

2023 KNOWLEDGE-BASED HOME-STUDY CE FOR PHARMACISTS AND PHARMACY TECHNICIANS

Perioperative Antimicrobial Stewardship and Pharmacy's Role

Speaker: Sara Jordan Hyland, PharmD, BCCCP, Clinical Pharmacist, Grant Medical Center, Columbus, OH

Learning Objectives

Upon completion of this CE activity, the Pharmacist will be able to do the following:

- Explain key pharmacokinetic and mechanistic concepts of antibiotics as they pertain to surgical antimicrobial prophylaxis
- Select optimal surgical antimicrobial prophylaxis regimens based on patient- and surgery-specific factors
- Identify antibiotic applications unique to surgical patients, including irrigations, antibiotic-laden bone cement (ALBC), spacers, pastes, and powders
- Recognize best practices and clinical pharmacist roles in perioperative antimicrobial stewardshipIdentify patient awareness opportunities related to safe pharmaceutical waste disposal

Upon completion of this CE activity, the Pharmacy Technician will be able to do the following:

- Describe the role of prophylactic antibiotics in surgical patients
- Locate clinical practice guidelines pertaining to antibiotic use in surgical patients
- Identify antibiotic applications unique to surgical patients, including irrigations, antibiotic-laden bone cement (ALBC), spacers, pastes, and powders
- Recognize medication stocking needs and medication safety implications pertaining to antibiotic use in surgery

Accreditation Information:

ACPE UAN: 0122-0000-23-039-H01-P/T

CPE Credit Hours: 1.0 hour (0.1 CEU) of home-study CE credit

Release Date: June 5. 2023

Expiration Date: June 5, 2026

Disclosures: Dr. Jordan Hyland, the moderators and planners for this activity do not have relevant financial relationships to disclose with ineligible companies.

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- 2. View and listen to the recorded activity
- 3. Complete both the learning assessment questions and the activity evaluation.

<u>Note</u>: A passing grade of 65% or better is required to receive CE credit. Three opportunities will be provided to successfully complete the post-test. Once all steps above are completed, participation will be provided to CPE Monitor.



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PERIOPERATIVE ANTIMICROBIAL STEWARDSHIP AND PHARMACY'S ROLE

Speaker:

Sara Jordan Hyland, PharmD, BCCCP

Clinical Pharmacist Grant Medical Center Columbus, OH

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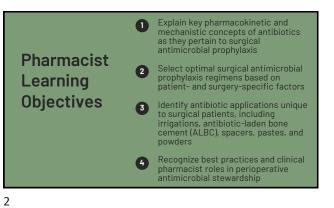
Speaker Bio:

Sara J Hyland, PharmD, BCCCP is a clinical pharmacist for Perioperative Services and Emergency Medicine at Grant Medical Center (OhioHealth), a large community teaching hospital and Level 1 Trauma Center in Columbus, OH. She holds Bachelor of Science in Pharmaceutical Sciences and PharmD degrees from The Ohio State University College of Pharmacy. Sara completed a PGY1 Pharmacy Practice Residency at Grant Medical Center. She is a Board-Certified Critical Care Pharmacist. She precepts PGY1 and PGY2 pharmacy residents in research rotations and in clinical perioperative/orthopedics rotations. Her practice and research interests include analgesia/sedation, neuromuscular blockade/reversal, hemostasis, VTE prophylaxis, pharmacy practice, and implementation science related to emergency and perioperative medicine. She has published or presented numerous professional papers on a variety of topics. Dr. Hyland serves on the research committee for the ACCP Emergency Medicine Practice and Research Network.

Perioperative Antimicrobial Stewardship and Pharmacy's Role

Sara J. Hyland, PharmD, BCCCP

Clinical Pharmacist - Perioperative Care and Emergency Medicine OhioHealth Grant Medical Center | Columbus OH | April 2023 sara.jordan@ohiohealth.com | @SaraJPharmD



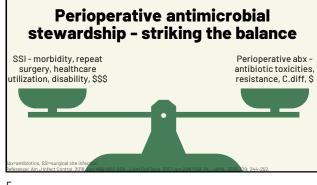
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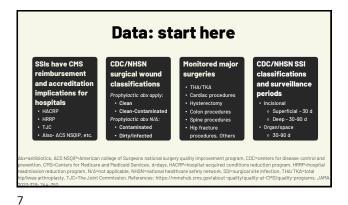
Disclosures and Disclaimers

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 Much of perioperative antibiotic use is not driven by high-quality evidence
- Comprehensive perioperative antimicrobial stewardship programs
- (Periop ASPs) appear to be in their infancy in published literature
- and consensus guidelines appear nonexistent at this time
- I'll be drawing from both published evidence and personal experience (>10 years practicing in this space)
 → "Tips" and "Best Practices" in this presentation are my personal recommendations for Periop ASPs



