelia, 34
Heparin Induced
Thrombocytopenia, 46
Hepatic Amoebiasis, 134
Hepatomenopathy, 241
Herpes genital, 395
Herpes simplex, 431
Herpes zoster, 432

Herpetic Keratitis, 309

Hiccup, 512 HIV, 162

HIV/AIDS in Pregnancy, 270 Hyperosmolar Non-Ketotic

Coma (HONK), 119 Hyperparasitaemia, 341 Hypertension, 52-56 Hypertensive Disorders in Pregnancy, 259 Hypoglycemia, 128 Hypoglycemia, 264
Hypothermia, 300
Hypothyroidism, 130

Immune Thrombocytopenic

Purpura, 36

Immunoglobulins, 439 Immunosuppression, 360

Impetigo, 417

Infectious and Parasitic

Conditions, 79

Infectious Diseases, 162

Insect bite, 450 Insect sting, 450 Insulin Dependent (Type 1), 123 Insulin, 120, 126

Intra-dermal regimes, 441

Intrauterine Death

(IUD), 277 Iodine Deficiency Disorders, 132

Iron Deficiency Anaemia, 25

Iron poisoning, 470

Iron, 470

Ischemic Heart Disease, 59 Kaposi's Sarcoma, 230

Keratitis 307
Lactic Acidosis, 352
Late syphilis, 414
Leprosy, 188
Leprosy, 436
Leukemia, 242
Low Birth Weight
Babies, 299
Lower abnominal
pain, 400, 403

Lower Respiratory Tract

Infections, 380 Lung Abscess, 390 Lyell's disease, 422 Lymphadenopathy, 191 Lymphadenopathy, 241 Lymphoma, 242

Lymphoma, 242 Malaria, Non-Severe, Uncomplicated, 334 Malnutrition, 145 Malnutrition, 475

Mania, 569

Mania, 569 Mastisis, 288 Mastoiditis, 95

Maxillofacial/Mandibular

Trauma, 332

Measles vaccination, 438

Meconium-stained

Liquor, 295

Medical Conditions in Pregnancy, 270 Medicine-Resistant Tuberculosis, 209

Megoblastic Anaemia, 27

Meningitis, 194 Meningitis, 264 Meningococcal, 197 Mental Disorders, 177 Metabolic Acidosis, 352

Migraine, 74

Mild Local Anaesthetic

Toxicity, 329

Mild malnutrition, 476 Mild to Moderate Pneumonia, 388

Milder Asthma Attacks, 383 Moderate Local Anaesthetic

Toxicity, 329

Moderate malnutrition, 476

Mood disorders, 563 Mouth Ulcers (Sores), 330 Movement disorders, 83 Mucocutaneous Lesions, 231

Multiple Sclerosis, 92 Musculoskeletal Pain and Trauma, 250 Muskuloskeletal Disorders, 246

Mycardial infarction, 61

Myelodysplastic Syndromes, 32 Myoclonus, 86

Neonatal conjunctivitis, 411 Neonatal Infections, 297 Neonatal Jaundice, 301 Neonatal pustulosis, 420 Nephrotic syndrome, 459 Neurocysticercosis, 82 Neurosyphilis, 416 Non Severe Anaemia, 5 Non-Insulin Dependent

(Type 2), 129

Non-specific Inflammatory

Arthritis, 246

Nosocomial Pneumonias, 389

Nueropathy, 87

Nutritional disorder, 475 Obstetric and Gynaecogic

Disorders, 257

Obstetric Conditions, 257 Obstetrical Fistula, 289

Odontogenic and Maxillofacial

Infections, 326 Odynophagia, 237 Oesophagus Cancer, 237

Onchocerciasis (River Blindness), 362 Onchocerciasis, 436 Open wound, 448

Ophthalmia Neonatorum, 305 Ophthalmic Conditions, 304 Oral and Maxillofacial Conditions, 320 Oral Trauma, 330 Orbital Cellulitis, 319

Organic mental disorders, 521

Organophosphate poisoning, 471

Organophosphates, 471 Osteomyelitis, 251 Otitis Externa, 96 Otitis Media, 95

Otitis, 96

Pain management, 504 Palliative care, 504

Paracetamol poisoning, 473

Paracetamol, 473

Parkison's disease, 83,84,85 Passive immunizations, 439

Pellagra, 502

Peptic Ulcer Disease, 159 Periodontal Abscess, 325 Periodontitis, 326

Peripheral Arterial disease, 61 Peritonsillar Abscess, 377 Persistent Diarrhoea, 156

Petit mal, 64

Pharyngitis, 104 Pharyngitis, 375 Platelets, 7

Pneumocystis Jirovecii Pneumonia, 172 Poisoning 465 Polio, 437 Post Abortal

Haemorrhage, 284
Post Abortal Sepsis, 283
Post Exposure Prophylaxis, 165

Post exposure

vaccination, 440, 443
Postoperative Bacterial
Endophthalmitis, 314
Postpartum depression, 568

Postpartum Haemorrhage

(PPH), 280

Pre-Eclampsia, 260

Pre-exposure immunization, 440

Pregnancy-Induced Hypertension, 259

Prelabour/Premature Rupture

of Membranes, 266 Premature Babies, 299 Prepregnancy Care, 273 Primary Neurological

Disorders, 177
Primary Neurological

Disorders, 180

Primary Open Angle Glaucoma, 316

Primary PPH, 280 Progressive Multifocal

Leukoencephalopathy (PML), 179

Prolapsed Cord, 295

Prurigo, 433 Pruritis, 433

Psychiatric conditions, 520
Psychotic disorder, 520
Psychotropics, 520
Puerperal psychosis, 558
Pulmonary Edema, 47
Pulmonary Embolism, 392
Pulmonary Tuberculosis, 214

Renal colic, 461 Renal failure, 461, 463

Resomal, 144

Respiratory Conditions, 371

Respiratory Distress Syndrome, 301

Retropharyngeal Abscess, 377

Reye's syndrome, 432 Rheumatic Fever, 253 Rupture of Membranes, 295 Salicylate poisoning, 470

Salicylates, 470

Salivary Gland Disease, 327

Scabies, 433
Schistosomiasis
(Bilharzia), 364
Schistosomiasis
Haematobium, 364
Schistosomiasis
Mansonii, 364

Schizophrenia, 520,546 Secondary PPH, 281 Seizures, 62, 63, 64 Self harm, 539 Sepsis, 221 Severe Allergic Eye Disease, 307

Severe Anaemia, 3 Severe Anaemia, 351 Severe Asthma, 381

Severe Chemical Injuries, 318 Severe Local Anaesthetic

Toxicity, 329

Severe Malaria in Paediatric

In-Patients, 355
Severe Malaria in
Pregnancy, 270
Severe Malaria in
Pregnancy, 358
Severe Malaria, 340
Severe malnutrition, 475
Severe Pneumonia, 389
Severe Pre-Eclampsia, 261
Sexually transmitted

infections, 393 Shingles, 432 Shock, 108

Sickle Cell Anaemia, 17

Sinusitis, 101 Sinusitis, 374 Skin conditions, 417

Snake bite, 450

Solid Malignancies, 225, 241

Splenomegaly, 241

Spontaneous Bleeding, 352

Squashed stomach syndrome, 516 Stable Angina, 60

Staphylococcal scalding skin

syndrome, 422 Status Epilepticus, 64

Steven johnson's syndrome, 423

STI, 393 Stroke, 68

Stromal Necrotic Keratitis, 309 Strongyloides Stercoralis

Infestation, 369 Subacute eczema, 429

Subarachnoid haemorrhage, 71 Substance induced psychosis, 552

Superficial injury, 452

Suppurative Lung Disease, 390

Swallowed, 466

Syndromic management, 394

Syphilis, 412

Taenia Saginata/Solium (Tapeworm)

Infestation, 370 Tension Headache, 77

Tetanus, 198

Thrombotic Thrombocytopenic Purpura-Haemolythic Uraemic

Syndrome, 39 Tinea, 430 Tonsillitis, 104 Tonsillitis, 375

Toxic epidermal necrolysis, 424

Trichiasis, 307

Trichuris Infestation, 368

Trigeminal and Glossopharyngeal

Neuralgia, 328

Trigeminal neuralgia, 77 Tropical ulcer, 435 Trypanosoiasis (Sleeping

Sickness), 365

Tuberculosis Meningitis, 208

Tuberculosis, 202

Uncomplicated Malaria, 270 Upper GIT Candidiasis, 177 Urethral discharge, 397

Urethritis, 397 Urticaria, 425 Use of activated charcoal, 468 Vaccination

recommendations, 442 Vaccinations, 437

Vaccine administration, 441

Vaccine, 440 Vaginal Candidiasis (Monoliasis, 286) Vaginal, 398 Varicella, 431

Venous Thromboembolism, 41

Vestibulitis, 101

Viral Meningoencephalitis, 79 Viral skin infections, 431 Vitamin A deficiency, 502

Vomiting, 160 Wernicke Korsakoff syndrome, 531

Worm Infestations, 368

Wounds, 439 Xerophthalmia, 502



MALAWI ESSENTIAL MEDICINES LIST (MEML) 2015



Introduction

The MEML lists all those medicines considered to be most suitable for current use in the country. Although primarily intended for public sector application, it is equally appropriate for private sector prescribers, who are strongly encouraged to select medicines on the MEML whenever possible.

1. Categorisation of medicines

- medicines items have again been categorised using a three letter coding system according to the:
 - approved level of use (H,D or C)
 - therapeutic priority (V or E)
 - procurement system (A or B)
- Details of this are given in the Presentation of Information section. Such
 categorisation is intended to facilitate the prioritisation of selection and
 subsequent procurement of drugs both by Central Medical Stores Trust
 (CMST) and user units.

2. Procedure for amendments of the MEML

- Prescribers are encouraged to continually and critically review the relevance of the MEML to current clinical practice.
- Suggestions for amendments should be made through submission of a proposal in writing to:

HTSS-Pharmaceuticals

PO Box 30377, Lilongwe 3, Malawi

Tel: (265) 01 788 371 Fax: (265) 01 788 502

- An amendment form which may be used for this purpose has been included at the end of this booklet.
- Amendments can include the addition or deletion of an item, change of dose presentation, change of categorisation code (s), change of layout of the publication, etc.)
- When sufficient proposals have been received or if urgent amendments are required, a special NMC will be convened to review these and agree on any changes to the MEML.
- Otherwise it is planned for the MEML to be revised once every 3 years in conjunction with the MSTG.

3. For adverse reactions:

Please contact the Pharmacy, Medicines and Poisons Board at info@pmb.mw



Contents

| | ductionduction | |
|-------|--|-----|
| Prese | entation of Information | V |
| 1. | General | V |
| 2. | Categorisation/Prioritisation of drug items | vi |
| 3. | Emergency orders | ix |
| 4. | Listing by category/priority within sections | ix |
| 5. | Abbreviations | х |
| 1. | Anaesthetics | 1 |
| 1.1 | General anaesthetics and oxygen | 1 |
| 1.1.1 | Inhalation Medicines | 1 |
| 1.1.2 | . Injectable Medicines | 1 |
| 1.2 | Local anaesthetics | 1 |
| 1.3 | Preoperative medication and sedation for short-term procedures | 2 |
| 2. | Medicines for Pain and Palliative Care | 3 |
| 2.1 | Non-opioids and non-steroidal anti-inflammatory medicines (NSAIMs) | 3 |
| 2.2 | Opioid analgesics | 3 |
| 2.3 | Medicines for other common symptoms for palliative care | 3 |
| 3. | Antiallergics and medicines used in Anaphylaxis | 4 |
| 3.1 | Antihistamines | 4 |
| 3.2 | Medicines used in nasal allergy | 4 |
| | Antidotes and other medicines used in poisonings | |
| 5. | Anticonvulsants/Antiepiletics | 7 |
| 6. | Anti-infective medicines | 8 |
| 6.1 | Antihelmintics | 8 |
| | Intestinal anthelmintics | |
| 6.1.2 | . Antifilarials | 8 |
| 6.1.3 | Antischistosomals and other antitrematode medicines | 8 |
| 6.2 | Antibacterials | 8 |
| 6.2.1 | .Penicillins and cephalosporins (Beta Lactam medicines) | 8 |
| 6.2.2 | Other antibacterials | 9 |
| 6.2.3 | Antileprosy medicines | .10 |
| 6.2.3 | Antileprosy medicines | 10 |
| 6.3 | Antifungal medicines | 10 |
| 6.4 | Antiviral medicines | 11 |
| 6.4.1 | Antiherpes medicines | 11 |
| 6.4.2 | Antiretrovirals | 11 |
| 6.5 | Antiprotozoal medicines | 13 |
| | Antiamoebics and antigiardiasis medicines | |
| 6.5.2 | Antileishmaniasis medicines | .13 |
| 6.5.3 | Antimalarials | 13 |
| 6.5.4 | Antipneumocytosis and Anti-toxoplasmosis medicines | .14 |
| 6.5.5 | Antitrypanosomal medicines (African) | 14 |
| 7. | Anti-migraine medicines | .15 |
| 7.1 | For treatment of acute attack | 15 |
| 7.2 | For prophylaxis | 15 |
| 8. | Antineoplastic and immunosuppressant medicines | 16 |
| 8.1 | Immunosuppressive medicines | 16 |
| 8.2 | Cytotoxic and adjuvant medicines | 16 |
| | Hormones and antihormones | |
| 9 | Antiparkinsonism medicines | 17 |
| 10 | Medicines affecting the blood | |
| 10.1 | Antianaemic medicine | |
| | Medicines affecting coagulation | |
| | Medicines to treat hyperkalaemia | |
| | | |

| 11. | Blood products and plasma substitutes | .19 |
|-------------|--|-----|
| 12 | Cardiovascular medicines | .20 |
| 12.1 | Antianginal medicines | .20 |
| 12.2 | Antiarrhythmic medicines | .20 |
| 12.3 | Antihypertensive medicines | .20 |
| 12.4 | Medicines use in heart failure | .20 |
| 12.5 | Antithrombotic medicines | .21 |
| 12.6 | Lipid-lowering agents | .21 |
| 12.7 | Antihypotensive medicines | .21 |
| 13 | Dermatological medicines (Topical) | .22 |
| 13.1 | Antifungal medicines | .22 |
| 13.2 | Anti-infective medicines | .22 |
| 13.3 | Anti-inflammatories and antipruritic medicines | .23 |
| 13.4 | Medicines affecting skin differentiation and proliferation | .23 |
| 13.5 | Scabicides and pediculocides | .24 |
| 13.6 | Other topical preparations | .24 |
| 14. | Diagnostic agents | .25 |
| 14.1 | Ophthalmic medicines | .25 |
| 14.2 | Radiocontrast media | .25 |
| 15 | Disinfectants and Antiseptics | .27 |
| 15.1 | Antiseptics | 27 |
| 15.2 | Disinfectants | .27 |
| 16. | Diuretics | .28 |
| 17 . | Gastrointestinal medicines | .29 |
| 17.1 | Antacids and other antiulcers medicines | .29 |
| 17.2 | Antiemetics | .29 |
| 17.3 | Anti-inflammatory medicines | .29 |
| 17.4 | Laxativesand Cathartics | .29 |
| 17.5 | Medicines used in diarrhoea | .30 |
| 17.5 | 1 Oral rehydration preparations | .30 |
| 17.5 | 2 Medicines for diarrhoea in children | .30 |
| 17.5 | 3 Antimotility medicines | .30 |
| 17.6 | Antihaemorrhoidals | .30 |
| 17.7 | Antispasmodics | .30 |
| | Hormones, other endocrine medicines and Contraceptives | |
| 18.1 | Adrenal hormones and synthetic substitutes | .31 |
| 18.2 | Androgens | .31 |
| 18.3 | Contraceptives | .31 |
| 18.3 | 1. Oral Hormonal contraceptives | .31 |
| 18.3 | 2 Injectable Hormonal contraceptives | .31 |
| 18.3 | 3 Intra-uterine devices (IUD) | .31 |
| 18.3 | 4 Barrier contraceptives | .32 |
| 18.3 | | |
| 18.4 | Oestrogens | |
| | Insulins and other medicines used for diabetes | |
| | Ovulation inducers | |
| | Progestogens | |
| | Thyroid hormones and antithyroid medicines | 33 |

