

Transatlantic Taskforce on Antimicrobial Resistance (TATFAR)

Summary the modified Delphi process for common structure and process indicators for hospital antimicrobial stewardship programs

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Executive Summary

The Transatlantic Task Force on Antimicrobial Resistance (TATFAR) fosters cooperation between the European Union (EU) and the United States (US) on the issue of antimicrobial resistance. The first TATFAR recommendation refers to appropriate use of antimicrobials in human medicine through hospital Antimicrobial Stewardship Programs (ASPs) and, specifically, to the development of common structure and process indicators of ASP. These indicators should allow characterization of programs and comparisons among healthcare systems in EU and US.

To this end, a multidisciplinary expert panel, coordinated by the European Centre for Disease Prevention and Control (ECDC) and US Centers for Disease Control and Prevention (CDC), was formed. The group consisted of 20 experts including representation of nine EU member states and six US states. The expert panel participated in a structured consensus process (modified Delphi method) to facilitate the international collaboration and ensure the equal involvement of all experts. The process was conducted between March and May 2014 and was concluded by a group consensus meeting in June 2014. An initial list of indicators was developed based on previous indicators, available guidance and a review of the literature, including published systematic reviews. The domains assessed were: Governance and Management; Human Resources; Laboratory; Information Technology; Education; Policies for Appropriate Use; Guidelines, Activities and Interventions; and Monitoring of Appropriate Use. The indicators were rated for feasibility, clinical importance and relevance to minimizing antimicrobial resistance. Three rounds of rating followed by the in-person meeting led to a final set of 33 indicators. Among them 17 indicators were considered essential to characterize an ASP and therefore were included in a core set of indicators. The remaining 16 indicators were considered optional indicators and included in a supplemental set.

Implementation of the TATFAR-developed core indicators in multiple nations would contribute to a comprehensive, comparative description of infrastructure, policies, and practices of ASPs internationally. These findings could, in turn, lead to an understanding of best practices of ASPs through further investigation into the relation of different ASP approaches to antimicrobial use and resistance. Current public health surveillance systems or special studies may also be candidates for the addition of ASP questions to baseline surveys. Furthermore these indicators are envisaged as drivers for improvement and alignment of best practices. Piloting, implementation and evaluation of the impact of the indicators constitute important next steps for the optimization of antimicrobial use.

Expert Panel

Names	Institutional Affiliations	Country	
Anastasia Antoniadou	University Hospital Attikon	Athens, Greece	
Bojana Beovic	University Medical Centre	Ljubljana , Slovenia	
Franky Buyle	Ghent University Hospital	Ghent, Belgium	
Sara Cosgrove	Johns Hopkins Medical Institutions	Baltimore (MD), USA	
Peter Davey	Medical Research Institute	Dundee, UK	
Elizabeth S. Dodds Ashley	University of Rochester Medical Center	Rochester (NY), USA	
Catherine Dumartin	Bordeaux University Hospital	Bordeaux, France	
Alison Holmes	Department of Medicine, Imperial College London	London, UK	
Winfried Kern	University of Freiburg Medical Centre	Freiburg, Germany	
Maria Luisa Moro	Regional Agency for Health and Social Care of Emilia-Romagna	Bologna, Italy	
Dilip Nathwani	Department of Medicine, University of Dundee	Dundee, UK	
Jeanne Negley	Georgia Department of Public Health	Atlanta (GA), USA	
Melinda Neuhauser	VHA Pharmacy Benefits Management Services	Hines (IL), USA	
Christopher A. Ohl	Wake Forest University School of Medicine	Winston-Salem (NC),	
		USA	
Diamantis Plachouras	European Centre for Disease Prevention and Control (ECDC)	Stockholm, Sweden	
Lori A. Pollack	Centers for Disease Control and Prevention (CDC)	Atlanta (GA), USA	
Jeroen Schouten	Senior Researcher, Scientific Institute for Quality of Healthcare	Nijmegen, Netherlands	
Ed Septimus	HCA Healthcare System	Houston (TX), USA	
Marc Struelens	European Centre for Disease Prevention and Control (ECDC)	Stockholm, Sweden	
Agnes Wechsler- Fördös	Department of Antibiotic and Infection Control	Wien, Austria	

Coordinators

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Final Set of Core and Supplemental Indicators for Hospital Antimicrobial Stewardship Programs

