Antimicrobial Stewardship:

Infection Prevention and Control Project Management

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Disclosures

Today's speaker has no financial relationships with an ineligible company relevant to this presentation to disclose.

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Objectives

- Define infection prevention (IP) and antimicrobial stewardship (AMS)
- Describe the intersection between IP and AMS
- Discuss real world examples of AMS in the IP setting
- Learn to use project management to accelerate IP/AMS projects and utilize as a guide to project completion

IP and AMS

 Infection prevention and control: A practical, evidence-based approach preventing patients and health workers from being harmed by avoidable infections¹

• Antibiotic stewardship: The effort to measure and improve how antibiotics are prescribed by clinicians and used by patients²

Infection Preventionist (IP) Roles

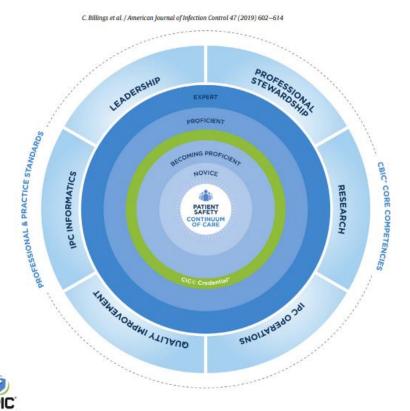


Table 2
Updated APIC Competency Model (2019) future-oriented competency domains and subdomains

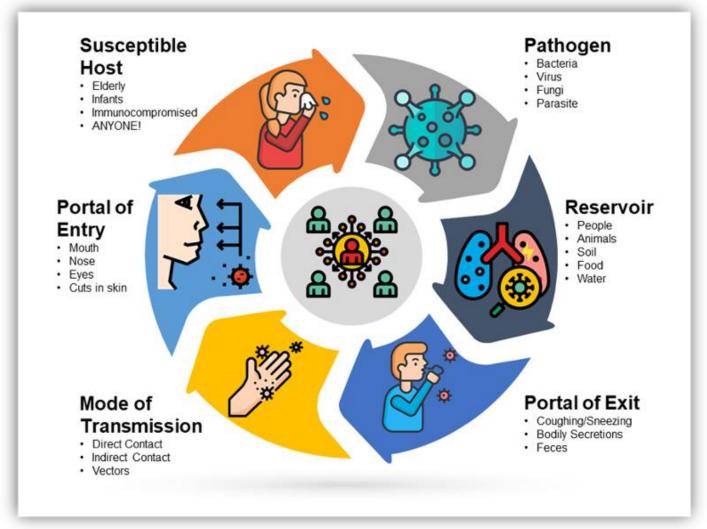
Leadership	Professional stewardship	Quality improvement	IPC operations	IPC informatics	Research
Communication Critical thinking* Collaboration Behavioral science Program management* Mentorship	Accountability Ethics Financial acumen Population health Continuum of care Advocacy	Infection preventionist as subject matter expert Performance improvement* Patient safety Data utilization Risk assessment and risk reduction	Epidemiology and surveillance* Education* IPC rounding Cleaning, disinfection, sterilization Outbreak detection and management Emerging technologies Antimicrobial stewardship* Diagnostic stewardship	Surveillance technology* Electronic medical records and electronic data ware- house* Data management, analysis, and visualization Application of diagnostic testing data and techniques	Evaluation of research Comparative effectiveness research Implementation and dissemina- tion science Conduct or participate in research or evidence-based practice

IPC, infection prevention and control.

*Future-oriented and updated definition of a subdomain that was also in the 2012 competency model.



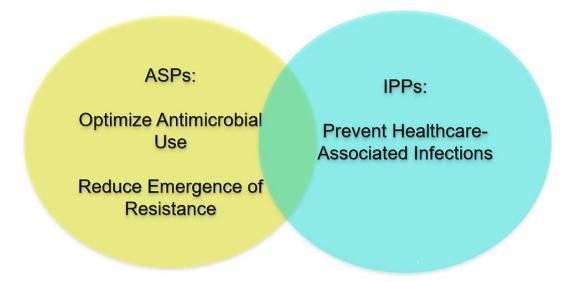
Breaking the Chain of Infection





Antimicrobial Stewardship (AMS)

"Furthermore, AS programs, when implemented alongside IPC measures, especially hand-hygiene interventions, were more effective than implementation of AS alone—verifying that a well-functioning IPC program is fundamental to a successful organizational AS strategy."