| 19. | Immunologicals | 34 |
|-------------|--|----|
| 19.1 | Immunological diagnostic agents | 34 |
| 19.2 | Sera and immunoglobulins | 34 |
| 19.3 | Vaccines | 34 |
| 19.3 | 1 Vaccines for universal immunisation | 34 |
| 19.3 | | |
| 20 | Muscle relaxants (peripherally acting) and cholinesterase inhibitors | 36 |
| 21. | Ophthalmological preparations | 37 |
| 21.1 | .Anti-infective agents | 37 |
| 21.2 | Anti-inflammatory agents | 37 |
| 21.3 | Local anaesthetics | 38 |
| 21.4 | Miotics and antiglaucoma medicines | 38 |
| 21.5 | Mydriatics | 38 |
| 21.6 | Others | 38 |
| 22 O | bstetric medicines | 39 |
| 22.1 | Oxytocics | 39 |
| 22.2 | Antioxytocics | 39 |
| 22.3 | Myometrial relaxants | 39 |
| 22.4 | Medicines used in severe PIH and eclampsia | 39 |
| | Medicines used in primary PPH | |
| 22.5 | Medicines used in secondary PPH | 39 |
| 23 | Peritoneal dialysis solutions | 40 |
| 24 | Medicines for mental and behavioural disorders | |
| 24.1 | Medicines used in psychotic disorders | 41 |
| | Medicines used in mood disorders | |
| 24.2 | 1 Medicines used in depressive disorders | 41 |
| 24.2. | · | |
| | Medicines for enxiety disorders | |
| 24.4 | Medicines used for obsessive-compulsive disorders | 42 |
| | Medicines for disorders due to psychoactive substance use | |
| | Other Medicines for mental disorder | |
| 25. | Medicines acting on the Respiratory Tract | 43 |
| 25.1 | Antiasthmatic and medicines for chronic obstructive | |
| | onary disease | |
| | Solutions correcting water, electrolytes and acid-based disturbances | |
| | Oral preparations | |
| | Parenteral preparations | |
| | Vitamins and minerals | |
| | Ear, Nose and throat medicines in Children | |
| | Preparations for the ear | |
| | Preparations for the oropharynx | |
| | Specific Medicines for Neonatal Care | |
| | Medicines administered to the neonate | |
| | Medicines administered to the mother | |
| | Medicines for Arthritis | |
| | Medicines used to treat gout | |
| | Medicines used to treat Nutritional Disorders | |
| MEN | 1L Index | 50 |

Presentation of Information

1. General

- 1.1 **Medicines names:** as in previous lists, each medicines item is described by its *generic name* which, according to the National Medicine Policy (NMP) 2014, should be used for all prescribing and dispensing.
- 1.2 Order of Sections: Medicines are arranged by pharmacological/therapeutic groups following the same basic format as the "WHO Model List of Essential Drugs (Eighteenth List)" April 2013, with the addition of an extra section (Number 31; Nutritional Disorders)
- 1.3 **Medicines numbering**: each item is numbered within its section and thus can be conveniently identified by a composite number, consisting of the section number and the medicines numbers (final number):
- 2. Categorisation/Prioritisation of medicines items

Each MEML item is categorised by the following three code letter system:

2.1 **Level of use code** (H, D, or C)

This indicates the level of Health Institution at which the item would normally be permitted for use:

- **H = Health Centre level** (i.e. for use throughout the health system at Health Centre, District Hospital and Central Hospital levels).
- **D = District Hospital level** (i.e. for use at District Hospital and Central Hospital levels only)
- **C = Central Hospital level** (i.e. for use at Central Hospital level only)

Thus C drugs should not normally be used at D and H levels, and D drugs should not normally be used at H level.

However, there are *possible exceptions* to this general rule:

(a) Additional clinical expertise: if this is available at H level (e.g. An Ophthalmic Medical Assistant, Psychiatric Enrolled Nurse/Midwife, Clinical officer, etc), certain D level drugs may be made available for use by that particular prescriber.

This however will be at the discretion of the District Health Officer (DHO). Written authorisation should be given to the named prescriber by the DHO, listing the drugs authorised and a copy of this sent to the Chief of Health Services (CHS).

Note: Such D level drugs will be ordered from the district hospital and will not be part of the LMIS report the health centre sends to the DHO and therefore will not be delivered directly to the health centre by Regional Medical Stores.

- (b) Special clinical expertise: certain DHOs may have access to specialised clinical expertise in one or more areas and must request their Regional Medical Stores to make available certain C level drugs for use at the particular district hospital. Authorisation stating name of prescriber, specified drugs etc should be sought in writing from the Chief Executive Officer of CMST.
- (c) Maintenance treatment: individual patients being treated for chronic conditions may have arrangements made for them to receive maintenance treatment with the required D or C level drugs at H level.

In these cases, such arrangements should be authorised by the DHO and formalised in writing (named patient specified drug/s and duration of treatment) with a copy retained at the district.

2.2 Therapeutic priority code (V or E)

This code identifies the therapeutic importance of each item using the VEN system as follows:

- (a) V (Vital) drugs which:
 - are potentially life-saving
 - have significant withdrawal side-effects making regular supply mandatory
 - are of major public health importance (e.g. needed by many patients for treatment of serious. contagious diseases, needed to control epidemics, etc)
- (b) E (Essential) drugs which:
 - are effective against less severe, but nevertheless significant forms of illness
- (c) The **VEN** system also has a third category of N (Non-essential) drugs which are:
 - used for minor or self-limiting illnesses
 - of questionable efficacy
 - have a high cost for a marginal therapeutic advantage Items in this category were not included in the MEML due to economic constraints that affect the drug supply system and the subsequent need to carefully prioritise/rationalise drug selection and procurement.

2.3 Procurement system code (A or B)

This third code specifies how items will be procured by CMST and by the user units.

- (a) A List items:
 - Are generally required for large numbers of patients
 - Will be routinely procured and stocked by CMST
 - Include all H level drugs

Note: Where funds for procurement are insufficient, first priority will be given to the procurement and supply of Vital) Alist items (see 2.2 above) i.e. those of the highest therapeutic importance.

If funds remain after securing such VA items, procurement of (Essential) A – list items will then be initiated.

Thus ensuring the availability of A-list items is primarily the responsibility of CMST.

(b) B – List items:

- are generally required for limited number of patients
- willnot be routinely procured and stocked by CMST
- estimates of annual requirements for these will have to be made well in advance by the hospitals and submitted to CMST through the Pharmaceutical section of the HTSS, according to a pre-agreed time schedule.
- payment for these must also be made in advance prior to procurement by CMST and subsequent supply to the hospitals.

Thus procurement of B-list items is primarily the responsibility of the user units.

Each user unit must carefully consider the total annual budget allocated for medicines and medical supplies and make appropriate allocations for other categories.

It should be clearly understood that the ultimate decision as to which items to select/procure lies with the user units.

The categorisation of items by the NMC by therapeutic priority (see 2.2 above) should facilitate this selection process and is intended to help ensure the continuous availability of the most important (ie. VA) items.

3. Emergency orders

- Because of the limited numbers of patients involved, some potentially life-saving MEDICINES on the MEMLL are categorised as VB items.
- Following established procurement priorities, CMST will not therefore
 routinely stock these (B-list) items. Ideally user units should keep small
 contingency stocks of such items to cover emergencies. However, if
 necessary, special emergency orders can be placed (initially by phone or
 fax and then followed up by a written requisition with CMST for
 identified life saving VB drugs.
- These orders, which must be authorised by the DHO and countersigned by the Regional Medical Stores Pharmacist-in-charge, will not require advance payment, as with other B-list drugs. User units will be promptly notified if CMST is unable to procure and deliver these items within 24 hours.

4. Listing by category/priority within sections

In order to make the MEML easier to use by different levels of health institution, drugs are listed by generic name within individual sections and subsections according to the following order of priority.

- 1. H level then D level then C level.
- 2. Within each level the listing is V (vital) then E (essential) drugs which are on the A list for procurement (i.e. always stocked by CMST) followed by V (vital) then E (essential) drugs which are on the B list for procurement (i.e. only stocked if ordered/paid for in advance)

5. Abbreviations

amp = ampoule
aq = aqueous
cap = capsule

CHSU = Community Health Sciences Unit, Lilongwe

CMST = Central Medical Stores Trust

COM = College of Medicine EO = emulsifying ointment

g = gram inj = injection

IU = international units
i/m = intramuscular
i/v = intravenous

L = litre

KCH = kamuzu Central Hospital

mg = milligram mixt = mixture mL = millilitre

MOH = Ministry of Health MU = mega (million) units

oint = ointment paed = paediatric

PFR = powder for reconstitution

PIH = pregnancy – induced hypertension

PMPB = Pharmacy, Medicines and Poisons Board,

QECH = Queen Elizabeth Central Hospital,

RMS = Regional Medical Stores

s/c = subcutaneous or sugar-coated as appropriate

soln = solution susp = suspension tab = tablet vag = vaginal

YSP = yellow soft paraffin

%v/v = percentage volume in volume

ZCH = Zomba Central Hospital ZMH = Zomba Mental Hospital

mcg = microgram

1. Anaesthetics

1.1 General anaesthetics and oxygen

1.1.1 Inhalation Medicines

| 1.1.1 | innaiation wiedicines | | |
|--|--|--|---|
| 1.2 | | | |
| 1 | Halothane | inhalation | DVA |
| 2 | Isoflurane | inhalation | CVA |
| 3 | Nitrous oxide | medical gas | CEB |
| 4 | Oxygen | inhalation (medicinal gas) | DVA |
| 1.1.2. | Injectable Medicines | | |
| 2 3 4 5 6 7 8 9 10 11 | Atracurium Fentanyl Ketamine HCI Morphine sulphate Neostigmine Pethidine Propofol Suxamethonium Thiopentone sodium Vecuronium | inj, 10mg/mL, 5mL amp inj, 50mcg/mL, 10mL inj, 50 mg/mL, 10 mL amp inj, 2mg/mL, 50mL vial inj, 2.5mg/mL, 1mL amp inj, 50mg/mL, 2mL amp inj, 2%, 50mL vial inj, 50mg/mL, 2mL amp inj, 0.5 g vial PFR inj, 10mg vial, PFR | CEB CEB DVA CVA DVA CVA DVA CVA DVA DVA DVA |
| | | | |
| 1.2 | Local anaesthetics | | |
| 1.2 | Local anaesthetics Lignocaine HCI | inj, 1%, 25 ml vial | HEA |
| | | inj, 1%, 25 ml vial dental cartridges, | HEA HEA |
| 1 | Lignocaine HCI | | |
| 1 | Lignocaine HCI Lignocaine HCI | dental cartridges, | |
| 1 2 | Lignocaine HCI Lignocaine HCI + adrenaline | dental cartridges, 2% + 1/80,000, 2.2 mL | НЕА |
| 1 2 | Lignocaine HCI Lignocaine HCI + adrenaline Lignocaine HCI | dental cartridges, 2% + 1/80,000, 2.2 mL inj, heavy spinal, | НЕА |
| 1 2 3 | Lignocaine HCI Lignocaine HCI + adrenaline Lignocaine HCI + glucose | dental cartridges, 2% + 1/80,000, 2.2 mL inj, heavy spinal, 5% + 7.5% | HEA |
| 1 2 3 | Lignocaine HCI Lignocaine HCI + adrenaline Lignocaine HCI + glucose Lignocaine HCI | dental cartridges, 2% + 1/80,000, 2.2 mL inj, heavy spinal, 5% + 7.5% gel, 2%, 30 g tube | HEA CVB DEA |
| 1 2 3 | Lignocaine HCI Lignocaine HCI + adrenaline Lignocaine HCI + glucose Lignocaine HCI Lignocaine HCI | dental cartridges, 2% + 1/80,000, 2.2 mL inj, heavy spinal, 5% + 7.5% gel, 2%, 30 g tube inj, 2%, 20 mL vial | HEA CVB DEA CVB |
| 1 2 3 4 5 6 | Lignocaine HCI Lignocaine HCI + adrenaline Lignocaine HCI + glucose Lignocaine HCI Lignocaine HCI Lignocaine HCI | dental cartridges, 2% + 1/80,000, 2.2 mL inj, heavy spinal, 5% + 7.5% gel, 2%, 30 g tube inj, 2%, 20 mL vial spray, 10% | HEA CVB CVB CEB |
| 1 2 3 4 5 6 | Lignocaine HCI Lignocaine HCI + adrenaline Lignocaine HCI + glucose Lignocaine HCI Lignocaine HCI Lignocaine HCI Bupivacaine (heavy) | dental cartridges, 2% + 1/80,000, 2.2 mL inj, heavy spinal, 5% + 7.5% gel, 2%, 30 g tube inj, 2%, 20 mL vial spray, 10% | HEA CVB CVB CEB |

1.3 Preoperative medication and sedation for short-term procedures

| 1 2 | Amiodarone Atropine sulphate | inj, 30mg/mL, 10mL inj, 600 mcg/mL, 1mL amp | CVA DVA |
|-----|---------------------------------|--|------------|
| | | | |
| 3 | Diazepam | inj, 5 mg/mL, 1 mL amp | DVA |
| 4 | Diazepam | tab, 5 mg | DEA |
| 5 | Dobutamine | inj, 12.5mg/mL, 20mL amp | CVA |
| 6 | Ephedrine | inj, 30mg/mL, 1mL amp | DVA |
| 7 | Epinephrine | inj, 100mcg/mL, 10 mL amp | DVA |
| 8 | Lorazepam | inj, 4mg/mL | CEB |
| 9 | Metoclopramide | inj, 5mg/mL, 2mL amp | CEB |
| 10 | Midazolam | inj, 1mg/mL, 2mL amp | CEB |
| 11 | Morphine sulphate | inj, 15 mg/mL, 1 mL amp | DVA |
| 12 | Morphine sulphate | tab, slow-release, 10 mg | DVB |
| 13 | Nitro glycerine | tab, 0.3mg | CVA |
| 14 | Pethidine HCI | inj, 50 mg.mL, 2 mL amp | DVA |
| 15 | Phenylephrine | inj, 10mg/mL, 1mL amp | CEB |
| 16 | Promethazine | tab, 25 mg | DEA |
| 17 | Promethazine HCI | elixir, 5 mg/5 mL | DEA |
| 18 | Promethazine HCI | inj, 25 mg/mL, 2 mL amp | DEA |
| 19 | Sodium citrate | soln, oral, 300mmol/L | DVA |

2. Medicines for Pain and Palliative Care

| 2.1 | Non-opioids and non-steroida | l anti-inflammatory medicines | (NSAIMs) |
|--------|-------------------------------------|---|------------|
| 1 | Aspirin | tab, 300 mg | HVA |
| 2* | Diclofenac sodium | tab, 50 mg | DEA |
| 3* | Ibuprofen | tab, 200 mg | DEA |
| 4 | Indomethacin | tab, 25 mg | HEA |
| 5 | Mefenamic acid | cap, 250mg | DVA |
| 6 7 | Paracetamol Paracetamol | elixir 125mg/ml tab, 500 mg | HVA HVA |
| 8 | Tranexamic acid | tab, 500 mg | DEB |
| | | , 0 | |
| 2.2 | Opioid analgesics | | |
| 1 | Codeine phosphate | tab, 15 mg | DVA |
| 2 | Dihydrocodeine tartrate | tab, 30 mg | DEA |
| 3 | Morphine sulphate | strong soln, 10mg/ml | DVA |
| 4 5 | Morphine sulphate Morphine sulphate | weak soln, 5mg/5ml | DVA |
| 6 | Morphine sulphate | inj, 15 mg/mL, 1 mL amp tab, slow-release, 10 mg | DVA DVB |
| 7 | Naloxone HCI | inj, neonatal, | DEA |
| 8 | 20 mcg/mL, 2 mL amp | , | |
| 9 | Pethidine HCI | inj, 50 mg/mL, 2 mL amp | DVA |
| 10 | Tramadol | cap, 50mg | DEB |
| 2.3 | Medicines for other common | symptoms for palliative care | |
| 1 | Amitryptiline | tab, 25mg | HVA |
| 2 | Baclofen | tab, 5mg | DEB |
| 3 | Bisacodyl | tab, 5mg | DEA |
| 4 | Bupivacaine | inj, 1mg/ml ampoule | CVA |
| 5 | Carbamazepine | liquid, 100mg/5ml | HVA |
| 6 | Chlopromazine | tab, 25mg | HVA |
| 7 | Dexamethasone | inj, 4mg/ml,1 ml ampoule | DEB |
| 8 | Dexamethasone | tab, 0.5mg | DEB |
| 9 | Diazepam | inj, 5mg/ml | DVA |
| 10 | Diazepam | tab, 5mg | DVA |
| 11 | Haloperidol | tab, 0.5mg | DEA |
| 12 | Hyoscine Hydrobromide | inj, 400mcg/ml | CEB |

| 13 | Lactulose | liquid, 3.1-3.7g/5ml | CVB |
|----|----------------|----------------------|-----|
| 14 | Metoclopramide | tab, 10mg | DEB |
| 15 | Phenytoin | tab, 100mg | HVA |
| 16 | Prednisolone | tab, 5mg | DVA |
| 17 | Senna | liquid, 7.5mg/5ml | DVB |
| 18 | Gabapentine | cap, 100mg | DVA |
| 19 | Gabapentine | tab, 600mg | DVA |

