

Stanford Health Care Antimicrobial Dosing Reference Guide

This document is also located on the SHC Intranet (<http://portal.stanfordmed.org/depts/AntimicrobialStewardshipProgram>) and <http://bugsanddrugs.stanford.edu> · ABX Subcommittee Approved: January 2025

Formulas for dosing weights: Ideal body weight IBW (male) = 50kg + (2.3 x height in inches > 60 inches) ·
 Ideal body weight IBW (female) = 45kg + (2.3 x height in inches > 60 inches) · Adjusted Body Weight ABW (kg) = IBW + 0.4 (TBW – IBW)
 Obesity: BMI ≥ 30 kg/m²

Drug	CrCl > 50 mL/min		CrCl 10 – 50 mL/min		CrCl < 10 mL/min		Intermittent Hemodialysis (IHD) <i>Assumes thrice weekly dialysis</i>	CRRT	
	CrCl > 50	CrCl 25 – 50	CrCl < 25	CrCl < 10	IHD	CRRT			
Acyclovir (IV) ¹⁻⁷ (Use adjusted BW for obesity)	Prophylaxis								
	BMT	250 mg/m ² IV q12h	125 mg/m ² IV q12h	125 mg/m ² IV q24h	62.5 mg/m ² IV q24h	62.5 mg/m ² IV q24h	125 mg/m ² IV q12h		
	Hematology/Oncology	2 mg/kg IV q12h	2 mg/kg IV q12h	2 mg/kg IV q24h	1 mg/kg IV q24h	1 mg/kg IV q24h	2 mg/kg IV q12h		
	Treatment								
	General (e.g. mucocutaneous HSV)	5 mg/kg IV q8h	5 mg/kg IV q12h	5 mg/kg IV q24h	2.5 mg/kg IV q24h	2.5 mg/kg IV q24h	5 – 10 mg/kg IV q12h		
Severe (e.g. CNS/ocular/disseminated HSV infections, Zoster)	10 mg/kg IV q8h	10 mg/kg IV q12h	10 mg/kg IV q24h	5 mg/kg IV q24h	5 mg/kg IV q24h	10 mg/kg IV q12h			
Acyclovir (PO) ^{1,2,7}	Prophylaxis								
	BMT	800 mg PO BID	400 mg PO BID	200 mg PO BID	200 mg PO daily	200 mg PO daily	No data		
	Hematology/Oncology	400 mg PO BID	400 mg PO BID	200 mg PO BID	200 mg PO daily	200 mg PO daily	No data		
	Treatment								
	Mucocutaneous HSV	400 mg PO q8h Alt: 200 mg 5x daily	200 mg PO q8h	200 mg PO q12h	200 mg PO q12h	200 mg PO q12h	No data		
VZV	800 mg PO q4h (or 5x daily) Consider valacyclovir for less frequent dosing	800 mg PO q8h	800 mg PO q12h	800 mg PO q12h	800 mg PO q12h	No data			
Amikacin (IV) ^{1,2,5,8,9} (Use adjusted BW for obesity)	See Aminoglycoside Dosing Protocol								
Amoxicillin (PO) ^{1,2}	Usual dose: 500 mg PO q8h or 1,000 mg PO q8-12h CAP: 1,000 mg PO q8h Procedural ppx: 2,000 mg PO x 1	Normal Dose	CrCl 10-29	CrCl <10	IHD	No data			
		1,000 mg PO q8h	1,000 mg PO q12h	500 mg PO q12h	500 mg PO q12h				
		875 - 1,000 mg PO q12h	500 mg PO q12h	500 mg PO q12-24h	500 mg PO q12-24h				
		500 mg PO q8h	500 mg PO q12h	500 mg PO q12-24h	500 mg PO q12-24h				
Amoxicillin/clavulanate (PO) ^{1,2,10-12} (See obesity dosing guide)	Usual dose: 500 mg PO q8h or 875 mg PO q12h CAP: 875 mg PO q12h IAI / Uncomplicated GNR bacteremia (oral step-down alternative): up to 875 mg PO q8h	CrCl 10 – 30: 500 mg PO q12h IAI / Uncomplicated GNR bacteremia (oral step-down alternative): up to 875 mg PO q12h	CrCl < 10: 500 mg PO q24h IAI / Uncomplicated GNR bacteremia (oral step-down alternative): up to 875 mg PO q24h	500 mg PO q24h; For q24h regimen, dose after dialysis or administer additional dose at the end of dialysis	No data				
Amphotericin B Liposomal (IV) ^{1,2} (See obesity dosing guide)	3 – 5 mg/kg/day	No change	No change	No change	No change				