"Similar data have also shown that the addition of AS interventions can enhance results of robust IPC measures, particularly when addressing an outbreak." APIC/SHEA/SIDP Antimicrobial Stewardship Position Paper

¹Manning, M. L., Septimus, E. J., Ashley, E. S., Cosgrove, S. E., Fakih, M. G., Schweon, S. J., Myers, F. E., & Moody, J. A. (2018). Antimicrobial stewardship and infection prevention—leveraging the Synergy: A position Paper update. *American Journal of Infection Control*, 46(4), 364–368. https://doi.org/10.1016/j.ajic.2018.01.001

Examples of AMS and IP Synergy

Day-to-Day/Regular Activities

- Daily isolation reviews with standardized language (.dotphrase)
 - MDROs
 - Changing isolation periods to be more similar to prevent confusion and unnecessary test out
 - Unit staff may not understand nuances and multiple isolation periods/test out protocols lead to issues with meeting isolation protocols
 - Removal of isolation for certain with carve outs if a unique population
- Monthly reviews of HAI data with key stakeholders
 - IP and ID providers, pharmacists, and others
 - Allow identification of one offs or concerning trends, explanations for unusual testing



UWMC Transmission-Based Precautions: Precautions A-Z

Type and Duration of Precautions Needed for Selected Infections and Condition

To find the organism or condition of interest in this document, use the Find Function: Ctrl-F or search subject by letter Subject by letter: A B C D E F G H I L M N O P R S T V X Y Z

Precaution Type/Duration	Abbreviation	Comment
Standard	S	Use personal protective equipment (mask, eye protection, gowns and gloves) whenever contact with secretions or excretions is anticipated during care. Transmission-based Precautions are used in addition to Standard Precautions.
Contact	С	Wear gown and gloves.
Contact Plus	СР	Wear gown and gloves for all staff, providers and visitors. Use dedicated equipment and disposable meal trays. EVS follows Contact Plus cleaning protocol. Staff clean high touch surfaces at least twice daily.
Contact Enteric	CE	In addition to Contact Precautions, wash hands with soap and water when exiting room and use hospital approved disinfectant appropriate for contact enteric precautions, such as bleach.
Droplet Contact	DC	Wear surgical mask, eye protection, gown and gloves. Additional requirements during aerosol generating procedures (AGP), see comments for specific infection/condition
Airborne Respirator	AR	Wear PAPR or fitted N95 respirator during care. Place patient in negative pressure Airborne Isolation Infection Room (AIIR).
Airborne Respirator Contact	AR/C	In addition to AR precautions above, Wear gown, gloves, eye protection.
Aerosol	Aerosol	Wear PAPR or fitted N95 respirator during care. Patient is placed in negative pressure Airborne Isolation Infection Room (AIIR) when possible. AIIR preferred for AGP

Precautions: Standard and Expanded Policy

Moderately immunocompromised: Individuals receiving chemotherapy for solid tumors, solid organ transplant recipients, HIV patients with CD4 counts <200, patients with acquired or genetic immunodeficiencies, patients on prolonged or high-level immunosuppression (e.g. cyclophosphamide, MMF) and those receiving prednisone > 20 mg/day for more than 14 days.

Highly Immunocompromised: Individuals who are receiving treatment for a hematologic malignancy (e.g. leukemia, lymphoma, multiple myeloma), all hematopoietic cell transplant, and those receiving CAR-T cell therapy.

NICU Visitation guidelines: uwm-ea-2127.pdf and uwm-ea-1367.pdf

Pregnant Healthcare Workers: Precautions during Pregnancy UWMC Employee Health Document