3. Antiallergics and medicines used in Anaphylaxis

| 3.1 | Antihistamines | | |
|-----|---------------------------------|-----------------------------|-----|
| 1 | Betamethasone | tab 0.5mg | DEB |
| 2 | Cetirizine | tab, 10mg | DEB |
| 3 | Chlorpheniramine maleate | tab, 4 mg | HEA |
| 4 | Chlorpheniramine maleate | inj, 10mg/ml, amp | DVA |
| 5 | Promethazine | tab, 25 mg | HVA |
| 6 | Promethazine HCI | elixir, 5 mg/5 mL | DEA |
| 7 | Promethazine HCI | inj, 25 mg/mL, 2 mL amp | DEA |
| | | | |
| 3.2 | Medicines used in nasal allergy | 1 | |
| 1 | Beclomethasone dipropionate | nasal spray, 50mcg/spray | DEB |
| 2 | Oxymetazoline hydrochloride | nasal drops 0.1%,spray 0.1% | DEB |

4. Antidotes and other medicines used in poisoning

| 4.1 No | n specific | | |
|--------|-----------------------|----------------------------|-----|
| 1 | Activated charcoal | powder | DVB |
| | | | |
| 4.2 Sp | ecific | | |
| 1 | Atropine sulphate | inj, 600 mcg/mL, 1 ML amp | DVA |
| 2* | Acetylcysteine | inj, 200 mg/mL, 10 mL amp | DVB |
| 3 | Darrows Half strength | infusion, 1L | CVB |
| | | with 5% glucose | |
| 4 | Desferrioxamine | inj, 500 mg vial (PFR) | DVB |
| 5 | Dextrose | infusion, 5% 1L | HVA |
| 6 | Ipecacuanha | emetic mixture, paediatric | HEA |
| 8 | Pralidoxime mesylate | inj, 200 mg/mL, 5 mL amp | CVB |
| 9 | Sodium Bicarbonate | tab, 600mg | CVB |
| 10 | Sodium Chloride | infusion, 0.9%, 1L | HVA |

| 5. | Anticonvulsants/Antiepiletic | cs | |
|----|------------------------------|--------------------------|-----|
| 1 | Carbamazepine | tab, 200 mg | DEB |
| 2 | Diazepam | inj, 5 mg/mL, 2 mL amp | DVA |
| 3 | Ethosuximide | cap, 250 mg | CEB |
| 4* | Magnesium sulphate | inj, 500 mg/m, 2 mL amp | DVA |
| 5 | Paraldehyde | inj, 10 mL amp | HVA |
| 6 | Phenobarbitone sodium | tab, 30 mg | HVA |
| 7 | Phenobarbitone sodium | inj, 200 mg/mL, 1 mL amp | HVA |
| 8 | Pheytoin sodium | tab, 100 mg | DVA |
| 9 | Phenytoin sodium | inj, 50mg/ml, amp | DVA |
| 10 | Sodium valproate | tab, 200 mg | CVB |

6. **Anti-infective medicines**

Antihelmintics 6.1

1

Intestinal anthelmintics 6.1.1 Albendazole

| 2 | Mebendazole | tab, 500mg | HEA |
|---|-------------|------------|-----|

tab, 200 mg

HEA

Niclosamide 3 tab, chewable, 500 mg DEB

6.1.2. Antifilarials

1 Ivermectin HVB tab, 6 mg

6.1.3 Antischistosomals and other antitrematode medicines

1 Praziquantel tab, 600 mg Н

6.2 **Antibacterials**

Penicillins and cephalosporins (Beta Lactam medicines) 6.2.1.

| Amoxycillin dispersible tab, 250mg HVA Amoxycillin + clavulanic acid tab, 500 + 125mg CEA Ampicillin sodium inj, 250 mg vial PFR DVA Benzylpenicillin (=2.4 MU) Benzylpenicillin inj, 4 g vial PFR (=5 MU) Cefotaxime inj, 500 mg PFR DVA |
|---|
| 4 Amoxycillin + clavulanic acid tab, 500 + 125mg CEA 5 Ampicillin sodium inj, 250 mg vial PFR DVA 6 Benzathine/ inj, 1.44 g vial PFR HVA Benzylpenicillin (=2.4 MU) 7 Benzylpenicillin inj, 4 g vial PFR (=5 MU) HVA |
| 5 Ampicillin sodium inj, 250 mg vial PFR DVA 6 Benzathine/ inj, 1.44 g vial PFR HVA Benzylpenicillin (=2.4 MU) 7 Benzylpenicillin inj, 4 g vial PFR (=5 MU) HVA |
| 6 Benzathine/ inj, 1.44 g vial PFR HVA Benzylpenicillin (=2.4 MU) 7 Benzylpenicillin inj, 4 g vial PFR (=5 MU) HVA |
| Benzylpenicillin (=2.4 MU) Benzylpenicillin inj, 4 g vial PFR (=5 MU) HVA |
| 7 Benzylpenicillin inj, 4 g vial PFR (=5 MU) HVA |
| |
| 8 Cefotaxime inj, 500 mg PFR DVA |
| |
| 9 Ceftazidime inj, 1g vial, PFR CVB |
| 10 Ceftriaxone inj, 1g PFR DVA |
| 11 Cephalexin cap, 250 mg DVA |
| 12 Cloxacillin cap, 250mg DEA |
| 13 Flucloxacillin capsule, 250 mg DVA |
| 14 Flucloxacillin elixir, 125 mg/5 mL DVA |
| 15 Flucloxacillin inj, 250 mg vial PFR DVA |
| 16 Imipenem inj 500mg vial PFR CVA |
| 17 Imipenem+ Cilastatin inj, 250mg+250mg PFR CVA |
| 18 Meropenem inj 500mg vial PFR CVA |
| 19 Piperacillin + Tazobactam inj, 4g + 500mg vial PFR CEB |
| 20 Procaine penicillin inj, 4.8 MU vial CVB |

| 6.2.2 | Other antibacterials | | |
|-------|---------------------------------------|--|-----|
| 1 | Azithromycin | cap, 250mg | DEA |
| 2 | Azithromycin | susp, 200mg/5 mL | DEA |
| 3 | Chloramphenicol sodium Succinate | inj, 1 g vial PFR | HVA |
| 4 | Co-trimoxazole | tab, 480 mg | HVA |
| 5 | Clarithromycin | tab, 500mg | CVA |
| 6 | Clarithromycin | susp, 125mg/5ml | DVA |
| 7 | Ciprofoxacin | tab, 250 mg | DVB |
| 8 | Clindamycin | cap, 150mg | DEB |
| 9 | Doxycycline | tab, 100 mg | HVA |
| 10 | Erythromycin | tab, e/c, 250 mg base | HVA |
| 11 | Erythromycin ethyl succinate | susp, 125 mg/5 mL (of erythromycin base) | DVA |
| 12 | Gentamicin | inj, 40 mg | HVA |
| | | (as sulphate) mL, 2 mL vial | |
| 13 | Gentamicin | paed inj, 10 mg (as sulphate)/mL, 2 mL vial | DEA |
| 14 | Metronidazole | tab, 200 mg | HVA |
| 15 | Metronidazole | inj, 5 mg/mL, 100 mL vial | DVA |
| | | (Iv infusion) | |
| 16* | Metronidazole | susp, 200 mg/5 mL | DVA |
| 17 | Nalidixic Acid | tab, 500 mg | DVA |
| 18 | Neomycin | tab, 500mg | DEB |
| 19 | Nitrofurantoin | tab, 50 mg | DEA |
| 20* | Nitrofurantoin | susp, 25 mg/5 mL | DVB |
| 21 | Sodium fusidate | tab, 250 mg | CVB |
| 22 | (Sulphamethoxazole + Trimethoprim) | (400 mg + 80 mg),960mg | DVA |
| 23 | Vancomycin | inj, 1g vial PFR, | CVA |
| 24 | Vancomycin | cap, 250mg | DVB |

6.2.3 Antileprosy medicines

| 6.2.3 | Antileprosy medicines | | |
|----------|---|-------------------------------------|------------|
| 1 | Clofazimine | cap, 50 mg | DVA |
| 2 | Dapsone | tab, 100 mg | DVB |
| 3 | Rifampicin | caps, 150 mg | DVB |
| 6.2.4 | Antituberculosis medicines | | |
| 0.2.4 | Antituberculosis medicines | | |
| 1 | Amikacin | inj 1g vial PFR | CVA |
| 2 | Capreomycin(Cm) | inj 1g, vial PFR | CVA |
| 3 4 | Cycloserine Ethambutol HCI | tab 250mg tab, 400 mg | CVB DVA |
| 5 | Ethionamide | tab 250mg | CVA |
| 6 | Isoniazid (H) + ethambutol (E) | tab, 150 mg + 400 mg | DVA |
| 7 | Isoniazid (INH) | tab, 100 mg | DVA |
| 8 | Kanamycin (Km) | inj 1g vial PFR | CVA |
| 9 | Levofloxacin (Lfx) | tab, 250mg | CVA |
| 10 | Moxifloxacin (Mfx) | tab, 400mg | CVA |
| 11 | Pyrazinamide (Z) | tab, 400 mg | DVA |
| 12 | Pyrazinamide (Z)Ethambutol (E | | DVA |
| 13 | RHZE | tab,150mg+75mg+400mg+275m | |
| 14 | Rifampicin (R) + isoniazid (H) | tab, 100 mg + 50 mg | DVA |
| 15 16 | rifampicin+isoniazid RH, Streptomycin sulphate | tab, 150mg+75mg inj 5 g vial PFR | DVA DVA |
| | | | |
| Paedia | tric | | |
| 17 | RHZ + E100 | tab, 60mg+30mg+150mg, | DVA |
| 18 | RH | tab, 60mg+60mg | DVA |
| | | | |
| 6.3 | Antifungalmedicines | | |
| 1 | Amphotericin B | inj, 50mg vial PFR | CVA |
| 2 | Clotrimazole | vagl tablets , 100mg | DEA |
| 3 | Fluconazole | cap, 250 mg | DVA |
| 4 | Fluconazole | i/v infusion, 2 mg/mL, 25mL | DVA |
| 5 | Fluconazole | oral liquid, 50mg/ml | DVA |
| 6 | Gentian violet | paint, aq, 0.5%, 500 mL | HEA |

| 7 | Griseofulvin | tab, 125 mg | DEA |
|---------------------------------------|--|---|--------------------------|
| 8 | Nystatin | pessary, 100,000 units ¹ | HVA |
| | | (with applicator) | |
| 9 | Ketoconazole | tab, 200 mg | DVA |
| 10* | Ketoconazole | susp, 100 mg/5 mL | DVA |
| 11 | Miconazole Nitrate | vag tablet, 200mg | CVA |
| | | | |
| 6.4 | Antiviral medicines | | |
| 6.4.1 | Antiherpes medicines | | |
| 1 | Acyclovir | tab, 200mg | HVA |
| 2 | Acyclovir | cream, 5%, 10g | HVA |
| 3 | Ganciclovir | inj, 500mg PFR vial | CEB |
| 4 | Ganciclovir | tab, 500mg | CEB |
| | | | |
| | | | |
| 6.4.2 | Antiretrovirals | | |
| | Antiretrovirals Nucleoside/Nucleotide reverse | transcriptase inhibitors | |
| | | transcriptase inhibitors tab, 300mg | DVA |
| 6.4.2.1 | Nucleoside/Nucleotide reverse | • | DVA DVA |
| 6.4.2.1 | Nucleoside/Nucleotide reverse Abacavir (ABC) | tab, 300mg | |
| 6.4.2.1 1 2 | Nucleoside/Nucleotide reverse Abacavir (ABC) Didanosine (ddl) | tab, 300mg tab, chewable, 50mg | DVA |
| 6.4.2.1 1 2 3 | Nucleoside/Nucleotide reverse Abacavir (ABC) Didanosine (ddI) Lamivudine (3TC) | tab, 300mg tab, chewable, 50mg tab, 150mg | DVA DVA |
| 6.4.2.1 1 2 3 4 | Nucleoside/Nucleotide reverse Abacavir (ABC) Didanosine (ddl) Lamivudine (3TC) Tenofovir (TDF) | tab, 300mg tab, chewable, 50mg tab, 150mg tab, 300mg | DVA DVA DVA |
| 6.4.2.1 1 2 3 4 5 | Nucleoside/Nucleotide reverse Abacavir (ABC) Didanosine (ddl) Lamivudine (3TC) Tenofovir (TDF) Zidovudine(AZT) | tab, 300mg tab, chewable, 50mg tab, 150mg tab, 300mg tab, 300mg | DVA DVA DVA HVA |
| 6.4.2.1 1 2 3 4 5 | Nucleoside/Nucleotide reverse Abacavir (ABC) Didanosine (ddl) Lamivudine (3TC) Tenofovir (TDF) Zidovudine(AZT) Zidovudine (AZT) | tab, 300mg tab, chewable, 50mg tab, 150mg tab, 300mg tab, 300mg syrup, 50mg/5ml | DVA DVA DVA HVA |
| 6.4.2.1 1 2 3 4 5 | Nucleoside/Nucleotide reverse Abacavir (ABC) Didanosine (ddI) Lamivudine (3TC) Tenofovir (TDF) Zidovudine(AZT) Zidovudine (AZT) | tab, 300mg tab, chewable, 50mg tab, 150mg tab, 300mg tab, 300mg syrup, 50mg/5ml | DVA DVA DVA HVA |
| 6.4.2.1 1 2 3 4 5 6 | Nucleoside/Nucleotide reverse Abacavir (ABC) Didanosine (ddl) Lamivudine (3TC) Tenofovir (TDF) Zidovudine(AZT) Zidovudine (AZT) Non nucleoside/Nucleotide re | tab, 300mg tab, chewable, 50mg tab, 150mg tab, 300mg tab, 300mg syrup, 50mg/5ml | DVA DVA HVA HVA |