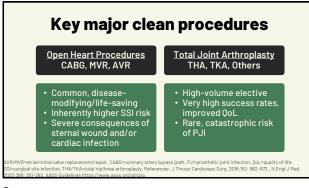






Recommended Best Practice: Pharmacists should collaboratively review every SSI case for antibiotic- and other medicationrelated opportunities

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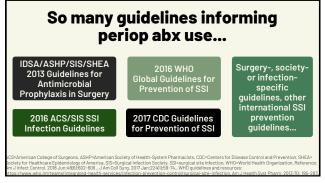


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Why is PJI such a big deal? Hopefully Success! TJA ~99% cured! May still aet lonaterm abx DAIR Lots of abx PJI Failure: Long-term abx, 1-Stage revision ~1% 2-Stage revision: reimplant 2-Stage revision: radical debridement Lots of eventually abx amputation /Girdlestone. More worse? abx 10

<u>Polling Question</u>: Which resource(s) are you relying on most heavily to guide your periop antibiotic recommendations at this time? Pick any that apply-

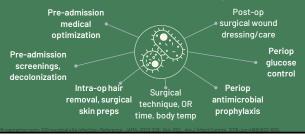
- A: 2013 IDSA/ASHP/SIS/SHEA Guidelines for Antimicrobial Prophylaxis in Surgery
- B: 2016-2017 Guidelines for SSI Prevention from WHO, CDC
- C: Surgery- or Society-specific guidelines
- D: Primary research/studies since guidelines are limited
- E: Many/multiple resources (I might be overwhelmed!)



<u>Tip</u>:

Don't just use these (what WE may like/care most about) - Make sure you know what the surgeons are reading/referencing (what THEY care most about) and integrate!

SSI mitigation is multimodal, interprofessional, and longitudinal



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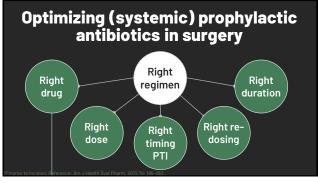
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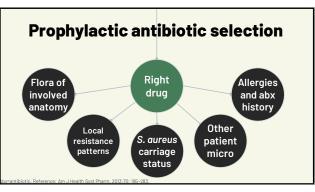
<u>Tip</u>:

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Today we'll focus on perioperative antibiotic use and stewardship, but pharmacists can help reduce SSIs in many ways! Polling Question: A 62yom weighing 102 kg with PMH of OA, DM2, and obesity presents for elective primary TKA. Pre-admission *S. aureus* nasal culture +MSSA. Allergy to "penicillins" (rash). Which pre-op antibiotic regimen is he currently most likely to receive at your institution?

A: Cefazolin 1000 mg
B: Cefazolin 2000 mg
C: Clindamycin 900 mg
D: Vancomycin 1000 mg
E: Vancomycin 2000 mg





<u>Tip</u>:

Optimizing preop antibiotic prescribing is challenging, but high-yield for improving SSI rates (and for making friends!)

Right drug: normal flora and other SSI pathogens by procedure

Skin flora - stop here for clean procedures (ortho, neuro, podiatry, CTS, breast]: S. aureus, coagulasenegative staphylococci, streptococci

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- <u>Head/neck</u>: polymicrobial, peptostreptococcus <u>Shoulder arthroplasty</u>: C. acnes
- <u>Gastroduodenal, hepatobiliary</u>: enteric GNB, less of-*Bacteroides* spp., enterococci
- <u>CRS, SOT</u>: as for upper GI, anaerobes more frequent
- <u>Hysterectomy</u>: polymicrobial including GPCs, enteric GNB, Bacteroides spp. <u>Uro/gyn (if in urinary tract)</u>: enteric GNB, if penile prosthesis - plus pseudomonas, candida

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Common pitfalls in pre-op abx ordering

• When cefazolin monotherapy isn't sufficient for the procedure

- When cefazolin monotherapy is sufficient, but clindamycin and vancomycin are not
- Minimally invasive procedures prophylactic abx may not be indicated

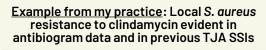
Right drug: consider local resistance patterns

Local antibiograms

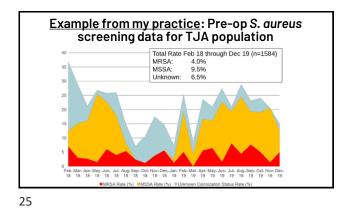
Local SSI pathogen history from case reviews