Precautions apply to all settings unless specifically indicated

MDROs not listed below (Includes multidrug-resistant E. coli, ESBL, Serratia, Enterobacter, Klebsiella, Acinetobacter, Pseudomonas, and all other MDROs that are not listed below) Carbapenemase-producing and other XDROs Carbapenemase-producing And other XDROs Carbapenemase-producing Carbapenemase-producing And other XDROs Sandard: Patients who are colonized (eg. Positive nares screening swab) or who have a history of MRSA and no active infection Contact: Active MRSA infection (e.g., bloodstream, lower respiratory culture (sputum), or wound) Until antibiotic treatment complete Exceptions: Prolonged antibiotic therapy (e.g., endocarditis, osteomyelitis, etc.): At least 2 weeks of therapy + negative blood cultures Standard: Patients with a history of VRE and no active infection Contact: Active VRE infection (e.g., bloodstream, lower respiratory culture (sputum or wound) keep in precautions until effective antibiotic treatment complete and no uncontained draining wounds	Infection/Condition/		Duration	Contact IP to Clear	
Carbapenemase-producing and other XDROs Methicillin Resistant Staphylococcus aureus (MRSA) S or C Vancomycin Resistant Enterococci (VRE) C Indefinitely C CPO/XDRO are considered pathogens of epidemiological concern Standard: Patients who are colonized (eg. Positive nares screening swab) or who have a history of MRSA and no active infection Contact: Active MRSA infection (e.g., bloodstream, lower respiratory culture (sputum), or wound) Until antibiotic treatment complete Exceptions: Prolonged antibiotic therapy (e.g., endocarditis, osteomyelitis, etc.): At least 2 weeks of therapy + negative blood cultures Standard: Patients with a history of VRE and no active infection Contact: Active VRE infection (e.g., bloodstream, lower respiratory culture (sputum or wound) keep in precautions until effective antibiotic treatment complete and no uncontained draining wounds	(Includes multidrug-resistant E. coli, ESBL, Serratia, Enterobacter, Klebsiella, Acinetobacter, Pseudomonas, and all other MDROs	С	after completed treatment for	-	Re-testing not required
Methicillin Resistant Staphylococcus aureus (MRSA) S or C Until treatment complete - Contact: Active MRSA infection (e.g., bloodstream, lower respiratory culture (sputum), or wound) Until antibiotic treatment complete Exceptions: Prolonged antibiotic therapy (e.g., endocarditis, osteomyelitis, etc.): At least 2 weeks of therapy + negative blood cultures Standard: Patients with a history of VRE and no active infection Contact: Active VRE infection (e.g., bloodstream, lower respiratory culture (sputum or wound) keep in precautions until effective antibiotic treatment complete and no uncontained draining wounds	Carbapenemase-producing	С		-	CPO/XDRO are considered pathogens of epidemiological concern
Vancomycin Resistant Enterococci (VRE) S or C Until treatment complete Contact: Active VRE infection (e.g., bloodstream, lower respiratory culture (sputum or wound) keep in precautions until effective antibiotic treatment complete and no uncontained draining wounds	Staphylococcus aureus	S or C	treatment		have a history of MRSA and no active infection Contact: Active MRSA infection (e.g., bloodstream, lower respiratory culture (sputum), or wound) Until antibiotic treatment complete Exceptions: Prolonged antibiotic therapy (e.g., endocarditis, osteomyelitis, etc.): At
do not require precautions	•	S or C	treatment	-	Contact: Active VRE infection (e.g., bloodstream, lower respiratory culture (sputum), or wound) keep in precautions until effective antibiotic treatment complete and no uncontained draining wounds Note: E. gallinarum and E. casseliflavus are intrinsically resistant to vancomycin and

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ISOVZVCONTACT	Localized Varicella Zoster Virus identified in skin rash on ***. Because the rash cannot be fully cov
ISOVZVCSFONLY	Varicella Zoster Virus in CSF only identified on ***. Please maintain standard precautions. If skin le
ISOVZVDISSEMINATED	Patient positive for disseminated Varicella Zoster Virus skin rash on ***. Please maintain airborne
ISOVZVLOCALIZEDIMMU	Localized Varicella Zoster Virus skin rash identified on ***. Because patient is immunocompromise
ISOVZVSTANDARDPREC	Localized Varicella Zoster Virus skin rash identified on ***. Since lesions can be fully covered/cont

Continued

- Two-step *C. diff* testing
- Blood culture protocols
- Shared definitions for hypotension for centralized bloodstream infections
- Shared protocols for tuberculosis

With so many competing priorities how can an IP get it all done?