¹ May be used for oral thrush (pessary is sucked)

| 6.4.2.3 | Protease | inhibitors |
|---------|----------|------------|
| | | |

| 6.4.2.3 | Protease inhibitors | | |
|----------------|--|------------------------------------|-------|
| 1 | Lopinavir + Ritonavir (LPV/r) | tab 200mg +50mg | DVA |
| 6.4.2.4 | Fixed dose combinations Stavudine (d4T) + Lamivudine (3TC) | tab, 30mg +150mg+200mg | HVA |
| - | + Nevirapine (NVP) | tub, 30111g - 130111g - 200111g | 11471 |
| 2 | Stavudine (d4T) + Lamivudine (3TC) | tab, 300mg +150mg+200mg | HVA |
| | + Nevirapine (NVP) | | |
| 3 | Zidovudine (AZT) + Lamivudine(3TC) HVA+Nevirapine (NVP) | tab, 300mg +150mg+200mg | |
| 4 | Zidovudine(AZT) + Lamivudine(3TC) | tab, 300mg +150mg | HVA |
| | Stavudine(d4T)+Lamivudine(3TC) | tab, 30mg+150mg | HVA |
| 5 | Stavudine(d4T)+Lamivudine(3TC) | tab, 6mg+30mg | HVA |
| 7 | TenofovirDisoproxilFumarate+ | | HVA |
| | Lamivudine+Efavirenz(TDF+3TC +EFV) | tab, 300mg+300mg+600mg | |
| 8 | TenofovirDisoproxilFumarate+Lamivu | dine(TDF/3TC) tab, 300mg+300mg | HVA |
| 9 | Atazanavir /Ritonavir (ATV/r) | tab, 300mg +100mg | HVA |
| 6.4.2.5 | Paediatric Anti-HIV medicines | | |
| 1 | Abacavir/Lamivudine (ABC/3TC) | tab, 600+300mg | HVA |
| 2 | Abacavir/Lamivudine (ABC/3TC) | tab, 60+30mg | HVA |
| 3 | Efavirenz (EFV) | tab, 200mg | DVA |
| 4 | Lopinavir/Ritonavir (LPV/r) | susp, 80/20mg, 60 ml | DVA |
| 5 | Lopinavir/Ritonavir(LPV/r) | tab, 100+25mg | HVA |
| 6 | Nevirapine | susp,10mg/ml, | |
| | | w/syringe,25ml | HVA |
| 7 | Nevirapine | susp,10mg/ml,w/syringe, | |
| | | 100ml | HVA |
| 8 | Nevirapine (NVP) | tab, 50mg | HVA |
| | | | |

| 9 | Nevirapine (AZT+3TC+NVP) | | |
|------------------------------------|--|--|--------------------------|
| 10 | Zidovudine+Lamivudine+ | tab, 60+30+50mg | HVA |
| 11 | Zidovudine+Lamivudine (AZT+3TC) | tab, 60+30mg | HVA |
| 6.5 | Antiprotozoalmedicines | | |
| 6.5.1 | Antiamoebics and antigiardiasis med | licines | |
| 1. | Metronidazole | tab, 200 mg | HVA |
| 2* | Metronidazole | susp, 200 mg/5 | DVA |
| 3 | Praziquantel | tabl, 600 mg | HEA |
| | | | |
| 6.5.2 | Antileishmaniasis medicines | | |
| 1 | Amphotericin B | inj, 50mg/vial | CEB |
| 2 | Fluconazole | tab, 200mg | DVA |
| | | | |
| | | | |
| 6.5.2 | Antimalarials | | |
| | Antimalarials For curative treatment | | |
| | | tab, 20mg + 120mg | HVA |
| 6.5.3.1 | For curative treatment | tab, 20mg + 120mg tab, 25mg + 67.5mg | HVA DVA |
| 6.5.3.1 I | For curative treatment Artemether+ Lumefantrine | | |
| 6.5.3.1 I | For curative treatment Artemether+ Lumefantrine Artesunate + Amodiaquine | tab, 25mg + 67.5mg | DVA |
| 6.5.3.1 I 1 2 3 | For curative treatment Artemether+ Lumefantrine Artesunate + Amodiaquine Artesunate + Amodiaquine | tab, 25mg + 67.5mg tab, 50mg + 135mg | DVA DVA |
| 6.5.3.1 I 1 2 3 4 | For curative treatment Artemether+ Lumefantrine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate + Amodiaquine | tab, 25mg + 67.5mg tab, 50mg + 135mg tab, 100mg + 270mg | DVA DVA DVA |
| 6.5.3.1 I 1 2 3 4 5 | For curative treatment Artemether+ Lumefantrine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate Bicarbonate | tab, 25mg + 67.5mg tab, 50mg + 135mg tab, 100mg + 270mg inj, 60mg vial | DVA DVA DVA CVB |
| 6.5.3.1 I 1 2 3 4 5 | For curative treatment Artemether+ Lumefantrine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate Bicarbonate Artesunate Rectal | tab, 25mg + 67.5mg tab, 50mg + 135mg tab, 100mg + 270mg inj, 60mg vial Suppository, 50mg | DVA DVA CVB HVA |
| 6.5.3.1 I 2 3 4 5 6 | For curative treatment Artemether+ Lumefantrine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate Bicarbonate Artesunate Rectal Artesunate Rectal | tab, 25mg + 67.5mg tab, 50mg + 135mg tab, 100mg + 270mg inj, 60mg vial Suppository, 50mg Suppository, 100mg | DVA DVA CVB HVA |
| 6.5.3.1 I 2 3 4 5 6 7 | For curative treatment Artemether+ Lumefantrine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate Bicarbonate Artesunate Rectal Artesunate Rectal Artesunate Rectal | tab, 25mg + 67.5mg tab, 50mg + 135mg tab, 100mg + 270mg inj, 60mg vial Suppository, 50mg Suppository, 100mg Suppository, 400mg | DVA DVA CVB HVA HVA |
| 6.5.3.1 I 2 3 4 5 6 7 8 9 | For curative treatment Artemether+ Lumefantrine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate + Amodiaquine Artesunate Bicarbonate Artesunate Rectal Artesunate Rectal Artesunate Rectal Halofantrine | tab, 25mg + 67.5mg tab, 50mg + 135mg tab, 100mg + 270mg inj, 60mg vial Suppository, 50mg Suppository, 100mg Suppository, 400mg susp, 100 mg/5 mL | DVA DVA CVB HVA HVA |

6.5.3.2 For prophylaxis

1 Sulphadoxine

+ Pyrimethamine (SP) tab, 25 mg + 500 mg HVA

2 Proguanil HCI tab, 100 mg DVA

3 Mefloquine hydrochloride tab, 250mg CEB

6.5.4 Antipneumocytosis and Anti-toxoplasmosis medicines

1 Co-trimoxazole tab, 480 mg HVA

6.5.5 Antitrypanosomal medicines (African)

1 Melarsoprol B inj, 3.6% solution, 5 mL amp DVB

2 Suramin sodium inj, 1 g vial PFR DVB

| 7. | Anti-migraine medicines | | |
|-----|------------------------------|---------------------------|-----|
| 7.1 | For treatment of acute attac | ck | |
| 1 | Paracetamol | tab, 500 mg | HVA |
| 2 | Ibuprofen | tab, 200 mg | DVA |
| 3 | Morphine Sulphate | tab, 10 mg (slow release) | DVA |
| | | | |
| 7.2 | For prophylaxis | | |
| 1 | Amitriptyline | tab, 25 mg | DVA |
| 2 | Carbamazepine | tab, 200 mg | DVA |
| 3 | Propranolol HCI | tab, 40 mg | DVA |

| 8. | Antineoplastic and immuno | suppressant medicines | | |
|----------|-----------------------------------|---|------------|--|
| 8.1 | Immunosuppressive medicines | | | |
| 1 | Azathioprine | inj 50mg vial PFR, tab 50mg, | CVB | |
| 2 | Cyclosporincap, 25mg | | CVA | |
| 3 | Cyclosporincap, 100mg | | CVA | |
| 4 | Prednisolonetab, 5mg | | DVA | |
| 8.2 | Cytotoxic and adjuvant med | dicines | | |
| 1 | Actinomycin D PFR (with mannitol) | inj, 500 microgram vial | CVB | |
| 2 | Adriamycin (doxorubicin) | inj 50mg vial PFR | CVA | |
| 3 | Bleomycininj | 15 IU vial PFR | CVA | |
| 4 | Busulphan | tab, 2 mg | CVB | |
| 5 | Carboplatin | inj, 50 mg/5 ml; 5Ml | CVB | |
| 6 | Chlorambucil | tab, 2 mg | CVB | |
| 7 | Cisplatin | inj 50mg/50ml,50ml bottle | CVB | |
| 8 | Cyclophosphamide | tab, 50 mg | CVA | |
| 9 | Cyclophosphamide | inj, 200 mg vial PFR | CVA | |
| 10 | Cyclosporin | cap, 100mg | CVB | |
| 11 | Cytarabine Darcabazine | inj, 100mg/mL, 5mL | CVB | |
| 12 13 | Etoposide | inj, 100mg/mL, PFR, 200mg vial inj, 20 mg/ml, 5-ml amp | CVB CVB | |
| 14 | Fluorouracil | inj 500mg/10ml,10ml amp | CVB | |
| 15 | Hydroxyureatab,500mg | mj 500mg/10mi,10m amp | CVB | |
| 16 | Ifosphamide | inj, 1 g vial, PFR. | CVB | |
| 17 | Leucovorin | inj, 10mg/mL 5mL vial | CVA | |
| 18 | Melphalan | tab, 2 mg | CVB | |
| 19 | Methotrexate | inj, 2.5. mg/mL, 1 mL amp | CVA | |
| 20 | Methotrexate | tab, 2.5 mg | CVA | |
| 21 | Paclitaxel | inj 300mg/50ml, vial | CVB | |
| 22 | Procarbazine | cap, 50mg | CVB | |
| 23 | Taxotere (Docetaxel) | inj, 40mg/ml, 2ml vial | CVB | |
| 24 | Vinblastine | inj, 1mg/mL, PFR, 10mL vial | CVB | |
| 25 | Vincristine sulphate | inj, 1 mg vial PFR | DVA | |
| 8.3 | Hormones and antihormon | es | | |
| 1 | Anastrazole | tab1mg | CVB | |
| 2 | Bicalutamide | tab,50mg | CVB | |
| 3 | Fludrocortisone | tab, 100mcg | CVB | |
| 4 | Tamoxifen | tab, 40mg | CVB | |
| 5 | Tamoxifen | susp, 10mg/5ml | CVB | |

| 9 | Antiparkinsonism medicines | | |
|---|----------------------------|---------------------|-----|
| 1 | Benzhexol HCI | tab, 5 mg | DEA |
| 2 | Bromocriptine | tab, 2.5. mg | CEB |
| 3 | Biperiden | tab 2mg | CEB |
| 4 | Levodopa + carbidopa | tab, 250 mg + 25 mg | CEB |
| 5 | Orphenadrinetab 50mg, | | CEB |

| 10 | Modicinos affactiva the blass | J | |
|------|-------------------------------|------------------------------|-----|
| 10 | Medicines affecting the blood | ι | |
| 10.1 | Antianaemic medicines | | |
| 1 | Ferrous sulphate + | tab, 200 mg + 0.5 mg | HVA |
| | Folic acid | | |
| 2 | Ferrous sulphate | mixt, paediatric, 60 mg/5 mL | HEA |
| 3 | Folic acid | tab, 5 mg | HVA |
| 4 | Hydroxocobalamin | inj, 1 mg/mL, 1 mLamp | CVB |
| 5 | Iron sorbitol | inj, 5% (50 mg/mL) | DVB |
| | | 2 mLamp | |
| | | | |
| 10.2 | Medicines affecting coagula | tion | |
| | | | |
| 1 | Heparin sodium | inj, 5,000, IU/mL, 5 mL vial | CEB |
| 2 | Phytomenadione | inj, 1 mg/0.5 mL amp | DVA |
| 3 | Phytomenadione | inj, 10mg/mL 1mL amp | CEA |
| | | | |
| 4 | Protamine sulphate | inj, 10 mg/mL, 5 mL amp | CEB |
| 5 | Tranexamic acid | tab, 250 mg | CEB |
| 6 | Tranexamic acid | tab, 500mg | CEB |
| 7 | Warfarin sodium | tab, 1 mg | CVB |
| 8 | Warfarin sodium | tab, 5 mg | CVB |
| | | | |
| 10.3 | Medicines to treat hyperkala | aemia | |
| 1 | Potassium binding resin pow | der | DVA |
| | (sodium polystyrene sulfonat | te | |
| | Kayexalate®) | | |
| | | | |

| 11. | Blood products and plasma substi | tutes | |
|------|-----------------------------------|---------------------------|-----|
| 11.1 | Blood Products | | |
| 1 | Cryoprecipitate | 30-40mls | DVA |
| 2 | Fresh frozen plasma | 200-300mls | DVA |
| 3 | Random donor platelet concentrate | s 50-70mls | CEB |
| 4 | Red cell suspensions adult | 280-420mls | DVA |
| 5 | Red cell suspensions paediatric | size 100-200mls | DVA |
| 6 | Whole blood adult | 450mls (405-495) | DVA |
| 7 | Whole blood Paediatric | size 100-225mls | DVA |
| | | | |
| 11.2 | Plasma Derived Blood Products | | |
| 1. | Factor viii | | CEB |
| 2. | Factor ix | | CEB |
| 11.3 | Transfusion alternatives | | |
| 1. | Erythropoietin | | CEB |
| 2. | IV Iron | | CEB |
| 3. | Gelatin (as polygeline) | i/v infusion, 500 mL pack | DVA |
| | (Haemaccel ®) | | |

| 12 | Cardiovascular Medic | ines | |
|----------|---------------------------|--|------------|
| 12.1 | Antianginal Medicine | s | |
| 1 | Glyceryltrinitrate | tab (sublingual), 500 mcg | DEB |
| 2* | Isosorbidedinitrate | tab, 10 mg | DEB |
| 3 | Nifedipine | cap, 10 mg | CVB |
| 4 | Nifedipine | tab, slow-release, 20 mg | CEB |
| 5 | Propranolol HCI | tab, 40 mg | DVA |
| 12.2 | Antiarrhythmic Medio | cines | |
| 1 | Propranolol HCI | tab, 40 mg | DVA |
| 2 | Lignocaine HCI | inj, 1%, 25 mL vial | CEB |
| 3 | Verapamil | tab, 40mg | СЕВ |
| 4 | Verapami | tab(modified release), 40mg | СЕВ |
| 12.3 | AntihypertensiveMedicines | | |
| 1 2 | Amlodipine Atenolol | tab, 5mg tab, 50mg or 100mg | DVA |
| 3 | Captopril | tab, 12.5 mg | DVA CVB |
| 4 | Enalapril | tab, 2.5 mg | DVA |
| 5 | Enalapril | tab, 5 mg | DVA |
| 6 | Hydralazine HCI | inj, 20 mg amp PFR | DVA |
| 7 | Hydralazine HCI | tab, 25 mg | DEA |
| 8 | Hydrochlorothiazide | tab, 25mg | DVA |
| 9 | Methyldopa | tab, 250 mg | DEA |
| 10 | Nifedipine | cap, 10 mg | DVA |
| 11 | Nifedipine | tab, slow-release, 20 mg | DVA |
| 12 | Nimodipine | tab, 30mg | CEB |
| 13 | Nimodipine | inj,0.2mg/ml, amp | CVB |
| 14 | Prazosin | tab, 1 mg | CEB |
| 15 | Propranolol HCI | tab, 40 mg | DVA |
| 16 17 | Reserpine Reserpine | tab 250 micrograms inj, 1 mg/mL, 1 mL amp | DVA DEA |
| 12.4 | Medicines used in hea | art failure | |
| 1 | Digoxin | tab, 250 mcg | DVA |
| 2 | Digoxin | inj, 250 mcg/mL | DVA |
| | | 2 mL amp | |
| | | MEML 2015 ————— | |

| 3 | Digoxin | tab, paed, 62.5 mcg | DVA |
|------|---------------------------|----------------------------|-----|
| 4 | Digoxin | elixir, 50 mcg/mL | DVA |
| 5 | Dopamine | inj, 40mg/ml, 5ml, vial | CEB |
| 6 | Enalapril | tab, 5mg | DEB |
| 7 | Epinephrine | inj, 0.001mg/ml, 10ml, amp | DEB |
| 8 | Frusemide | tab, 40mg | HVA |
| 9 | Furosemide | inj, 10mg/ml, 5ml, amp | DVA |
| 10 | Glyceryltrinitrate | tab (sublingual), 500 mcg | DEB |
| 11 | Isosorbidedinitrate | tab, 10 mg | DEB |
| 12 | Metoclopramide | inj, 5mg/ml, 2ml, amp | DEA |
| 13 | Metolazone | tab, 5mg | CEB |
| 14 | Morphine | inj, 1mg/ml, 2ml, amp | DVA |
| 15 | Spironolactone | tab, 25 mg | DVB |
| | | | |
| 12.5 | Antithrombotic medicine | | |
| | Acetylsalicylic acid | tab, 100 mg | DVA |
| | | | |
| 12.6 | Lipid-lowering agents | | |
| | Simvastatin | tab, 40mg | CVA |
| | | | |
| 12.7 | Antihypotensive medicines | | |
| 1 | Dopamine HCI | inj, 40 mg/mL, 5 mL amp | CVB |
| 2 | | | |
| | Ephedrine sulphate | inj, 30 mgmL, 1mL amp | DVA |