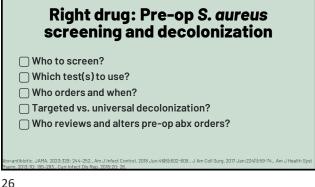
Local S. aureus colonization data

Example from my practice:		MSSA	MRSA
Local S. aureus resistance to	Cefazolin	100%	0%
clindamycin evident in antibiogram data and	Clindamycin	74%	58%
in previous TJA SSIs	Vancomycin	100%	100%



<i>S. aureus</i> Colonization	Pre-op Antibiotic	SSI Pathogen	SSI Pathogen Susceptibility to Clindamycin
Unknown	Clindamycin	MRSA	Resistant
Unknown	Cefazolin	MSSA	Resistant
Unknown	Cefazolin + gentamicin	MSSA	Susceptible
Unknown	Clindamycin + gentamicin	MRSA	Resistant
Unknown	Cefazolin	MRSA	Susceptible





Pre-op S. <i>aureus</i> screening, decolonization, targeted prophylaxis						
OR (95%CI) for GPC SSI for MRSA SSI for MSSA SSI						
Decolonization n=1730	0.41 (0.30-0.55)	0.30 (0.15-0.62)	0.50 (0.37-0.69)			
Glycopeptide prophylaxis 0.70 (0.40 0.40 (0.20-0.80) 1.47 (0.91-2.38)						
Decolonization + targeted ppx n=1129	0.41 (0.30-0.56)	0.22 (0.12-0.38)	0.45 (0.26-0.78)			

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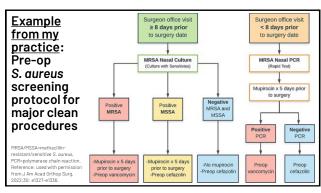
Right drug: Pre-op S. aureus screening and decolonization

Who to screen? - high-risk procedures, esp. TJA, CTS
 Which test(s) to use? - consider culture if have time vs. PCR
 Who orders and when? - surgeon office, PAT?
 Targeted vs. universal decolonization? - targeted please!
 Who reviews and alters pre-op abx orders?? - pharmacists!

ntibiotic. JAMA. 2023;329: 244–252., Am J Infect Control. 2018 Jun;46(6):602-609., J Am Coll Surg. 2017 Jan;224(1):59-74., Am J He

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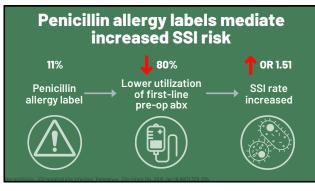
Recommended Best Practice: Pharmacists should prospectively review and optimize all preoperative antibiotic orders



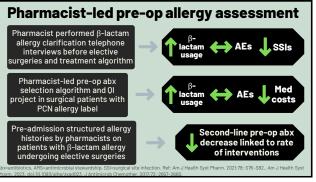


Right drug: Allergy assessment
Are allergies effectively evaluated at pre-op abx ordering?
Best penicillin delabeling strategy for periop?
Improved history-taking: who, when, how?
Pre-op allergist referral?
Augmenting/suppressing allergy warnings?
Again, who reviews and alters pre-op abx orders??
xx=antibiotic. Reference: Infect Control Hosp Epidemiol. 2022;43: 829–833. J Am Acad Orthop Surg. 2023;31: e107–e117. J Arthroplasty. 2017. 01–S108., Clin Orthop Relat Res. 2016;474: 1801–1808., Clin Infect Dis. 2018;66: 329–336., Clin Infect Dis. 2021;72: 1404–1412., Hosp Epidemiol. 27.4. 529. 817.

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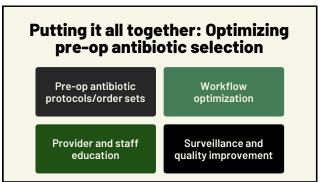


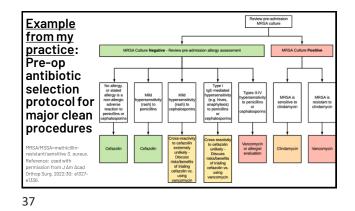


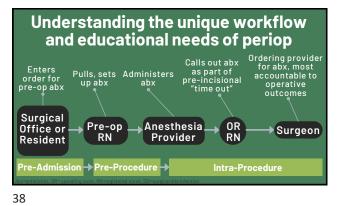


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Recommended Best Practice: Perioperative antimicrobial stewardship efforts should make increasing use of first-line pre-op antibiotics to improve patient and institutional outcomes a top priority