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD) <i>Assumes thrice weekly dialysis</i>	CRRT		
Ampicillin (IV) ¹⁻³ (See obesity dosing guide)	<u>Mild/uncomplicated:</u> 1 – 2 g IV q6h <u>Meningitis/endovascular/PJI:</u> 2 g IV q4h	<u>Mild/uncomplicated:</u> 1 g IV q6–8h <u>Meningitis/endovascular /PJI:</u> 2 g IV q6–12h	<u>Mild/uncomplicated:</u> 1 g IV q12h <u>Meningitis/endovascular /PJI:</u> 2 g IV q12–24h; or 1 g IV q8h	<u>Mild/uncomplicated:</u> 1 g IV q12h <u>Meningitis/endovascular/PJI:</u> 2 g IV q12–24h	CVVH: 2 g IV q8–12H CVVHDF: 2 g IV q6–8h <u>Meningitis/endovascular /PJI:</u> 2 g IV q6h		
Ampicillin/sulbactam (IV) ^{1-3,5,13}		CrCl >30:	CrCl 15-30:	CrCl < 15:	IHD	CRRT	
	<u>Mild/uncomplicated</u>	1.5 g IV q6h	1.5 g IV q12h	1.5 g IV q24h	1.5 g IV q24h	3 g IV q12h	
	<u>Systemic</u>	3 g IV q6h	3 g IV q12h	3 g IV q24h	3 g IV q24h	3 g IV q8h	
	<u>Acinetobacter baumannii</u> For more resistant <i>Acinetobacter baumannii</i> infections, consider higher dosing regimens	3 g IV q4h	3 g IV q8h	3 g IV q12h	3 g IV q12h	3 g IV q6h	
Azithromycin (IV/PO) ^{1,2}	500 mg IV/PO q24h	No change	No change	No change	No change		
Aztreonam (IV) ^{1-3,14} (See obesity dosing guide)	1 – 2 g IV q8h <u>Severe/Meningitis:</u> 2 g IV q6–8h	<u>CrCl < 30:</u> 1 g IV q8h <u>Severe/Meningitis:</u> 1 g IV q6–8h	500 mg IV q8h <u>Severe/Meningitis:</u> 1g IV q12h	1 g IV q24h <u>Severe/Meningitis:</u> 1 g IV q12h	2 g IV load, then 1 g IV q8h – or – 2 g IV q12h		
Caspofungin (IV) ^{1,2,15,15-17} (See obesity dosing guide)	70 mg IV x 1, then 50 mg IV q24h 70 mg IV q24h if on phenytoin, rifampin, other strong enzyme inducers <u>Endocarditis/Endovascular:</u> 150 mg IV q24h Dosage adjustments are not required for Child-Pugh B or C cirrhosis			No change	No change		
Cefazolin (IV) ^{1-5,18-20} (See obesity dosing guide)	<u>CrCl ≥ 35 mL/min:</u> <u>Mild/moderate:</u> 1 g IV Q8H <u>Severe:</u> 2 g IV Q8H	<u>CrCl 10 – 34 mL/min:</u> <u>Mild/moderate:</u> 1 g IV Q12H <u>Severe:</u> 2 g IV Q12H	<u>Mild/moderate:</u> 1 g IV Q24H <u>Severe:</u> 2 g IV Q24H	1 g IV Q24H <i>Dose daily, but after HD on HD days</i> <u>alt:</u> 2g/2g/3g IV post-HD only	2 g IV Q12H		
Cefepime (IV) ^{1-3,5,21-23} (See obesity dosing guide)	Extended Infusion (4-hour infusion)				0.5 – 1 g IV Q24H <i>Dose daily, but after HD on HD days</i> <u>alt:</u> 2 g IV post-HD only	2 g IV load, then 1 g IV Q8H (4-hour infusion)	
		CrCl > 60	CrCl 30 – 60	CrCl < 11-29			CrCl < 10
	General	1 g IV Q8H or 2 g IV Q12H	1 g IV Q12H or 2 g IV Q24H	1 g IV Q24H			500 mg IV Q24H
	Pulmonary/ Neutropenic Fever/ CNS/ confirmed Pseudomonal infection/ Severe infections	2 g IV Q8H	2 g IV Q12H	1 g IV Q12H	1 g IV q24h		
Cefiderocol (IV) ^{1,2} (SHC Restriction)	<u>CrCL > 120:</u> 2 g IV q6h <u>CrCL 60 -120:</u> 2 g IV q8h	<u>CrCL 30 – 60:</u> 1.5 g IV q8h <u>CrCL 15 – 30:</u> 1 g IV q8h	<u>CrCL < 15:</u> 750 mg IV q12h	750 mg IV q12h	Effluent Flow Rate	Dose	
					≤ 2L/hr	1.5 g IV q12h	
					2.1–3 L/hr	2 g IV q12h	
					3.1–4 L/hr	1.5 g IV q8h	
					≥4.1 L/hr	2 g IV q8h	
Shown as Effluent Dose (mL/kg/hr) in Epic							