13 Dermatological medicines (Topical)

| 13.1 | Antifungal Medicines | | |
|------|----------------------------------|--------------------------|------|
| 1 | Benzoic acid + | ointm, 6% + 3%, 500 g | HEA |
| - | Salicylic acid | omani, 070 + 370, 300 g | 1127 |
| 2* | Clotrimazole | cream, 1%, 20g | DEA |
| 2 | (or equivalent alternative) | Cream, 170, 20g | DEA |
| 3 | Econazole | cream, 1%,30g | DEA |
| 4 | Sodium thiosulphate | lotion, aq., 10%, 500 mL | DEA |
| | · | | |
| 5 | Miconazole | cream, 2%, 30g | DEA |
| 13.2 | Anti-infective medicines | | |
| 1 | Calamine lotion + | lotion, aqueous, 500 mL | HEA |
| | Sulphur 2% | | |
| 2 | Chloherxidinedicluconate | soln,0.2%,500ml | DVA |
| 3 | Gentian violet | paint, aq., 0.5%, 500 ml | HEA |
| 4 | Hydrogen peroxide | soln, 20 volume, 500 mL | DEB |
| 5 | lodine | soln, weak, 500 mL | DEA |
| 6 | Potassium permanganate | 3%, 500 mL | HEA |
| | Solution | (for dilution) | HEA |
| 7 | Povidone iodine | soln, 10%, 500ml | DEA |
| 8 | Salicylic acid + | ointt, 5% + 5%, 500g | HEA |
| | Sulphur | (in YSP base) | |
| 9 | Salicylic acid + | ointn, 5% + 5%, 500 g | CEB |
| | Sulphur | (In EO base) | |
| 10 | Silver sulphadiazine | cream, 1%, 500 g | CEB |
| 11 | Zinc ointment + Sulphur 5% | ointn, 500 g | DEA |
| 12 | Zinc paste compound + Sulphur 5% | paste, 500 g | DEA |

| 13.3 | Anti-inflammatories and antipruritic Medicines | | | |
|------|--|------------------------------|-----|--|
| 1 | Calamine lotion + | lotion, aq, 500 mL | HEA | |
| | Sulphur 2% | | | |
| 2 | Hydrocortisone | oint, 1%, 15 g | DEA | |
| 3 | Betamethasone | oint, 0.15, 15 g | CEB | |
| | (asvalerate) | | | |
| 4 | Calamine | lotion, aq, 500 mL | CEB | |
| 5 | Clobetasol propionate | oint, 0.05%, 30g | CVB | |
| | | | | |
| 13.4 | Medicines affecting skin differ | entiation and proliferation | | |
| 1* | Benzoyl peroxide | gel, 5%, 30 g | DEA | |
| 2 | Crude coal tar 10% | oint, 500 g | CEB | |
| 3 | Dithranol 0.5% in zinc + | paste, 500g | CEB | |
| | Salicylic acid paste | | | |
| 4 | Podophyllum resin | paint, alcoholic, 15%, 20 mL | DEA | |
| | | (compound benzoin tincture) | | |
| 5 | Salicylic acid | lotion, 5%, 500 mL | DEA | |
| | | (in alcohol 70%) | | |
| 6 | Salicylic acid | oint, 10%, 500 g | DEA | |
| 7 | Salicylic acid | oint, 20%, 500 g | CEA | |
| | (in YSP base) | | | |
| 8 | Salicylic acid 2% + | shampoo, 500 mL | DEA | |
| | Coal tar solution 15% + | (in soap spirit base) | | |
| | Sulphur 2% | | | |
| 9* | Salicylic acid | collodion, 12% | DEA | |
| 10 | Salicylic acid + | oint, 500 g | CEB | |
| | crude coal tar 5% | (in YSP base) | | |
| 11 | Zinc paste compound + paste, | 500 g | CEB | |
| | Crude coal tar 5% | | | |

13.5 Scabicides and pediculocides

| 1 | Benzyl benzoate | application, 25%, 500 mL | HEA |
|---|-----------------|--------------------------|-----|
| 2 | Lindane | cream/lotion 1% | CEB |

13.6 Other topical preparations

| 1. | Emulsifying ointment | oint, 500 g | HEA |
|----|-------------------------|-----------------------|-----|
| 2* | Ethyl Chloride | spray, 3% | HEA |
| 3* | Silver nitrate | pencil toughened, 40% | DEA |
| 4 | Yellow soft paraffin | oint, 500g | DEA |
| 5 | Zinc oxide (in EO base) | oint, 15%, 500 g | HEA |

| 14. | Diagnostic agents | | |
|------|---|---|-----|
| 14.1 | Ophthalmic Medicines | | |
| 1 | Fluorescein sodium | eye drops, 1% (Minims) | DEA |
| 14.2 | Radiocontrast media | | |
| 1. | Barium sulphate | susp oral, 98%, 340 g pack PFR, | DEA |
| 2. | Barium sulphate | susp oral for CT; 21% package btle | CEB |
| 3. | Barium sulphate | susp, for CT colonoscopy 40%;kit | CEB |
| 4. | Barium sulphate | enema, disposable, 93%, | |
| | | 400 g pack | CEB |
| 5. | Effervescent agent (carbex) | granules(oral), 25 g sachet | CEB |
| 6. | Effervescent agent (carbex) | soln(oral), bottle | CEB |
| 7. | Feridex(Ferumoxide; | MRI-TAgent) | CEB |
| 8. | Gastrografin | susp, (oral/rectal) 100g | CEB |
| 9. | GoLYTELY | electrolyte solution | |
| | | (polyethylene glycol(PEG) | CEB |
| 10. | Magnevist | inj, 0.5mmol/mL, 100mL vial | CEB |
| 11. | ProHance | (gadopentetatedimeglumine; MRI-T1 Agent) (gadoteridol 379; MRI- T1 Agent) | СЕВ |
| 12. | Resolvist | (Iron oxide anoparticles | |
| | | MRI-T2 Agent) | CEB |
| 13. | NulLytely electrolyte solution (sodium amidotrizoate+ | (polyethylene glycol(PEG) 660 g meglumineamidotrizoate) | CEB |
| 14. | Urografin (sodium amidotrizoate | 60% inj 0,08 g | CEB |
| | + meglumineamidotrizoate | 0,52 g/ml). | |
| 15. | Urografin | 76% inj | CEB |
| | (sodium amidotrizoate | 0,10 g | |
| 16. | Ultravist | 300 inj (iopromide 300 mg mL) | CEB |
| 17. | Ultravist | 370inj (iopromide 370 mg/mL) | CEB |

| 18. Visipaque | 320 inj (lodixanol 652mg/mL) | CEB |
|-----------------------------|--|-----|
| 19. +meglumineamidotrizoate | 0,66 g/ml) | |
| 20. Xenetix | 250 inj (lobitridol,250 mg iodine/mL) | CEB |
| 21. Xenetix | 300 inj,(lobitridol,300 mg iodine/mL) | CEB |
| 22. Xenetix | 350,inj (lobitridol ,350 mg iodine/mL) | CEB |
| 23. X-prep liquid | (130mg sennocides, bowel evacuant) | |
| | bottle | CEB |

15 **Disinfectants and Antiseptics** 15.1 Antiseptics 15.2 Disinfectants 1 Cetrimide + chlorhexidine soln, 15% + 1.5% HVA (For dilution) 2 Black disinfectant soln for dilution) HVA 3 Glutaraldehyde soln, buffered, 2% DEA

soln, 10%, 500mg, bottle

DVA

4

Povidone Iodine

| 16. | Diuretics | | |
|-----|--------------------------|---------------------------------------|------------|
| 1 | Bendrofluazide | tab, 5 mg | DVA |
| 2 | Furosemide Furosemide | tab, 40 mg inj, 10 mg/mL, 2 mL amp | DVA DVA |
| 4 | Hydrochlorothiazide | tab, 25 mg | DVA |
| 5 | Mannitol | inj, 20%, 250 mL bottle | DEA |
| 6 | Spironolactone | tab, 25 mg | DEA |

17. Gastrointestinal medicines

| 17.1 | Antacids and other antiulcers medicines | | | |
|--------|---|-------------------------------------|------------|--|
| 1 | Cimetidine | tab, 400 mg | CVB | |
| 2 | Bismuth chelate | liquid, 120 mg/5 mL, 560 mL | CEB | |
| 3 | Magnesium trisilicate | tab, chewable | HEA | |
| 5 | Omeprazole | tab, 10mg | DVA | |
| 4 | Ranitidine | tab, 150 mg | CEB | |
| | | (tripotassium dictratobism uthat | e) | |
| 17.2 | Antiemetics | | | |
| 1 2 | Cyclizine Metoclopramide HCI | tab, 25mg inj, 5 mg/mL, 2 mL amp | CEB DEA | |
| 3 | Promethazine HCI | tab, 25 mg | DEA | |
| 4 | Promethazine HCI | elixir, 5 mg/5 mL | DEA | |
| 5 | Promethazine HCI | inj, 25 mg/mL, 2 mL amp | DEA | |
| 6 | Prochlorperazine | tab, 5mg | CEB | |
| | | | | |
| 17.3 | Anti-inflammatory medicines | | | |
| 1 | Hydrocortisone | inj, 50mg/ml, 2ml vial | DEA | |
| | | | | |
| 17.4 | Laxatives and Cathartics | | | |
| 1 | Bisacodyl | tab, 5mg | DEA | |
| 2 | Glycerol | suppository (child) 2 g | DEA | |
| 3 | Lactulose | soln, 3.1mg/5ml | CVA | |
| 4 | Liquid paraffin | soln | HVA | |

17.5 Medicines used in diarrhoea

| 17.5.1 | Oral rehydration preparations | | | |
|---------------|--|----------|-----------------------|-----|
| 1 | Oral rehydration salts low osmolarity powder in sachet | | | |
| | | | for 1 litre | HVA |
| | (ORS) | | (WHO citrate formula) | |
| 2 | ReSoMal | | powder for 1 litre | DVA |
| 17.5.2 | Medicines for diarrhoea in chil Zinc | ldren | tab, 20mg | HVA |
| 17.5.3 | Antimotility medicines | | | |
| 1 | Codeine phosphate | tab, 15 | mg | DVA |
| 2* | Loperamide HCI | tab, 2 r | mg | DEA |
| 47.6 | | | | |
| 17.6 | Antihaemorrhoidals | | | |
| 1 | Bismuth subgallate co. | suppos | itory, 5g | DEA |
| 17.7 | Antispasmodics | | | |
| 1 | Atropine sulphate | inj, 600 |) mcg/mL | DVA |
| | | | 1 mL amp | |
| 2 | Hyoscinebutylbromide | inj, 20 | mg/mL, 1mL amp | DEA |

18. Hormones, other Endocrine Medicines and Contraceptives

| 18.1 | Adrenal hormones and synthetic substitutes | | | |
|---------|--|------------------------------|-----|--|
| 1 | Dexamethasone | inj, 5 mg/mL, 5mL vial D | | |
| | | as sodium phosphate) | | |
| 2 | Dexamethasone | tab, 500 mcg | DEA | |
| 3 | Hydrocortisone | inj, i/v, 50 mg/mL, 2 mL amp | DVA | |
| | | (as sodium succinate) | | |
| 4 | Prednisolone | tab, 5 mg | DVA | |
| 5 | Hydrocortisone acetate | tab, 20 mg | CVB | |
| 6 | Hydrocortisone acetate | inj, aq susp, | CVB | |
| | (i/m or intra-articular) | 25 mg/mL, 5 mL vial | | |
| | | | | |
| 18.2 | Contraceptives | | | |
| 18.2.1. | Oral Hormonal contraceptives Combined, low-oestrogen | | | |
| 1 | Levonorgestrel | surgical implant, 75mg | DVA | |
| 2 | Levonorgestrel | tab, 750 mcg | DEB | |
| 3 | Norgestrel + ethinylestradiol | tab, 0.3mg + 0.03mg | HVA | |
| 4 | Norgestrel | tab, 0.75mg | HVA | |
| | Progestogen – only | | | |
| | | | | |
| 18.2.2 | Injectable Hormonal contracep | tive | | |
| 1 | Medroxyprogesterone | inj, aq susp, | HVA | |
| | acetate | 150 mg/mL, 10 mL vial | | |
| | | | | |
| 18.3.3 | Intra-uterine devices (IUD) | | | |
| 1 | Copper containing IUD | wire, 176mg | DVB | |

| 18.3.4 | Barrier contraceptives | | |
|--------|-----------------------------------|----------------|-----|
| 1 | Female Condoms | pack, each | HVA |
| 2 | Male Condoms | pack, 100 | HVA |
| | Implantable Contraceptives | | |
| 1 | Levonorgestrel-releasing implant | rod, 75mg | DVA |
| 18.4 | Oestrogens | | |
| 1 | Oestrogens, conjugated | tab, 625 mcg | CEB |
| 18.5 | Insulins and other medicines used | for diabetes | |
| 1 | Glibenclamide | tab, 5 mg | DVA |
| 2 | Gliclazide | tab, 80mg | CEB |
| 3 | Insulin, soluble | inj, 100 iu/mL | DVA |
| | (Human Actrapid ®) | 10 mL vial | |
| 4 | Insulin zinc suspension | inj, 100 lu/mL | DVA |
| | (Human Monotard ®) | 10 mL vial | |
| 5 | Metformin HCI | tab, 500 mg | DVA |
| 6 | Metformin HCl | tab,850mg | DVA |
| 18.6 | Ovulation inducers | | |
| | | tab 50 mg | CED |
| 1 | Clomiphene citrate | tab, 50 mg | CEB |
| 18.7 | Progestogens | | |

1

Norethisterone

tab, 5 mg

CEA

18.8 Thyroid hormones and antithyroid medicines

| 1 | lodine | aqueous soln, oral, 30 mL | DEA |
|---|------------------|---------------------------|-----|
| | | (Lugol's iodine) | |
| 2 | Carbimazole | tab, 5 mg | CVA |
| 3 | Thyroxine sodium | tab, 100 mcg | CVA |
| 4 | Levothyroxine | tab, 0.05mg | CVB |

| 19. | Immunologicals | | |
|-------|---|--------------------------------|-----|
| 19.1 | Immunological diagnostic ager | nts | |
| 1 | Tuberculin purified | inj soln | DVA |
| | Protein derivative (PPD) | 1 mL amp/vial | |
| 19.2 | Sera and immunoglobulins | | |
| 1 | Anti D (RH _o) | inj, 250 mcg/mL, | DVA |
| | Immunoglobulin (Human) | 1 mL amp | |
| 2 | Antirabies serum | inj, 30IU/ml, 1ml, vial | DVA |
| 3 | Diphtheria antitoxin | inj, 20,000 IU/vial | DVA |
| 4 | Tetanus antitoxin | inj, 20,000 IU/vial | DVA |
| 5 | Tetanus antitoxin | inj, 1,500 IU/vial | DVA |
| 19.3 | Vaccines | | |
| 19.3. | 1 Vaccines for universal immunis | ation | |
| 1 | Diphtheria-pertissis- | inj, 20-dose (10 mL) vial | HVA |
| | tetanus (DPT) vaccine | (triple vaccine) | |
| 2 | BCG vaccine | inj, 20 dose vial PFR | HVA |
| | (adsorbed) | | |
| 3 | Measles vaccine, live | inj, 10-dose (5mL) vial | HVA |
| | | PFR | |
| 4 | Poliomyelitis vaccine | oral susp, 20-dose | HVA |
| | live | dispenser | |
| 5 | Tetanus toxoid vaccine | inj, 10 mL vial | HVA |
| 6 | Pentavalent vaccine (diphtheria, tetanus, pertussis Hepatitis B, heamophilus Influenza) | inj, 2 dose vial | HVA |
| 7 | Pneumococcal Conjugate Vaccine | e inj, 0.5mL prefilled syringe | HVA |
| 8 | Rotavirus vaccine | inj, PFR | HVA |