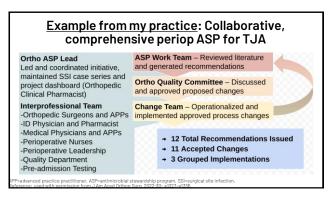


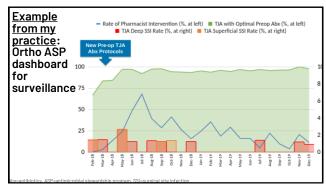
Understanding the unique workflow and educational needs of periop "Patient is PCN-allergic, are we sure we want cefazolin PTI?" and dies from anghylavis?" May auto- "Dr. B, you "You want matically ordered me to give change to cefazolin when what? What second-line the pt has a about my about my liability?!" the pt has a PCN allergy!" and dies from anaphylaxis?!' abx Surgical Pre-op Anesthesia. V OR Office or Surgeon RN Provider RN Resident Pre-Admission → Pre-Procedure → Intra-Procedure

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Recommended Best Practice: Periop antimicrobial stewardship programs should only be implemented after collaborative development and broad educational interventions spanning the surgical continuum of providers and staff

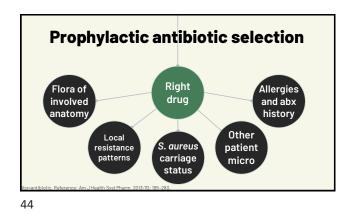


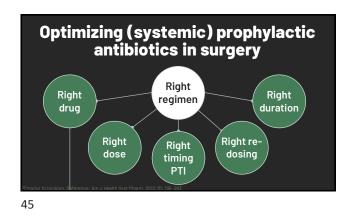


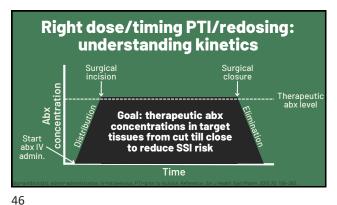


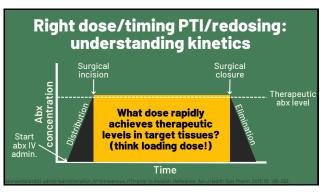


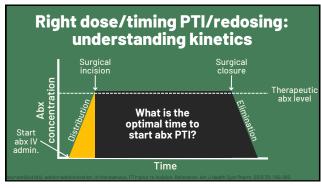
<u>Example</u> from my	• AKI F		uperficial	SSI Rate	• Deep S	
			Start		Start	Phase 3 Start
practice:	8%					
Quality	6%		•		•	•
improvemen	L 4%					
	490					
report for				-		
report for Ortho ASP	2%			-		
				+		
Ortho ASP		-A FY1	18-B F	Y19-A F	Y19-B	FY20-A
Crtho ASP	2%	-A FY1 7/17 - 12/17	18-B F	Y19-A F 7/18 - 12/18	TY19-B	
Critho ASP	2% FY18					FY20-A 7/19 - 12/19 1.33%
Crtho ASP	2% FY18 Outcome	7/17 - 12/17	1/18 - 6/18	7/18 - 12/18	1/19 - 6/19	7/19 - 12/19

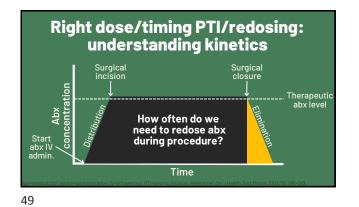




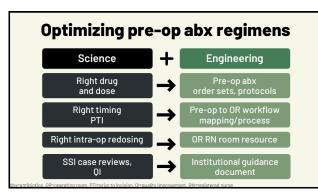








- no co	Cefazolin vs. alternatives - no contest for ideal pre-op antibiotic!					
	Cefazolin	Clindamycin/Vancomycin				
	Ø	\checkmark				
	$\mathbf{\nabla}$	×				
	V					





Example from my practice: Periop workflow /process mapping for pre-op vancomycin				
Pre-op holding area	Operating room			
Pre-op RN identifies vancomycin as ordered abx, notifies OR circulator RN	Circulator RN notes patient needs pre-op vancomycin start time coordinated			
*	*			
Pre-op RN pulls vancomycin from ADC and sets up on smart pump, leaving paused	Circulator RN works with scrub and anesthesia teams to identify when patient is ~60 mins PTI and notifies pre-op RN			
Pre-op RN starts vancomycin and documents administration				
Patient transferred to OR with vanco running, pre-op to circulator h/o	Anesthesia team receives patient, confirms info, continues vanco and documents infusion stop			

Example from		Start intra-op dose at this time after start of pre-op dose
my practice:	Cefazolin	4 hours*
Intra-op redosing OR resource -	Clindamycin	6 hours*
laminated copy	Vancomycin	None needed
on every OR white board, used by OR	Cefotetan	6 hours*
circulator RN	Gentamicin	None needed
EBL=estimated blood loss,	Metronidazole	None needed
OR=operating room, RN=registered nurse. Reference: Am J Health Syst Pharm. 2013;70: 195-283.,	*also give	redose if/when 1500 mL EBL