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD) <i>Assumes thrice weekly dialysis</i>	CRRT				
Cefpodoxime (PO) ^{1,2}	<u>Uncomplicated cystitis:</u> 100 mg PO q12h <u>CAP/bronchitis:</u> 200 mg PO q12h <u>Skin/soft tissue:</u> 400 mg PO q12h	<u>CrCl < 30:</u> same dose q24h		Same dose, administered post-HD only	No data				
Ceftaroline (IV) ^{1,2,24} (SHC Restriction)		CrCl > 50	CrCl 30 – 50	CrCl 15 – 30	CrCl < 15	200 mg IV q8–12h <u>Endocarditis/S. aureus bacteremia/ SDD:</u> 200 mg IV q8–12h administered over 2-hr	No data		
	General	600 mg IV q12h	400 mg IV q12h	300 mg IV q12h	200 mg IV q12h				
	Endocarditis/ <i>S. aureus</i> bacteremia, Susceptible-dose dependent (SDD)	600 mg IV q8h administered over 2-hr	400 mg IV q8h administered over 2-hr	300 mg IV q8h administered over 2-hr	200 mg IV q8h administered over 2-hr				
Ceftazidime (IV) ^{1–3,25}	<u>Usual dose:</u> 1 – 2 g IV q8h <u>Severe:</u> 2 g IV q8h	<u>CrCl 30 – 50:</u> 1 – 2 g IV q12h <u>CrCl 16 – 30:</u> 1 – 2 g IV q24h <u>CrCl 6 – 15:</u> 0.5 – 1 g IV q24h	<u>CrCl < 5:</u> 0.5 g IV q24h	0.5 – 1 g IV q24h <i>Dose daily, but after HD on HD days</i> <u>alt:</u> 1 – 2 g IV q48–72h or 1 g IV post-HD only TIW	2 g IV load, then 1 g IV q8h – or – 2 g IV q12h				
Ceftazidime/avibactam (IV) ^{1,2,26–29} (SHC Restriction)	2.5 g IV q8h	<u>CrCl 31 – 50:</u> 1.25 g IV q8h <u>CrCl 16 – 30:</u> 0.94 g IV q12h <u>CrCl 6 – 15:</u> 0.94 g IV q24h	<u>CrCl < 5:</u> 0.94 g IV q48h	0.94 g IV q24–48h <i>Dose daily, but after HD on HD days</i>	1.25 g IV q8h 2.5g IV q8h if MIC > 4 mcg/mL or deep-seated				
Ceftolozane/tazobactam (IV) ^{1,2,30–33} (SHC Restriction)		CrCl > 50	CrCl 30 – 50	CrCl 15 – 29	CrCl < 15	IHD	CRRT		
	Cystitis	1.5 g IV q8h	750 mg IV q8h	375 mg IV q8h	750 mg IV load, then 150 mg IV q8h	750 mg IV load, then 150 mg IV q8h	1.5 g IV q8h		
	HAP, VAP, Systemic pseudomonal infection, CF exacerbation	3 g IV q8h	1.5 g IV q8h	750 mg IV q8h	2.25 g IV load, then 450 mg IV q8h	2.25 g IV load, then 450 mg IV q8h	3 g IV q8h		
Ceftriaxone (IV) ^{1,2,34}	1 – 2 g IV q24h <u>Endovascular/osteomyelitis/PJI:</u> 2 g IV q24h <u>Meningitis, E. faecalis endocarditis:</u> 2 g IV q12h		No change		No change		No change		
Cephalexin (PO) ^{1,2,35}	250 – 1000 mg PO Q6H <u>Uncomplicated cystitis:</u> 500 mg PO Q12H <u>Complicated cystitis/ Cellulitis/ SSTI:</u> 500 mg PO Q6H	<u>CrCl 15 – 29:</u> 250 mg PO Q8–12H <u>CrCl 5 – 14:</u> 250 mg PO Q24H		500 mg PO Q24H <i>Dose daily, but after HD on HD days</i>		No data			
Ciprofloxacin (IV/PO) ^{1–4,28,36} (See obesity dosing guide)		CrCl > 50	CrCl 30 – 50	CrCl < 30		200 – 400 mg IV q24h 250 – 500 mg PO q24h <i>Dose daily, but after HD on HD days</i>		400 mg IV q12h 500 mg PO q12h <u>Severe infection with <i>A. baumannii</i> or <i>P. aeruginosa</i>:</u> 400 mg IV q8–12h	
	General infections	400 mg IV q12h 500 mg PO q12h	Same	400 mg IV q24h 500 mg PO q24h					
	Pseudomonas, severe	400 mg IV q8h 750 mg PO q12h	400 mg IV q8–12h 500 mg PO q12h	400 mg IV q24h 500 mg PO q24h					
Clindamycin (IV/PO) ^{1,2} (See obesity dosing guide)	600 – 900 mg IV q8h 150 – 450 mg PO q6h	No change		No change		No change		No change	