19.3.2 Vaccines for specific individuals

| 1 | Polyvalent Snake Antivenom | inj | DVA |
|---|----------------------------|---------------------------|-----|
| 2 | Rabies vaccine | inj, 1-dose (0.5 mL) vial | DVA |
| | | (PFR + diluent amp) | |
| 3 | Yellow fever vaccine | inj, 10-dose (5 mL) vial | CEB |
| | | (PFR + diluent amp) | |

| 20 | Muscle relaxants (peripherally acting) and cholinesterase inhibitors | | |
|----|--|-------------------------|-----|
| 1 | Alcuronium chloride | inj, 5 mg/mL, 2mL amp | CEB |
| 2 | Edrophonium chloride | inj, 10 mg/mL, 1 mL amp | CVB |
| 3 | Neostigmine | inj, 2.5 mg/mL, 1 mL | CEB |
| | methylsulphate | | |
| 4 | Suxamethonum chloride | inj, 50 mg/mL, 2 mL amp | DVA |
| 5 | Vecuronium bromide | inj, 10 mg vial PFR | CEB |

21. Ophthalmological preparations

| 21.1. | Anti-infective Agents | | |
|-------|----------------------------|----------------------------|-----|
| 1 | Acyclovir | tab, 200mg | DVA |
| 2 | Acyclovir | tab, 400mg | DVA |
| 3 | Acyclovir | oint, 3%, 4.5g | DVA |
| 4 | Amphotericin B | eye drops, 0.15% | CEB |
| 5 | Econazole | eye oint, 1% 15g | CEB |
| 5 | Erythromycin | eye oint, 0.5%, 3.5g tube | CEB |
| 6 | Ganciclovir | inj, 500mg PFR vial | CEB |
| 7 | Ganciclovir | tab, 500mg | CEB |
| 8 | Foscarnet | inj, 24mg/ml, 250ml bottle | CEB |
| 9 | Gentamicin | eye drops, 0.3%, 5 mL | HVA |
| | (as sulphate) | | |
| 10 | Idoxuridine | eye drops, 0.1%, 5 mL | CEB |
| 11 | Miconazole | eye drops, 1%, 10 mL | CEB |
| 12 | Natamycin | ophthalmic susp, 5% | CEB |
| 13 | Tetracycline HCI | eye oint, 1%, 3.5 g tube | HVA |
| | | | |
| 21.2 | Anti-inflammatory agents | | |
| 1 | Dexamethasone | eye drops, 0.1%, 5 mL | CVA |
| 2 | Emadastine | eye drops, 0.05%,5ml | CEB |
| 3 | Fluoromethalone | eye drops 0.1%, 5ml | CEB |
| 4 | Flucinoloneacetonide | implants, 0.59mg | CEB |
| 5 | Ketotifen | eye drops,0.025ml | CEB |
| 6 | Methylprednisolone acetate | inj, 40 mg/mL, 2 mL vial | CEB |
| 7 | Acetate | (for sub-conjunctival) | CED |
| 7 | Nedocromil | eye drops, 2%, 5ml | CEB |
| 8 | Prednisolone | eye drops, 0.5%, 5ml | CEB |
| 9 | Triamcinolone acetonide | inj, 40mg/ml, 1ml vial | CEB |

21.3 Local anaesthetics

| 1 | Amethocaine HCI | eye drops, 1%, 10 mL | DEA |
|--------|-----------------------------|---------------------------|-----|
| 21.4 | Miotics and antiglaucoma me | dicines | |
| 1 | Acetazolamide | tab, 250 mg | DVA |
| 2 | Acetazolamide | inj, 500 mg vial PFR | CEB |
| 3 | Bimatroprost | eye drops, 0.03%, 3ml | CEB |
| 4 | Brinzolamide | eye drops, 1%, 5ml | CEB |
| 5 | Betaxolol | eye drops, 0.25%, 0.25ml | CEB |
| 6 | Brimonidine | eye drops, 0.2%, 5ml | CEB |
| 7 | Dorzolamide | eye drops, 2%, 5ml | CEB |
| 8 | Glycerol | oral soln, 50% | CEB |
| 9 | Latanoprost | eye drops, 0.005%, 2.5ml | CEB |
| 10 | Pilocarpine HCI | eye drops, 1%, 10 mL | CVA |
| 11 | Timolol maleate | eye drops, 0.25% | CVA |
| | | 5 mL metered dose unit | |
| 12 | Travoprost | eye drops, 0.004%, 2.5ml | CEB |
| | | | |
| 21.5 | Mydriatics | | |
| 1 | Atropine sulphate | eye oint,0.5%, 3.5 g tube | DVA |
| 2 | Cyclopentolate HCI | eye drops, 0.5%, 5 mL | DEA |
| 3 | Tropicamide | eye drops, 0.5%, 5 mL | CEB |
| 4 | Phenylephrine | eye drops, 2.5% | CEB |
| 5 | Phenylephrine | eye drops, 10%,10ml | CEB |
| | | | |
| 21.6 0 | thers | | |
| 1 | Ascorbic acid | tab, 200mg | DEB |
| 2 | Carmellose | eye drops, 0.5%, 10ml | DEB |
| 3 | Sodium ascorbate | eye drops, 10% | DEB |

22 Obstetric medicines

| 22.1 | Oxytocics | | |
|---------------------------------------|---|--|--------------------------|
| 1 | Dinoprostone | vag, gel, 200 mcg/mL | CVB |
| | | 2.5. mL (500mcg) | |
| 2 | Dinoprostone | vag tab, 3 mg | CEB |
| 3 | Ergometrine maleate | inj, 500 mcg | HVA |
| 4 | Misoprostol | tab, 200mcg | DVA |
| 5 | Oxytocin | inj, 10 IU/mL, 1 mLamp | HVA |
| | | | |
| 22.2 | Antioxytocics | | |
| 1 | Nifedipine | tab, 10mg, SR | DVA |
| 22.3 | Myometrialrelaxants | | |
| 1 | • | tab 4 mg | HVA |
| | Salbutamolsulphate | tab, 4 mg | |
| 2 | Salbutamol sulphate | inj, 1 mg/mL, 5 mL amp | DVA |
| | | | |
| 22.4 | Medicines used in severe Pre- | Eclampsia and Eclampsia | |
| 22.4 1 | Medicines used in severe Pre- Magnesium sulphate | Eclampsia and Eclampsia inj, 500 mg/mL, 2 mL amp | HVA |
| | | | HVA |
| | Magnesium sulphate | | HVA |
| | Magnesium sulphate | inj, 500 mg/mL, 2 mL amp | HVA |
| 1 | Magnesium sulphate (MgSO ₄ .7H ₂ O) | inj, 500 mg/mL, 2 mL amp | HVA |
| 22.5 | Magnesium sulphate (MgSO ₄ .7H ₂ O) Medicines used in primary PPI | inj, 500 mg/mL, 2 mL amp | |
| 1 22.5 1 2 | Magnesium sulphate (MgSO ₄ .7H ₂ O) Medicines used in primary PPI Oxytocin Misoprostol | inj, 500 mg/mL, 2 mL amp H inj, 10 IU/mL, 1 mL amp tab, 200mcg | HVA |
| 1 22.5 1 | Magnesium sulphate (MgSO ₄ .7H ₂ O) Medicines used in primary PPI Oxytocin | inj, 500 mg/mL, 2 mL amp H inj, 10 IU/mL, 1 mL amp tab, 200mcg | HVA |
| 1 22.5 1 2 | Magnesium sulphate (MgSO ₄ .7H ₂ O) Medicines used in primary PPI Oxytocin Misoprostol | inj, 500 mg/mL, 2 mL amp H inj, 10 IU/mL, 1 mL amp tab, 200mcg | HVA |
| 1 22.5 1 2 | Magnesium sulphate (MgSO ₄ .7H ₂ O) Medicines used in primary PPI Oxytocin Misoprostol Medicines used in secondary I | inj, 500 mg/mL, 2 mL amp H inj, 10 IU/mL, 1 mL amp tab, 200mcg | HVA DVA |
| 1 22.5 1 2 22.6 | Magnesium sulphate (MgSO ₄ .7H ₂ O) Medicines used in primary PPI Oxytocin Misoprostol Medicines used in secondary II Amoxycillin | inj, 500 mg/mL, 2 mL amp H inj, 10 IU/mL, 1 mL amp tab, 200mcg PPH tab, 500mg | HVA DVA |
| 1 22.5 1 2 22.6 1 2 | Magnesium sulphate (MgSO ₄ .7H ₂ O) Medicines used in primary PPI Oxytocin Misoprostol Medicines used in secondary II Amoxycillin Erythromycin | inj, 500 mg/mL, 2 mL amp H inj, 10 IU/mL, 1 mL amp tab, 200mcg PPH tab, 500mg tab, 250mg | HVA DVA HVA |
| 1 22.5 1 2 22.6 1 2 3 | Magnesium sulphate (MgSO ₄ .7H ₂ O) Medicines used in primary PPI Oxytocin Misoprostol Medicines used in secondary II Amoxycillin Erythromycin Gentamicin | inj, 500 mg/mL, 2 mL amp H inj, 10 IU/mL, 1 mL amp tab, 200mcg PPH tab, 500mg tab, 250mg inj, 40 mg | HVA DVA HVA HVA |

| 23 | Peritoneal dialysis solutions | | |
|----|-------------------------------|--------------------------------|-----|
| 1 | Dianeal + dextrose | 1.5% intraperitoneal dialysis | CVB |
| | | Soln, 1 L bottle | |
| 2 | Dianeal + dextrose | 4.25% intraperitoneal dialysis | CVB |
| | | Soln, 1 L bottle | |

24 Medicines for mental and behavioural disorders

| 24.1 | Medicines used in psychotic disorders | | |
|--------|---------------------------------------|------------------------------|-----|
| 1 | Chlorpromazine HCI | tab, 100 mg | DVA |
| 2 | Chlorpromazine HCI | inj, 25 mg/mL, 2 mL amp | HEA |
| 3 | Chlorpromazine HCI | tab, 25 mg | HEA |
| 4 | Fluphenazinedecanoate | inj, oily, 25 mg/mL | DVA |
| | mp 2 mL a | Э | |
| 5 | Fluoxetine | cap, 20mg | DVA |
| 6 | Fluoxetine | tab, 10mg | DVA |
| 7 | Haloperidol | tab, 1.5 mg | DEA |
| 8 | Haloperidol | tab, 5mg | DEA |
| 9 | Haloperidol decanoate | inj, oil, 50 mg/mL, 1 mL amp | DEB |
| 10 | Risperidone | tab, 0.5 mg | DVA |
| 11 | Risperidone | tab, 1 mg | DVA |
| 12 | Risperidone | tab, 2 mg | DVA |
| 13 | Sodium Valproate | tab(crushable), 100mg | CEB |
| 14 | Sodium Valproate | tab, 500mg | CEB |
| 24.2 | Medicines used in mood diso | rders | |
| 24.2.1 | Medicines used in depressive | disorders | |
| 1 | Amitriptyline HCI | tab , 25 mg | HVA |
| 2 | Amitriptyline HCI | inj, 10 mg/mL | CVA |
| 3 | Fluoxetine | tab, 20mg | DVA |
| | | | |
| 24.2.2 | Medicines used in bipolar dis | orders | |
| 1 | Carbamazepine | tab, 200 mg | DVA |
| 2 | Sodium Valproate | tab, 200mg | CVA |

| 24.3 | Medicines for anxiety disorders | ;- | |
|--------|-------------------------------------|------------------------|-----|
| 1 | Chlordiazepoxide hydrochloride | cap, 5mg | CVA |
| 2 | Chlordiazepoxide hydrochloride | cap, 10mg | CVA |
| 3 | Chlordiazepoxide hydrochloride | cap, 25mg | CVA |
| 4 | Diazepam | inj, 5 mg/mL, 2 mL amp | DVA |
| 5 | Diazepam | tab,5 mg | DEA |
| 6 | Lorazepam | tab, 1 mg | HEA |
| | | | |
| 24.4 N | ledicines used for obsessive-comp | ulsive disorders | |
| 1 | Fluoxetine | cap, 20mg | CVA |
| | | | |
| 24.5 N | ledicines for disorders due to psyc | hoactive substance use | |
| 1 | Chlordiazepoxide hydrochloride | cap, 5mg | CVA |
| 2 | Chlordiazepoxide hydrochloride | cap, 10mg | CVA |
| 3 | Chlordiazepoxide hydrochloride | cap, 25m | CVA |
| 4 | Thiamine | inj, 100mg/ml | DEB |
| | | | |
| 24.6 C | ther Medicines for mental disorde | er | |
| 1 | Benzhexol | tab, 5mg | DEB |
| 2 | Procyclidine HCI | inj, 5 mg/mL, 2 mL amp | CEB |

CEB

3

Procyclidine HCI

tab, 5mg

25. Medicines acting on the Respiratory Tract

25.1 Antiasthmatic and medicines for chronic obstructive pulmonary disease

| 1 | Aminophylline | tab, 100 mg | HVA |
|----|---------------------|-------------------------------|-----|
| 2 | Adrenaline | inj, 1/1,000, 1 mL amp | HVA |
| 3 | Aminophylline | inj, 25 mg/mL, 10 mL amp | DVA |
| 4 | Beclomethasone | aerosol inhalation, | CVB |
| | Diproprionate | 50 mcg/dose,200-dose unit | |
| 5 | Salbutamol sulphate | tab, 4 mg | HVA |
| 6 | Salbutamol Sulphate | inj, 1 mg/mL, 5 mL amp | DVA |
| 7 | Salbutamol Sulphate | aerosol inhalation | DVA |
| | | 100 mcg/dose,200 – dose unit | |
| 8 | Salbutamol Sulphate | respirator soln, 1 mg/mL | DVA |
| | | single dose nebuliser amps | |
| 9 | Sodium cromoglycate | spincap, 20 mg | CEB |
| | | (for use with an insufflator) | |
| 10 | Theophylline | tab, 250mg | CEB |

26. Solutions correcting water, electrolytes and acid-based disturbances

| 26.1 | Oral preparations | | |
|------|---|-------------------------------|-----|
| 1. | Oral rehydration salts, | powder in sachet for I L | HVA |
| | (ORS) | (WHO low osmolarity) | |
| 2 | Potassium chloride | tab, slow release, 600 mg | DVA |
| 26.2 | Parenteral preparations | | |
| 11 | Dextrose Normal Saline (D5NS) | i/v infusion, 0.9% NaCl + | |
| | | 5% dextrose, 1L pack | DEB |
| 1 | Glucose (dextrose) | inj, 50%, 20 mL amp | HVA |
| 2 | Glucose (dextrose) | i/v infusion, 5%, 1L pack | HVA |
| 3 | Glucose (dextrose) | i/v infusion, 10%, 100 L pack | DVA |
| 4 | Sodium lactate comp | i/v infusion, 1L pack | HVA |
| | (Ringer-lactate or Hartmann's solution) | | |
| 5 | Water for injections | for i/v use, 10 mL amp | HVA |
| 6 | Potassium chloride | inj, 20%, 10 mL amp | CVA |
| 7 | Sodium bicarbornate | inj, 4%, 50 mL vial | DVA |
| 8 | Sodium chloride | i/v infusion, 0.9%, 1L pack | HVA |
| 9 | Sodium lactate + glucose | i/v infusion, 1L (adult) pack | DEB |
| | (Darrow's ½ strength in dextrose 5%) | | |
| 10 | Sodium lactate + glucose | i/v infusion, 200 mL (paed) | DVA |

| 27 | Vitamins and minerals | | |
|----|---------------------------|------------------------|-----|
| 1 | Vitamin A | cap, 200,000 IU | HVA |
| | | (liquid or gel filled) | |
| 2 | Vitamin B Co. strong | tab | HEA |
| 3 | Nicotinamide | tab, 50 mg | DEA |
| 4 | Pyridoxine HCI | tab, 20 mg | HVA |
| 5 | Thiamine | inj, 100mg/ml | DEB |
| 6 | Vitamins, multiple | syrup | HEA |
| 7 | Vitamins, multiple | tab | HEA |
| 8 | Calcium gluconate | tab, chewable, 500 mg | CEB |
| | | | |
| 9 | Vitamins, multiple | inj, i/v, high-potency | DEB |
| | | 10 mL (in 2 amps) | |
| 10 | Calciferol, high-strength | tablet, 10,000 IU | CVB |
| 11 | Calcium gluconate | inj, 10%, 10 mL, amp | CVB |
| 12 | Vitamin B ₁₂ | tab, 0.005mg | HEA |
| 13 | Vitamin K | tab, 10mg | DEB |
| 14 | Vitamin K | inj, 10mg/ml, 1mL amp | DEB |