<u>Example from</u> <u>my practice</u> : Guidance	Pre-op Abx	Recommended Start Time PTI	Recommended Redosing Interval
document for	Cefazolin	5-60 min	4 hr
periop abx timing for QI	Clindamycin	15-120 min	6 hr
purposes	Vancomycin	15-60 min	Not indicated
Abx=antibiotic(s). IVP=intravenous push, PTI=prior to incision, DI=quality improvement. Reference:	Cefotetan	15-60 min	6 hr
01–quanty improvement, Receiptice: Am J Health Syst Pharm. 2013;70: 195–283., Ann Surg. 2009;250: 10– 16., Medicine (Baltimore). 2017;96: e6903., Antimicrob Agents	Gentamicin	15-120 min	Not applicable
Chemother. 2002;46: 3026-3030., JAMA. 1980;244: 1353-1354., lin Orthop Relat Res. 2011;469: 3486-	Metronidazole	30-120 min	Not indicated
3494., J Antimicrob Chemother. 2006;58: 645-650.			

Tip:

When developing pre-op antibiotic protocols and procedures, consideration must be given to:

- infusion times
- necessity of smart pumps
- compatibility of abx and infusion apparatus with anesthetics and anesthesia setup
- other logistical/human factors

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Polling Question: Clinical case - A 62yom with NKDA was admitted to your hospital last night with septic shock. He was started on empiric vancomycin 1750 mg q12hr, cefepime 2 g IV q12hr, IV fluids and vasopressors. This morning, he is being rushed to OR for emergent ex-lap with possible subtotal colectomy for suspected ischemic bowel. The OR RN calls you to ask what pre-op abx they should give PTI, considering the vancomycin was last given 4 hours ago and cefepime 2 hours ago. What would you advise?

- A: No additional abx are needed PTI

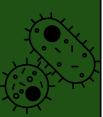
- B: Get an order for cefazolin 2 g IV to give PTI C: Get an order for cefotetan 2 g IV to give PTI D: Get an order for metronidazole 500 mg IV to give PTI

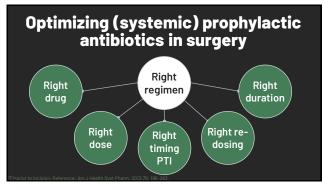
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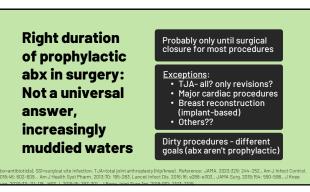
Special circumstances - inpatients already on abx who need surgery

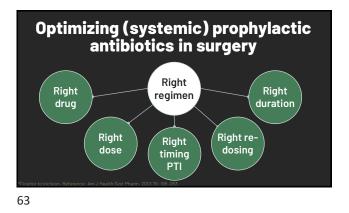
- Goal = ensure abx regimen maintains therapeutic need while also providing sufficient spectrum AND timing for duration of surgical prophylaxis
- Requires patient-specific evaluation, optimization, and communication
 - · Consider length of procedure and also if current abx at steady-state yet
 - Ensuring floor doses are given intra-op requires "manual override" of usual processes
- If therapeutic goal is met, no clinical or compliance reason to give additional abx PTI (if documented infection)!

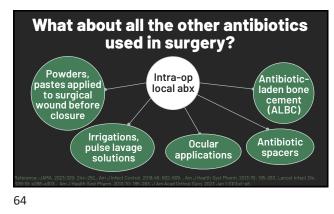
Recommended Best Practice: Clinical pharmacists should target inpatient surgeries for patient-specific antibiotic optimization interventions

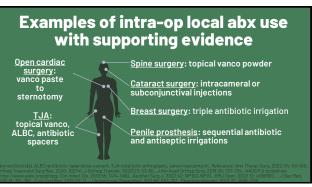


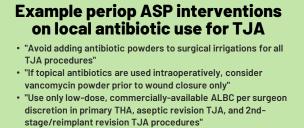


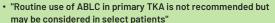












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Example periop ASP interventions on local antibiotic use for TJA

 "Adding antibiotic powder to bone cement to compound highdose ABLC in 1st-stage/spacer revision TJA procedures is appropriate, but consider using vancomycin preferentially for this purpose due to best likely benefit/risk ratio of studied antibiotics. Additionally, postoperative therapeutic drug level monitoring should be considered before further systemic antibiotic dosing to avoid overexposure"

ALBC=antibiotic-laden bone cement, THA/TJA/TKA=total hip/ioint/knee arthroplasty. Reference: J Am Acad Orthop Suri

<u> Tip</u>:

Systemic exposure and toxicity can occur from local intra-op antibiotics, especially those loaded into spacers or bone cement during major orthopedic procedures! <u>Polling Question</u>: Does your institution have vials of antibiotic powder(s) stocked in your OR automated dispensing cabinets (ADCs)?

ycin

A: I have no idea
B: I think so?
C: Yes, vancomycin
D: Yes, vancomycin and tobram
E: Yes, tons of them!