5 Phases of the Project Life Cycle



Step 1: Initiation

Project goals & feasibility

Stakeholder register

Project charter

Kickoff meeting



Step 2: Planning

Scope & budget

Deadlines

Team roles

Communication plan

Milestones



Step 3: Execution

Task completion

Team collaboration

Efficient workflows

Status reports & meetings



Step 4: Monitoring & control

Budget & timeline

Project goals

Quality control

Team performance

Risk management



Step 5: Closure

Retrospective meeting

Project closure report

Team celebration

=teamgantt

Examples from Previous Work

Shared Hypotension Definition

- 1. Initiation
- 2. Planning
- 3. Execution
- 4. Monitoring & Control
- 5. Closure

Tuberculosis

- 1. Initiation
- 2. Planning
- 3. Execution
- 4. Monitoring & Control
- 5. Closure

Tuberculosis Documentation Example

Tuberculosis (TB) System and Facility Comparison

5/13/25; most recent update 7/7/25

Scope: This document is intended to be used as a tool to compare TB practices across system partners, identify disparate practices, and determine where alignment is possible. In areas where alignment is not possible, the request is to have agreement on which facility policy should be followed and when. Note: Policies and content reviewed are located within the appendix at the end of document.

<u>Directions</u>: Please answer all <u>bolded</u> questions for your facility in the <u>ALIGNMENT QUESTIONS/DISCUSSION</u> column.

- Do not answer those that are greyed out. These will be answered as a group and/or informed by previous questions. Will be updated later.
- · Less than 20 minutes to complete, 15 questions total.
- · Please have physician lead and IP lead review and respond
- Due: 5/21/25

Yellow- Request more information; potential disparate practice

If all responses are <u>not in alignment</u>, proposed and alignment language will be below responses or in comment box

Detailed Algorithm and Clearance Criteria Comparison

TB Categories Found in A-Z Disease Surveillance Documents and Others				
нмс	UWMC	FRED HUTCH (FH)	POTENTIAL DISPARATE PRACTICES	ALIGNMENT QUESTIONS/DISCUSSION
Tuberculosis (M. tuberculosis) Extrapulmonary, (draining lesion) Extrapulmonary, no draining lesion, meningitis Pulmonary or laryngeal disease,	Mycobacterium tuberculosis Extra-pulmonary TB TB Meningitis included Latent TB Mycobacteria, nontuberculosis (atypical)	Tuberculosis (M. tuberculosis, MTB, TB) Extrapulmonary No draining lesion, including meningitis Draining lesion Latent (LTBI),	Coordinate outpatient evaluation and notify public health Notify Public Health-Seattle & King County PHSKC-TB Clinic if a King County resident. Not a resident of King County? Notify the Public Hearth Department of that jurisdiction. Place mask on patient and initiate Airborne Respirator Isolation - Negative Pressure room required - Respirators for all HCW - If outpatient, surgical mask when entering/Exiting the clinic room NO Does patient require hospitalization?	(1) Do TB policies only apply to inpatient care? HMC and UWMC differentiate per decision tree example on left. No noted clearance criteria for outpatient specificstates to coordinate Qutpt and notify PH. Hope to clarify if Qutpt follows the same policies and clearance as Inpt. See Appendix for more. a. HMC: N b. UWMC: N, applies to outpatient as well. Also depends on how public health advises.
 confirmed Pulmonary or laryngeal disease, suspected Skin-test positive 	Pulmonary and Laryngeal TB	including blood or skin test positive without TB signs or symptoms • Pulmonary or	(2) Latent TB (ALL) a. Skin test positive without TB s/s (HMC, FH only with specific call out)	c. FH: NA, Specific outpatient policies (2) Does a positive skin test w/o TB s/s fall under Latent TB?
with no evidence of current active disease Neisseria meningitidis (meningococcal) known or suspected • M. tuberculosis		Laryngeal O R/O or suspected TB O Active TB, including miliary	(3) Active TB, including miliary (FH only)	a. HMC: Y b. UWMC: Y c. FH: Y-include both IGRA and positive skin test (TST/PPD) (3) Would facilities consider specific call out for miliary TB under pulmonary and laryngeal TB category?

Mycobacterium,	Mycobacteria,	a. HMC: Y
nontuberculosis	nontuberculosis (atypical;	b. UWMC: Y
(atypical)	NTM)	c. FH: Y
 Pulmonary 		
 Wound 		

QUESTIONS?