28. Ear, Nose and throat medicines in Children

| 28.1 | Preparations for the ear | | |
|------|---------------------------------|------------------------------|------------|
| 1 | Acetic acid | ear drops, 2% | HEA |
| 2 | Ampicillin | cap, 250mg | DVA |
| 3 | Amoxycillin | cap, 250mg | HVA |
| 4 | Betamethasone | oint, 0.15, 15 g | CEB |
| 5 | Beclomethasonedipropionate | nasal spray, 50mcg/spray | DEB |
| 6 | Ciprofloxacin | eardrops CVA | |
| 7 | Chloramphenicol | ear drops, 5%, 10 mL | DVA |
| 8 | Ceftriaxone | inj, 1g PFR | DVA |
| 9 | Cloxacillin | cap, 250mg | DEA |
| 10 | Cetirizine Erythromycin | tab, 10mg tab, 250mg | DEB HVA |
| 11 | Flucloxacillin | cap, 250 mg | DVA |
| 12 | Metronidazole | tab, 200mg | HVA |
| 13 | Hydrocortisone | oint, 1%, 15 g | DEA |
| 14 | Liquid paraffin | soln HVA | |
| 15 | Oxymetazoline hydrochloride | nasal drops 0.1%, spray 0.1% | DEB |
| 28.2 | Preparations for the oropharynx | | |
| 1 | Gentian violet | paint, aq., 0.5%, 500 mL | HEA |
| 2 | Nystatin | oral susp | DEA |
| | | 100,000 IU/mL, 20 mL | |
| | | (with graduated dropper) | |

29. Specific Medicines for Neonatal Care 29.1 Medicines administered to the neonate 1. Chlorhexidine digluconate soln, 7.1%, HVA 29.2 Medicines administered to the mother 1. Betamethasone sodium inj, 4mg/ml, 1ml amp CEB phosphate 2. Dexamethasone inj, 4mg/ml, 2ml vial DVA

30. Medicines for Arthritis

30.1 Medicines used to treat gout

| 1 | Allopurinol | tab, 100 mg | DEA |
|----|-------------------------------------|---------------------------|-----|
| 2 | Colchicine | tab, 500mcg | DVB |
| | | | |
| | | | |
| 31 | 1 Medicines used to treat Nutrition | ial Disorders | |
| 1. | Albendazole | tab, 200 mg | HEA |
| 2. | Amoxicillin | cap, 250 mg | HVA |
| 3. | Benzylpenicillin | inj, 4 g vial PFR (=5 MU) | HVA |
| 4. | Cotrimoxazole | tab, 480 mg | HVA |
| 5. | Ciprofloxacillin | tab,250mg | DVA |
| 6. | Ciprofloxacillin | tab, 500mg | DVA |
| 7. | Dextrose 50% | inj, 50%, 50mL | DVA |
| 8. | Folic Acidtab, 5 mg | | HEA |
| 9. | Furosemide | inj, 10mg/ml, 5ml, amp | DVA |
| 10 | Gentamycininj, 40 mg | | HVA |
| | (as sulphate) n | nL, 2 mL vial | |
| 11 | . Iron +Folic Acid | tab, 200 mg | HEA |
| 12 | Metronidazole | tab, 200 mg | HVA |
| 13 | Mebendazole | tab, 500mg | HEA |
| 14 | Nicotinamide | tab, 50 mg | DEA |
| 15 | . Potassium chloride | tab, slow release, 600 mg | DVA |
| 16 | ReSoMal | powder for 1 litre | DVA |
| 17 | . Vitamin A | cap, 100 000iu | HEA |
| 18 | Vitamin A | cap, 200,000IU | HEA |
| 19 | Vitamins, | syrup, multiple vitamins | HEA |
| | | | |

31.2 Non-medicinal therapeutic products used to treat Nutritional Disorders

Ready to use therapeutic food (RUTF) 260g DVA

Ready to use therapeutic food (RUTF) sachet, 92 g DVA

Corn Soya Blend++

LikuniPhala DVA

F-75 milk based formulation DVA

F-100 milk based formulation DVA

MEML Index

Abacavir (ABC) Tablet, 11
Abacavir/Lamivudine (ABC/3TC) Tablet,
Abacavir/Lamivudine (ABC/3TC) Tablet,
Acetazolamide Injection, 38
Acetazolamide Tablet, 38
Acetic acid Ear Drops, 46
Acetylcysteine Injection, 6

Acetylsalicylic acid Tablet, 21
Actinomycin D Injection, 16
Activated charcoal Powder, 6

Acyclovir Tablet, 37 Acyclovir Cream, 11 Acyclovir Ointment, 37 Acyclovir Tablet 11 Acyclovir Tablet, 37 Adrenaline Injection, 43

Adriamycin (doxorubicin) Injection, 16

Albendazole Tablet, 48 Albendazole Tablet, 8

Alcuronium chloride Injection, 36

Allopurinol Tablet, 48

Amethocaine HCI Eye drops, 38

Amikacin Injection, 9

Aminophylline Injection, 43
Aminophylline Tablet, 43
Amiodarone Injection, 2
Amitriptyline Tablet, 15
Amitriptyline HCI Injection, 41
Amitriptyline HCI Tablet, 41
Amitryptiline Tablet, 3
Amlodipine tablet, 20
Amoxicillin Capsule, 48

Amoxycillin dispersible Tablet, 8 Amoxycillin + clavulanic acid Tablet, 8

Amoxycillin Capsule, 46 Amoxycillin Elixir, 8 Amoxycillin Tablet, 39 Amoxycillin Tablet, 8

Amphotericin B Injection, 13 Amphotericin B Eye drops, 37 Amphotericin B Injection, 9 Ampicillin Capsule, 46

Ampicillin sodium Injection, 8

Anastrazole Tablet,16 Anti D (RH_o) Injection, 34 Antirabies serum Injection, 34

Artemether+ Lumefantrine Tablet, 13

Artesunate + Amodiaquine Tablet, 13 Artesunate + Amodiaquine Tablet, 13 Artesunate Bicarbonate Injection, 13

Artesunate + Amodiaguine Tablet, 13

Artesunate Rectal Suppository, 13 Artesunate Rectal Suppository, 13 Artesunate Rectal Suppository, 13

Ascorbic acid Tablet, 38

Aspirin Tablet, 3

Atazanavir /Ritonavir (ATV/r) Tablet, 12

Atenolol Tablet, 20 Atracurium Injection, 1

Atropine sulphate Eye ointment, 38
Atropine sulphate Injection, 30
Atropine sulphate Injection, 6
Atropine sulphate Injection, 2
Azathioprine Injection, 16
Azithromycin Capsule, 9
Azithromycin Suspension, 9

Baclofen Tablet, 3

Barium sulphate Suspension oral, 25 Barium sulphate Enema, disposable, 25 Barium sulphate Suspension oral, 25 Barium sulphate Suspension, 25 BCG vaccine Injection, 34

Beclomethasone Inhalation aerosol,43

Beclomethasonedipropionate

Nasal Spray, 46

Beclomethasonedipropionate

Nasal Spray, 5

Bendrofluazide Tablet, 28
Benzathine Injection, 8
Benzhexol HCI Tablet, 17
Benzhexol Tablet, 42
Benzoic acid +Salicylic acid

Ointment, 22

Benzoyl peroxide Gel, 23
Benzyl benzoate Application, 24
Benzylpenicillin Injection, 8
Benzylpenicillin Injection, 48
Betamethasone Ointment, 23
Betamethasone Ointment, 46
Betamethasone Tablet, 5

Betamethasone sodium Injection, 47

Betaxolol Eye drops,38 Bicalutamide Tablet,16 Bimatroprost Eye drops,38 Biperiden Tablet, 17 Bisacodyl Tablet, 29 Bisacodyl Tablet, 3

Bismuth chelate Liquid, 29

Bismuth subgallate co. Suppository, 30

Black disinfectant Solution, 27 Bleomycininj International Units, 16

Brimonidine Eye drops, 38 Brinzolamide Eye drops, 38 Bromocriptine Tablet, 17

Bupivacaine (heavy) +glucose Injection, 1

Bupivacaine Injection, 3

Bupivacaine(plain) Injection, 1

Busulphan Tablet, 16

Calamine Aqueous, lotion, 23
Calamine lotion + Sulphur

Aqueous, lotion, 22 Calamine lotion + Sulphur Aqueous, lotion, 23

Calciferol, high-strength Tablet, 45
Calcium gluconate Injection, 45
Calcium gluconate Tablet, 45
Capreomycin(Cm) Injection, 9

Captopril Tablet, 20
Carbamazepine Liquid, 3
Carbamazepine Tablet, 15
Carbamazepine Tablet, 41
Carbamazepine Tablet, 7
Carbimazole Tablet, 33
Carboplatin Injection, 16
Carmellose Eye drops, 38
Cefotaxime Injection, 8
Ceftazidime Injection, 8
Ceftriaxone Injection, 8
Ceftriaxone Injection, 8

Cetrimide + chlorhexidine Solution, 27

Chloherxidinedicluconate

Cephalexin Capsule, 8

Cetirizine Tablet, 46

Cetirizine Tablet, 5

Solution, 22

Chlopromazine Tablet, 3
Chlorambucil Tablet, 16
Chloramphenicol Ear Drops, 46
Chloramphenicol Sodium

Succinate Injection, 10

Chlordiazepoxide hydrochloride

Capsule, 42

Chlorhexidine digluconate

Solution, 47

Chlorpheniramine maleate

Tablet, 5

Chlorpheniramine maleate

Injection, 5

Chlorpromazine HCl Injection, 41 Chlorpromazine HCl Tablet, 41 Chlorpromazine HCl Tablet, 41

Cimetidine Tablet, 29
Ciprofloxacillin Tablet, 48
Ciprofloxacillin Tablet, 48
Ciprofloxacin Ear Drops, 46
Ciprofoxacin Tablet, 10
Cisplatin Injection, 16

Clarithromycin Suspension, 10 Clarithromycin Tablet, 10 Clindamycin Tablet, 10

Clobetasol propionate Ointment, 23

Clofazimine Capsule, 9

Clomiphene citrate Tablet, 32
Clotrimazole Cream, 22
Clotrimazole Tablets Vagl, 9
Cloxacillin Capsule, 46
Cloxacillin Capsule, 8
Co-trimoxazole Tablet, 14
Co-trimoxazole Tablet, 10
Codeine phosphate Tablet, 30
Codeine phosphate Tablet, 3

Colchicine Tablet, 48

Copper containing IUD wire, 31

Corn Soya Blend++ , 49 Cotrimoxazole Tablet, 48 Crude coal tar Ointment, 23 Cryoprecipitate Suspension, 19

Cyclizine Tablet, 29

Cyclopentolate HCI Eye drops, 38 Cyclophosphamide Injection, 16

Cyclophosphamide Tablet, 16

Cycloserine Tablet, 9
Cyclosporin Capsule, 16
Cyclosporin Capsule, 16
Cyclosporin Capsule, 16
Cytarabine Injection, 16
Dapsone Tablet, 9

Darcabazine Injection, 16

Darrows Half strength Infusion, 6

Desferrioxamine Injection, 6
Dexamethasone Eye drops, 37
Dexamethasone Injection, 3
Dexamethasone Injection, 31
Dexamethasone Injection, 47
Dexamethasone Tablet, 3
Dexamethasone Tablet, 31

Dextrose Infusion, 6
Dextrose Injection, 48
Dextrose Normal Saline
(D5NS) i/v infusion, 44

Dianeal + dextrose Solutionn, 40
Dianeal + dextrose Solution, 40

Diazepam Injection, 2
Diazepam Injection, 3
Diazepam Injection, 42
Diazepam Injection, 7
Diazepam Tablet, 2
Diazepam Tablet, 3
Diazepam Tablet, 42
Diclofenac sodium Tablet, 3

Didanosine (ddl) Tablet, 11

Digoxin Injection, 20 Digoxin Elixir, 21 Digoxin Tablet, 21 Digoxin Tablet, 20

Dihydrocodeine tartrate Tablet, 3 Dinoprostone Gel, vag, 39 Dinoprostone Tablet vag, 39 Diphtheria antitoxin Injection, 34 Diphtheria-pertissis- Injection, 34

Diproprionate, 43 Dithranol 0.5% in zinc

+ Salicylic acid paste Paste, 23 Dobutamine Injection, 2 Dopamine HCI Injection, 21 Dopamine Injection, 21 Dorzolamide Eye drops, 2%, Doxycycline Tablet, 10 Econazole Cream, 22 Econazole Eye Ointment, 37

Edrophonium chloride Injection, 36

Efavirenz (EFV) Tablet, 11 Efavirenz (EFV) Tablet, 12 Effervescent agent (carbex)

Granules(oral), 25

Effervescent agent (carbex)

Solution(oral), 25

Emadastine Eye drops, 37
Emulsifying ointment Ointment, 24

Enalapril Tablet, 20 Enalapril Tablet, 20 Enalapril Tablet, 21 Ephedrine Injection, 2

Ephedrine sulphate Injection, 21

Epinephrine Injection, 2 Epinephrine Injection, 21

Ergometrine maleate Injection, 39 Erythromycin Eye Ointment, 37 Erythromycin ethyl Suspension, 9

Erythromycin Tablet, 9 Erythromycin Tablet, 39 Erythromycin Tablet, 46

Erythropoietin, 19 eserpine Tablet, 20 Ethambutol HCI Tablet, 9 Ethionamide Tablet, 9 Ethosuximide Capsule, 7 Ethyl Chloride Spray, 24 Etoposide Injection, 16

F-100 milk based formulation , 49 F-75 milk based formulation, 49 Female Condoms Pack, 32 Fentanyl Injection, 1

Feridex(Ferumoxide; MRI-TAgent), 25 Ferrous sulphate +

Ferrous sulphate Mixture,

Paediatric, 18

Flucinoloneacetonide

Implants, 37

Flucloxacillin Capsule, 46
Flucloxacillin Capsule, 8
Flucloxacillin Elixir, 8
Flucloxacillin Injection, 8
Fluconazole Tablet, 13
Fluconazole Intravenous, 9

Fluconazole Oral Liquid, 9
Fluconazole Capsule, 9
Fludrocortisone Tablet,16
Fluorescein sodium Eye drops, 25
Fluoromethalone Eye drops, 37
Fluorouracil Injection, 16
Fluoxetine Tablet, 41
Fluoxetine Capsule, 42
Fluoxetine Tablet, 41

Injection, 41

Folic acid Tablet, 18
Folic acid Tablet, 18
Folic Acid Tablet, 48
Foscarnet Injection, 37
Fresh frozen plasma
Suspension,19

Frusemide Tablet, 21
Furosemide Injection, 21
Furosemide Injection, 48
Furosemide Tab, 40 mg
Furosemide Injection, 28
Gabapentine Tablet, 4
Gabapentine Capsule, 4
gadopentetatedimeglumine;

MRI-T1 Agent), 25 Ganciclovir Injection, 11 Ganciclovir Injection, 37 Ganciclovir Tablet, 11 Ganciclovir Tablet, 37

Gastrografin Suspension, 25

Gelatin (as polygeline) Intravenous Infusion, 19 Gentamicin Eye drops, 37 Gentamicin Injection, 10 Gentamicin Injection, 39

Gentamicin Paediatric Injection, 10

Gentamycin Injection, 48

Gentian violet Aqueous, paint, 22 Gentian violet Paint, Aqueous, 9 Gentian violet Paint, Aqueous, 46