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Intra-op local/topical antibiotic use: Opportunities unidentified

- Not "ordered" no clear order trail, no pharmacist verification
- Surgeon preference cards
- Override ADC to sterile field
- Admixed in OR?
- Allergy warnings?
- "One-step" documentation instead of MAR entries

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Recommended Best Practice: Pharmacy technicians and pharmacists should routinely collaborate to surveillance antibiotic stock in perioperative ADCs to identify potential opportunities to optimize medication safety and stewardship

Closing thoughts why is perioperative antimicrobial stewardship so hard?

- Many targets spanning phases of care
- Complex orchestra of players and processes
- Collaboration is essential, but can be <u>challenging</u>
- Need patient outcomes data, not just med use
- Need to pay for it
- Incremental wins

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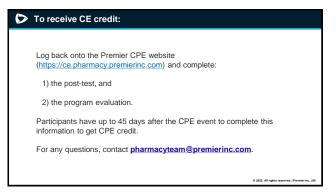
Summary and Discussion

- Perioperative antimicrobial stewardship has unique challenges
 - Collaboration and integration of clincial + operational aspects are keys to success
- Pharmacists and pharmacy technicians can play active roles in improving perioperative antibiotic use to improve patient and institutional outcomes

Perioperative Antimicrobial Stewardship and Pharmacy's Role

Sara J. Hyland, PharmD, BCCCP

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Торіс	CDC Guideline for Prevention of SSI, 2017	ACS/SIS: Surgical Site Infection Guidelines, 2016 Update	WHO Global Guidelines on SSI Prevention, WHO Recommendations on Intra/Postop SSI Prevention- 2016	IDSA/SIS/ASHP/SHEA Guidelines for AMP in Surgery, 2013
Pre-operative staph aureus screening and decolonization; use of pre-op vancomycin	Did not directly address	 Implementation of MRSA screening and decolonization should depend on baseline SSI and MRSA rates Screening and decolonization is recommended for patients undergoing total joint replacement and cardiac procedures No standard decolonization protocol supported by literature Vancomycin should not be administered as prophylaxis to MRSA-negative patients 	 Patients undergoing cardiothoracic or orthopedic surgery with known nasal carriage of <i>S. aureus</i> should receive perioperative intranasal applications of mupirocin 2% ointment with or without chlorhexidine body wash (strong recommendation, moderate quality of evidence) No recommendations on frequency or duration of mupirocin administration The panel suggests <i>considering</i> treating patients with known nasal carriage of <i>S. aureus</i> undergoing other types of surgery with intranasal mupirocin 2% with or without chlorhexidine body wash (conditional recommendation, moderate quality of evidence) Take into account local rates of <i>S. aureus</i> and MRSA, and patient-specific factors (past <i>S. aureus</i> infection, known carrier of CA-MRSA, or colonized by <i>S. aureus</i> in sites other than the nose **Screening for S. aureus was not assessed as part of the intervention so no recommendations can be made 	Vancomycin prophylaxis should be considered in patients with known MRSA colonization or at high risk for MRSA colonization in the absence of surveillance data. In institutions with SSIs attributed to CA-MRSA, other antimicrobial agents with activity against MRSA should be considered. Screening for both MSSA and MRSA colonization is recommended to help identify candidates for decolonization and to assist in the selection of optimal antibiotic therapy. Decolonization with intranasal application of 2% mupirocin is recommended in many patients colonized with <i>S. aureus</i> , especially those undergoing cardiac and orthopedic surgeries.