		CORE INDICATORS for hospital antimicrobial stewardship programs	
Infrastructure	C1	Does your facility have a formal antimicrobial stewardship program accountable for ensuring	
		appropriate antimicrobial use?	
	C2	Does your facility have a formal organizational structure responsible for antimicrobial stewardship	
		(e.g., a multidisciplinary committee focused on appropriate antimicrobial use, pharmacy	
		committee, patient safety committee or other relevant structure)?	
	C3	Is an antimicrobial stewardship team available at your facility (e.g., greater than one staff member	
		supporting clinical decisions to ensure appropriate antimicrobial use)?	
	C4	Is there a physician identified as a leader for antimicrobial stewardship activities at your facility?	
<u>T</u>	C5	Is there a pharmacist responsible for ensuring appropriate antimicrobial use at your facility?	
	C 6	Does your facility provide any salary support for dedicated time for antimicrobial stewardship	
		activities (e.g., percentage of full-time equivalent (FTE) for ensuring appropriate antimicrobial use)?	
	C7	Does your facility have the IT capability to support the needs of the antimicrobial stewardship	
		activities?	
	C8	Does your facility have facility-specific treatment recommendations based on local antimicrobial	
e		susceptibility to assist with antimicrobial selection for common clinical conditions?	
ctic	C9	Does your facility have a written policy that requires prescribers to document an indication in the	
Pra		medical record or during order entry for all antimicrobial prescriptions?	
pu	C10	Is it routine practice for specified antimicrobial agents to be approved by a physician or pharmacist	
c v		in your facility (e.g., pre-authorization)?	
Policy and Practice	C11	Is there a formal procedure for a physician, pharmacist, or other staff member to review the	
		appropriateness of an antimicrobial at or after 48 hours from the initial order (post-prescription	
		review)?	
	C12	Has your facility produced a cumulative antimicrobial susceptibility report in the past year?	
ack	C13	Does your facility monitor if the indication is captured in the medical record for all antimicrobial	
Feedback		prescriptions?	
Monitoring and Fee	C14	Does your facility audit or review surgical antimicrobial prophylaxis choice and duration?	
	C15	Are results of antimicrobial audits or reviews communicated directly with prescribers?	
	C16	Does your facility monitor antimicrobial use by grams [Defined Daily Dose (DDD)] or counts [Days	
		of Therapy (DOT)] of antimicrobial(s) by patients per days?	
	C17	Has an annual report focused on antimicrobial stewardship (summary antimicrobial use and/or	
	ore Ind	practices improvement initiatives) been produced for your facility in the past year?	

C = Core Indicator

		SUPPLEMENTAL INDICATORS
		for hospital antimicrobial stewardship programs
Infrastructure	NA	S1. Does your facility have a named senior executive officer with accountability for antimicrobial leadership?
	C3	Is an antimicrobial stewardship team available at your facility (e.g., greater than one staff
		member supporting clinical decisions to ensure appropriate antimicrobial use)?
		S2. If YES, Is an infection preventionist or hospital epidemiologist involved in stewardship activities?
		S3. If YES, Is a microbiologist (laboratory staff) involved in stewardship activities?
		S4. Is clinical infectious disease (ID) consultation available at your facility?
	C4	 Is there a physician identified as a leader for antimicrobial stewardship activities at your facility? S5. If YES, are stewardship duties included in the job description and/or annual review? S6. If YES, has this physician had specialized training in infectious diseases, clinical microbiology and/or antimicrobial stewardship?
	C5	Is there a pharmacist responsible for ensuring antimicrobial use at your facility?
		S7. If YES, has this pharmacist had specialized training in infectious disease management or stewardship?
	C9	Does your facility have facility-specific treatment recommendations based on local antimicrobial
		susceptibility to assist with antimicrobial selection for common clinical conditions:
Policy and Practice		S8 . If YES, for surgical prophylaxis ?
		S9. If YES, for community-acquired pneumonia ?
		S10. If YES, for urinary tract infection ?
		S11. If YES to any of the clinical conditions above, are these treatment recommendations easily accessible to prescribers on all wards (printed 'pocket guide' or electronic summaries at workstations)?
	C11, C12	Are any of the following actions implemented in your facility to improve antimicrobial prescribing?
		S12. Standardized criteria for changing from intravenous to oral antimicrobial therapy in appropriate situations?
		S13. Dose optimization (pharmacokinetics/pharmacodynamics) to optimize the treatment of organisms with reduced susceptibility?
		S14 . Discontinuation of specified antimicrobial prescriptions after a pre-defined duration ?
ing	NA	S15 . Does your facility measure the percentage of antimicrobial prescriptions that are consistent with the local treatment recommendations for either UTI or CAP?
tor	C15	Does your facility audit or review surgical antimicrobial prophylaxis choice and duration?
Monitoring		S16. If YES, are antimicrobial prescriptions for surgical prophylaxis compliant with facility- specific guidelines in >80% of sampled cases in your facility?

C = Core Indicator

S = Supplemental Indicator

NA = Not applicable to a specific Core Indicator