Glibenclamide Tablet, 32 Gliclazide Tablet, 32

Glucose (dextrose) i/v infusion, 44 Glucose (dextrose) i/v infusion, 44 Glucose (dextrose) Injection, 44 Glutaraldehyde Solution, 27 Glycerol Oral solution, 38 Glycerol Suppository ,29 Glyceryltrinitrate Tablet, 21 Glyceryltrinitrate Tablet, 20 GoLYTELY Solution electrolyte, 25

Griseofulvin Tablet, 11

Haemaccel ®,19

Halofantrine Suspension, 13 Haloperidol Tablet, 41

Haloperidol decanoate

Injection, 41

Haloperidol Tablet, 3
Haloperidol Tablet, 41
Halothane Inhalation, 1
henylephrine eye drops,
Heparin sodium Injection, 18

Human Actrapid ®, 32 Human Monotard ®, 32 Hydralazine HCI Injection, 20 Hydralazine HCI Tablet, 20 Hydrochlorothiazide Tablet, 20 Hydrochlorothiazide Tablet, 28 Hydrocortisone Injection, 31 Hydrocortisone Ointment, 23 Hydrocortisone Ointment, 46

Hydrocortisone acetate Injection, Aqueous Suspension, 31

Hydrocortisone acetate

Injection, 31

Hydrogen peroxide

Solution, 22

Hydroxocobalamin

Injection, 18,

Hydroxyureatab, 16 Hyoscinebutylbromide

Injection, 30

HyoscineHydrobromide

Injection, 3

Ibuprofen Tablet, 15 Ibuprofen Tablet, 3 Idoxuridine Eye drops, 37 Ifosphamide Injection, 16 Imipenem Injection, 8

Imipenem+ Cilastatin Injection, 8

Indomethacin Tablet, 3

Insulin zinc suspension Injection, 32

Insulin, soluble Injection, 32

Iodine Solution, 22

Iodine Solution aqueous, oral, 33

Ipecacuanha Emetic Mixture, Paediatric, 6 Iron +Folic Acid Tablet, 48

Iron IV, 19

Iron sorbitol Injecton, 18
Isoflurane Inhalation, 1
Isoniazid (H) + ethambutol

(E) Tablet, 9

Isoniazid (INH) Tablet, 9 Isosorbidedinitrate Tablet, 21 Isosorbidedinitrate Tablet, 20

Ivermectin Tablet, 8

Kanamycin (Km) Injection, 9

Kayexalate®),18

Ketamine HCI Injection, 1
Ketoconazole Suspension, 11
Ketoconazole Tablet, 11
Ketotifen Eye drops, 37
Lactulose Liquid, 4
Lactulose Solution, 29
Lamivudine (3TC) Tablet, 11
Lamivudine+Efavirenz(TDF+3TC

+EFV) Tablet, 12

Latanoprost Eye drops, 38 Leucovorine Injection, 16

Levodopa + carbidopa Tablet, 17

Levofloxacin (Lfx) Tablet, 9

Levonorgestrel Surgical implant, 31

Levonorgestrel Tablet, 31 Levonorgestrel-releasing

implant Rod, 32

Levothyroxine Tablet,33 Lignocaine HCI Injection, 20 Lignocaine HCI + adrenaline

dental cartridges , 1 Lignocaine HCI + glucose

Injection, 1

Lignocaine HCI gel, 1 Lignocaine HCI Injection, 1 Lignocaine HCI Injection, 1 Lignocaine HCI spray, 1

LikuniPhala, 49

Lindane Cream/lotion,24 Liquid paraffin Solution, 29 Liquid paraffin Solution, 46 Loperamide HCI Tablet, 30 Lopinavir + Ritonavir (LPV/r)

Tablet, 12

Lopinavir/Ritonavir (LPV/r)

Suspension, 12

Lopinavir/Ritonavir(LPV/r)

Tablet, 12

Lorazepam Tablet, 42 Lorazepam Injection, 2

Magnesium sulphate Injection, 7 Magnesium sulphate Injection, 39 Magnesium trisilicate Tablet, 29

Magnevist Injection, 25 Male Condoms Pack 32 Mannitol Injection, 28

Measles vaccine, live Injection, 34

Mebendazole Tablet, 48 Mebendazole Tablet, 8

Medroxyprogesterone Injection

, Aqueous Suspension, 31 Mefenamic acid Capsule, 3

Mefloquine hydrochloride Tablet, 14

Melarsoprol B Injection, 14

Melphalan Tablet,16
Meropenem Injection, 8
Metformin HCl Tablet, 32
Metformin HCl Tablet, 32
Methotrexate Injection, 16
Methotrexate Tablet, 16
Methoxamine HCl Injection, 21

Methyldopa Tablet, 20

Methylprednisolone acetate

Injection, 37

Metoclopramide Injection, 2 Metoclopramide HCI Injection, 29 Metoclopramide Injection, 21 Metoclopramide Tablet, 4 Metolazone Tablet, 21 Metronidazole Injection, 10 Metronidazole Suspension, 10 Metronidazole Tablet, 39 Metronidazole Tablet, 46 Metronidazole Tablet, 10 Metronidazole Suspension, 13 Metronidazole Tablet, 13 Metronidazole Tablet, 48 Miconazole Cream, 22 Miconazole Eye drops, 37

Miconazole Nitrate Tablet Vag, 11

Midazolam Injection, 2 Misoprostol Tablet, 39 Misoprostol Tablet, 39 Morphine Injection, 21

Morphine sulphate Injection, 3 Morphine sulphate Tablet,

Slow-Release, 3

Morphine sulphate Injection, 1 Morphine sulphate Injection, 2 Morphine sulphate Solution

Strong, 3

Morphine Sulphate Tablet, 15 Morphine sulphate Tablet, 2

Morphine sulphate Weak Solution, 3

Moxifloxacin (Mfx) Tablet, 9 Nalidixic Acid Tablet, 10 Naloxone HCI Injection

Neonatal, 3

Natamycin Suspension

ophthalmic, 37

Nedocromil Eye drops, 37
Neomycin Tablet, 10
Neostigmine Injection, 1
Neostigmine Injection, 36
Nevirapine Suspension, 12
Nevirapine Suspension, 12
Nevirapine (AZT+3TC+NVP),12
Nevirapine (NVP) Tablet, 11
Nevirapine (NVP) Tablet, 12
Nevirapine Suspension, 11

Niclosamide Tablet, 8
Nicotinamide Tablet, 48
Nicotinamide Tablet, 45
Nifedipine Capsule 20
Nifedipine Capsule, 20
Nifedipine Tablet, 20
Nifedipine Tablet, 20
Nifedipine Tablet, 39
Nimodipine Injection, 20
Nimodipine Tablet, 20
Nitro glycerine Injection, 2
Nitrofurantoin Suspension, 10
Nitrofurantoin Tablet, 10

Nitrous oxide medical gas ,1

Norgestrel + ethinylestradiol

Norethisterone Tablet, 32

Tablet, 31

Norgestrel Tablet, 31

NulLytely electrolyte solution (polyethylene glycol(PEG), 25 Nystatin Oral Suspension, 46

Nystatin Pessary, 11

Oestrogens, conjugated Tablet, 32

Omeprazole Tablet, 29

Oral rehydration salts Powder, low

osmolarity in sachet ,30

Oral rehydration salts, Powder, 44

Orphenadrine Tablet, 17

Oxygen Inhalation (medicinal gas),1

Oxymetazoline hydrochloride

Nasal Drops, 46

Oxymetazoline hydrochloride

Nasal Drops, 5

Oxytocin Injection, 39
Oxytocin Injection, 39
Oxytocin Injection, 39
Paclitaxel Injection, 16
Paracetamol Tablet, 15
Paracetamol Elixir, 3
Paracetamol Tablet, 3
Paraldehyde Injection, 7

Pentavalent vaccine Injection, 34

Pethidine HCI Injection, 2 Pethidine HCI Injection, 3 PethidinePropofol Injection, 1 Phenobarbitone sodium

Injection, 7

Phenobarbitone sodium

Tablet, 7

Phenylephrine eye drops, 38 Phenylephrine Injection, 2 Phenytoin sodium Injection, 7

Phenytoin Tablet, 4
Pheytoin sodium Tablet, 7
Phytomenadione Injection, 18
Phytomenadione Injection, 18
Pilocarpine HCI Eye drops, 38
Piperacillin + Tazobactam

Injection, 8

Pneumococcal Conjugate Vaccine Injection, 34 Podophyllum resin Alcoholic paint, 23 Poliomyelitis vaccine

Injection, 34

Polyvalent Snake

Antivenom Injection, 35 Potassium binding resin

powder,18

Potassium chloride Injection, 44 Potassium chloride Tablet, 48 Potassium chloride Tablet,

slow release, 44

Potassium permanganate

Solution, 22

Povidone iodine Solution, 22 Povidone Iodine Solution, 27 Pralidoximemesylate Injection, 6

Praziquantel Tablet, 13
Praziquantel Tablet, 8
Prazosin Injection, 20
Prednisolone Eye drops, 37
Prednisolone Injection, 31
Prednisolone Tablet, 4
Prednisolonet Tablet, 16
Procaine penicillin Injection, 8

Prochlorperazine Tablet, 29 Procyclidine HCI Injection, 42 Procyclidine HCI Tablet, 42

Procarbazine Capsule, 16

Proguanil HCI Tablet, 14
ProHance (gadoteridol 379;

MRI- T1 Agent). 25
Promethazine Tablet, 2
Promethazine HCI Injection, 5
Promethazine HCI Elixir, 2
Promethazine HCI Elixir, 29
Promethazine HCI Elixir, 5
Promethazine HCI Injection, 2

Promethazine HCI Injection, 29 Promethazine HCI Tablet, 29 Promethazine Tablet, 5

Propanol Injection, 1

Propranolol HCI Tablet, 20 Propranolol HCI Tablet, 20 Propranolol HCI Tablet, 20 Propranolol HCI Tablet, 15 Protamine sulphate Injection, 18

Pyrazinamide (Z) Tablet, 9 Pyrazinamide (Z)Ethambutol

(E) Tablet 9

Pyridoxine HCI Tablet, 45

Quinine dihydrochloride Injection, 13

Quinine sulphate Tablet, 13

Rabies vaccine Injection, 35 Random donor platelet concentrates Suspension, 19

Ranitidine

(tripotassium dictratobism uthate)

Tablet, 29

Ready to use therapeutic food

(RUTF) sachet, 49

Ready to use therapeutic food

(RUTF), 49

Red cell suspensions adult

Suspension, 19

Red cell suspensions paediatric

Suspension,19

Reserpine Injection, 20

Resolvist (Iron oxide anoparticles

MRI-T2 Agent), 25 ReSoMal Powder, 48 ReSoMal Powder, 30

RH Tablet, 9

RHZ + E100 Tablet, 9

RHZE Tablet, 9

Rifampicin (R) + isoniazid

(H) Tablet, 9

Rifampicin Capsule, 9 rifampicin+isoniazid RH,

Tablet, 9

Risperidone Tablet, 41
Risperidone Tablet, 41
Risperidone Tablet, 41
Ropivacaine Injection, 1
Rotavirus vaccine Injection, 34

Salbutamol Sulphate

Aerosol inhalation, 43

Salbutamol sulphate

Injection, 39

Salbutamol Sulphate

Injection, 43

Salbutamol Sulphate

Respirator Solution, 43

Salbutamol sulphate

Tablet, 43

Salbutamolsulphate

Tablet, 39 Salicylic acid Ointment, 23 Salicylic acid Ointment, 23

Salicylic acid Lotion, 23

Salicylic acid + Sulphur (in YSP base) Ointment, 22 Salicylic acid +crude coal tar

Ointment, 23

Salicylic acid +Sulphur

(In EO base) Ointment, 22

Salicylic acid 2% + Coal

tar solution Shampoo, 23

Salicylic acid Collodion, 23

Senna Liquid, 4
Silver nitrate Pencil
toughened, 24
Silver sulphadiazine

Cream, 22

Simvastatin Tablet, 21 sodium amidotrizoate

+ meglumineamidotrizoate, 25 Sodium ascorbate Eye drops, 38 Sodium Bicarbonate Tablet, 6

Sodium bicarbornate

Injection, 44

Sodium chloride i/v infusion, 44 Sodium Chloride Infusion, 6 Sodium citrate Solution, 2

Sodium cromoglycate Spincap, 43

Sodium fusidate Tablet, 10 Sodium lactate + glucose

i/v infusion, 44

Sodium lactate + glucose

i/v infusion, 44 Sodium lactate comp i/v infusion, 44 Sodium thiosulphate Aqueous, lotion, 22

Sodium Valproate Tablet, 41 Sodium Valproate Tablet, 41 Sodium valproate Tablet, 7 Sodium Valproate Tablet, 41 Spironolactone Tablet, 21 Spironolactone Tablet, 28 Stavudine (d4T) + Lamivudine

(3TC Tablet, 12

Stavudine (d4T) + Lamivudine

(3TC) Tablet, 12

Stavudine(d4T)+Lamivudine

(3TC) Tablet, 12

Stavudine(d4T)+Lamivudine

(3TC) Tablet, 12

Streptomycin sulphate Injection, 9 Sulphadoxine + Pyrimethamine (SP)

Tablet, 14

Sulphamethoxazole +Trimethoprim,10

Suramin sodium Injection, 14 Suxamethanium Injection, 1 Suxamethonum chloride

Injection, 36

Tamoxifen Suspension, 16 Tamoxifen Tablet, 16 Taxotere (Docetaxel)

Injection, 16

Tenofovir (TDF) Tablet, 11
TenofovirDisoproxilFumarate+

Tablet. 12

TenofovirDisoproxilFumarate +Lamivudine(TDF/3TC)

Tablet, 12

Tetanus antitoxin Injection, 34 Tetanus antitoxin Injection, 34

Tetanus toxoid vaccine

Injection, 34

Tetracycline HCI Eye Ointment, 37

Theophylline Tablet, 43 Thiamine Injection, 42 Thiamine Injection, 45 Thiopentone sodium

Injection, 1

Thyroxine sodium Tablet, 33 Timolol maleate Eye drops, 38

Tramadol Capsule, 3

Tranexamic acid Tablet, 18 Tranexamic acid Tablet, 18 Tranexamic acid Tablet, 3 Travoprost Eye drops, 38 Triamcinolone acetonide

Injection, 37

Tropicamide Eye drops, 38 Tuberculin purified Injection

solution, 34

Ultravist Injection, 25 Ultravist Injection, 25 Urografin Injection, 25
Urografin Injection, 25
Vancomycin Capsule, 10
Vancomycin Injection, 10
Vecuronium Injection, 1

Vecuronium bromide Injection, 36

Verapami Tablet, 20 Verapamil Tablet, 20 Vinblastine Injection, 16

Vincristine sulphate Injection, 16

Visipaque

+meglumineamidotrizoate

Injection, 26
Vitamin A , 48
Vitamin A Cap, 48
Vitamin A Capsule, 45
Vitamin B Co. strong

Tablet, 45

Vitamin B_{12} Tablet, 45 Vitamin K Injection, 45 Vitamin K Tablet, 45

Vitamins, multiple Injection, 45 Vitamins, multiple Syrup, 45 Vitamins, multiple Tablet, 45

Vitamins, Syrup, 48

Warfarin sodium Tablet, 18 Warfarin sodium Tablet, 18 Water for injections for

i/v use, 44

Whole blood adult Suspension, 19

Whole blood Paediatric

Suspension, 19 X-prep Liquid, 26 Xenetix Injection, 26 Xenetix Injection, 26 Xenetix Injection, 26

Yellow fever vaccine Injection, 35 Yellow soft paraffin Ointment, 24 Zidovudine (AZT) Syrup, 11 Zidovudine (AZT) + Lamivudine

(3TC) Tablet, 12

Zidovudine(AZT) Tablet, 11 Zidovudine(AZT) + Lamivudine

(3TC) Tablet, 12

Zidovudine+Lamivudine (AZT+3TC) Tablet, 13 Zidovudine+Lamivudine+

Tablet, 13 Zinc Tablet, 30

Zinc ointment + Sulphur

Ointment, 22

Zinc oxide (in EO base)

Ointment, 24

Zinc paste compound + Crude coal tar Paste, 23 Zinc paste compound + Sulphur 5% Paste, 22