Торіс	CDC Guideline for Prevention of SSI, 2017	ACS/SIS: Surgical Site Infection Guidelines, 2016 Update	No recommendations about which antibiotics to use or dosing were made WHO Global Guidelines on SSI Prevention, WHO Recommendations on Intra/Postop SSI Prevention- 2016	IDSA/SIS/ASHP/SHEA Guidelines for AMP in Surgery, 2013
Duration of antimicrobial prophylaxis (AMP)	In <u>clean and</u> <u>clean-contaminated</u> <u>procedures</u> , do not administer additional prophylactic antimicrobial agent doses after the surgical incision is closed in the operating room, even in the presence of a drain. (Category IA-strong recommendation; high-quality evidence.) The available data examined the following comparisons for different postoperative AMP durations: 1. All surgeries—No post-op AMP vs. ≤ 24 hours 2. Cardiac a. No post-op AMP vs. ≤ 24 hours b. No post-op AMP vs. < 96 hours c. No post-op AMP vs. 72-96 hours d. ≤ 24 vs. 72 hours	Antibiotics should be discontinued at time of incision closure, with the exception of; Implant-based breast reconstruction 0 24-48hrs Joint arthroplasty 0 24hrs Cardiac procedures 0 48hrs In general, there is no evidence that antibiotic administration after incision closure decreases SSI risk across a range of procedures, including <u>clean</u> , <u>clean-contaminated, and</u> <u>contaminated</u> wound classes.	The panel recommends against the prolongation of surgical antibiotic prophylaxis (SAP) administration after completion of the operation for the purpose of preventing SSIs (strong recommendation, moderate quality of evidence) Meta-analysis showed continuation might be beneficial in • Cardiac (OR= 0.43; 0.25 – 0.76) • Orthognathic (OR= 0.3; 0.1 – 0.88) **Considering the low quality of the evidence and the results of the overall meta-analysis (moderate quality), the expert panel decided to strongly recommend against SAP prolongation, also because of the widespread risk of antimicrobial resistance. The panel suggests not continuing perioperative antibiotic prophylaxis because of the presence of a wound drain (conditional recommendation, low quality of evidence). Seven RCTs were identified. The meta-analysis showed that prolonged antibiotic prophylaxis in the presence of a wound	Post-operative antimicrobial administration is not recommended for most cases, and the duration of antimicrobial prophylaxis should be less than 24 hours for most procedures. A prophylaxis duration of up to 48 hours has been accepted for cardiothoracic procedures (expert panel consensus) Several studies have shown that extended use of antimicrobial prophylaxis (>48 hours) fails to reduce the risk of SSI and leads to antimicrobial resistance to selected antibiotics

Торіс	3. Thoracic—No post-op AMP vs. 2 days CDC Guideline for Prevention of SSI, 2017	ACS/SIS: Surgical Site Infection Guidelines, 2016 Update	drain has no benefit in reducing SSI compared with perioperative prophylaxis alone (OR 0.79; 95% CI 0.53–1.20). WHO Global Guidelines on SSI Prevention, WHO Recommendations on Intra/Postop SSI Prevention- 2016	IDSA/SIS/ASHP/SHEA Guidelines for AMP in Surgery, 2013
Intra-operative topical/local antibiotic use	No recommendation regarding antimicrobial irrigation for the prevention of SSI was made. Topical antimicrobial agents should not be applied to the surgical incision for the prevention of SSI (Category 1B – strong recommendation, low quality evidence)	Topical antibiotics can reduce SSI for specific cases, including spine surgery, total joint arthroplasty, and cataract surgery, but there is insufficient evidence to recommend routine use at this time.	The panel considers that there is insufficient evidence to recommend for or against saline irrigation of <u>incisional</u> wounds before closure for the purpose of preventing SSI The panel suggests considering the use of irrigation of the <u>incisional</u> wound with an aqueous povidone-iodine (PVP-I) solution before closure for the purpose of preventing SSI, particularly in clean and clean-contaminated wounds The panel suggests that antibiotic <u>incisional</u> wound irrigation before closure should not be used for the purpose of preventing SSI. (Conditional recommendation, low quality of evidence) • Evidence from 5 RCTs shows that the antibiotic irrigation of the incisional wound has neither benefit nor harm in reducing SSI when compared to no irrigation or to saline irrigation.	 Routine use of topical antimicrobials is not recommended. Prophylactic topical administration of antimicrobials in the surgical incision is superior to placebo but not to parenteral administration, and it does not increase the efficacy of parenteral antimicrobials when used in combination. Limited high quality data assessing safety and efficacy is available

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Resources and References

CMS Quality Programs - <u>https://mmshub.cms.gov/about-quality/quality-at-CMS/quality-programs</u> TJC SSI Info - <u>https://www.jointcommission.org/resources/patient-safety-topics/infection-prevention-and</u> <u>-control/surgical-site-infections/</u> NSOIR https://www.facs.org/guality.programs/data.and.registrics/acs.psgip/

NSQIP - <u>https://www.facs.org/quality-programs/data-and-registries/acs-nsqip/</u> CDC/NHSN SSI Definitions - <u>https://www.cdc.gov/nhsn/pdfs/pscmanual/9pscssicurrent.pdf</u>

Sternal wound infection prevention guidance - [1]

Recent PJI reviews and guidelines - [2-4]

Guidelines informing general SSI prevention and surgical antimicrobial prophylaxis - [5–9], also WHO book/resources available-

https://www.who.int/teams/integrated-health-services/infection-prevention-control/surgica I-site-infection

Recent SSI prevention review - [10]

Presumptive antibiotics in trauma - [11]

Pre-op S. aureus screening and	decolonization (se	elect pieces, see S	SSI guidelines	and reviews
too): [1,12–17]				

Pre-op asymptomatic bacteriuria: [18–21]

Periop PCN allergy delabeling/increasing use of first-line abx: [22–42]

PK deep dive into optimal start time of pre-op abx PTI (in addition to review contained in 2013 guidelines above): [43–64]

Postop abx: [65–75]

Bone cement/spacer abx (select examples): [76–88] Other loca/topical intra-op abx (select examples): [89–96]

Comprehensive ortho surgery ASP I lead: [97] Other periop ASPs: [98–101] Periop clinical pharmacy practice: [102]

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Question Report	
Report Generated:	
Торіс	Actual Duration (minutes)
Premier April 4, 2023 Antimicrobial Stewardship Forum	84
Question Details	
Question	Answer
we need help with stopping post-op antibiotics but ASHP has not updated their guidelines. Is there a good guideline that states which surgeries should have NO post-op abx use and which ones should and for how long?	Great question. The ACS/SIS guidelines and the CDC guidelines discuss this in the greatest detail among current guidelines, but procedure-specific assessment of primary literature is still important in my opinion. See my supplemental handout "antimicrobial prophylaxis guideline comparison" for a summary of what the various guidelines say on this specific topic. I recommend working with surgical teams more longitudinally to assess their antibiotic use for a particular type of procedure to include the optimal duration of prophylaxis, rather than trying to get all surgeons/procedures to stop postop antibiotic prescribing. This strategy will help you work with a more focused group of providers where you can collaboratively review the evidence for risk/benefit in that particular patient population. For example, in TJA, there is mounting evidence that extended postop antibiotics (i.e. oral antibiotics at discharge after TJA) can effectively reduce PJI, though this comes at the cost of selecting for resistent pathogens in subsequent PJIs (see the references in slide 63 "Right duration"). We're not going to get the surgeons to completely abandon this practice for this procedure and there isn't a definite evidence-based answer. So we collaborated with them to review the studies and limit duration as much as we can and also limit to certain patients with risk factors rather than universal protocols.
For an ortho patient with an MRSA infection due to hardware in the ankle, would it be reasonable to treat with Daptomycin vs Vancomycin simply due to desire to avoid vanco monitoring and multiple daily dosing?	If treating an MRSA bone/joint infection then I think either could be reasonable based on your local stewardship practices and institutional resources. It may also be reasonable to treat with an oral agent based on the results of the OVIVA trial though too! (https://www.nejm.org/doi/full/10.1056/NEJMoa1710926)
Do you have any tips for reminding anesthesia to redose intraoperatively? Is there anything that can be incorporated into EMR? What works at your institution?	Yes we have for a while now had the circulator RNs (who manage the "time-out") also keep track of this and cue the anesthesia team, but now we have also build into Epic when these intervals are approaching and so the anesthesia providers do get an alert on their screen. So between these two mechanisms our compliance with intraop dosing appears very high
Not a question but just a comment - we have a surgeon who always wants a vial of polymyxin B for spinal surgeries!	This is likely for a common spinal surgery irrigation- this is an old practice among old spine surgeons in my experience! ;) All dates back to this study with "zero infection rate" https://pubmed.ncbi.nlm.nih.gov/9736080/
Are there other options for pre-op decolonization? At a confernce there was a vendor that promoting an intranasal alcohol product just before surgery for decolonization.	Yes there are, I have read some about the premade alcohol nasal swab product which does seem infinitely more convenient and comfortable, but I'm just not sure the comparative efficacy data is there yet to change practice for us. I have been assessing this though
In your practice, how do long do you see post op prophylaxis abx being given for? Because by the book, it would be 24hrs but in my practice, the practioners give beyond that time frame.	I would say we're pretty good at =24 hours for the vast majority of procedures, and 48hours for CTS, and total joints are their own constant evolution on this front (see the first question response above)</td
In your practice, have you see prolonged pre-op abx given for "high risk" patients prior to urological procedures like up to 1 month in advance?	that seems a bit excessive to me! Pretty much chronic suppression at that point, which doesn't seem ideal. My experience has been that the urologist checks urine Cx in office a couple/few weeks prior to procedure and treat for 7-10 day courses prior to surgery usually, and then we will still consider covering the organism in the preop abx regimen