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD) <i>Assumes thrice weekly dialysis</i>	CRRT								
Dalbavancin (IV) ^{1,37} (SHC Restriction)	Indication	CrCL > 30	CrCl < 30	IHD	CRRT								
	Skin/Soft Tissue	<u>Preferred:</u> 1,500 mg IV x 1 <u>Alternative:</u> 1,000 mg IV x 1 followed by 500 mg x1 1-week later	<u>Preferred:</u> 1,125 mg IV x 1 <u>Alternative:</u> 750 mg IV x 1 followed by 375 mg x1 1-week later	<u>Preferred:</u> 1,500 mg IV x 1 <u>Alternative:</u> 1,000 mg IV x 1 followed by 500 mg x1 1-week later	No data								
Daptomycin (IV) ^{1,2,23,38-45} (SHC Restriction) (Use adjusted BW for obesity)	Indication	CrCL > 30	CrCl < 30	IHD	CRRT								
	Skin/Soft Tissue	4 – 6 mg/kg IV q24h	4 – 6 mg/kg IV q48h	6 mg/kg post-HD only or 6/6/9 mg/kg post-HD only <u>alt:</u> 4 – 6 mg/kg IV q48h	6 mg/kg IV q24h								
	Bacteremia/Endovascular	8 mg/kg IV q24h	8 mg/kg IV q48h	8 mg/kg post-HD <u>alt:</u> 8 mg/kg IV q48h	6 – 8 mg/kg IV q24h								
E. faecium Infection – consult ID	10 – 12 mg/kg IV q24h	10 – 12 mg/kg IV q48h	8 – 10 mg/kg post-HD <u>alt:</u> 8 – 10 mg/kg IV q48h	8 mg/kg IV q24h Doses > 8 mg/kg q24h increase the risk of CPK elevations and myopathy. Caution, clinical judgment, and frequent CPK monitoring, including a baseline value, should be used if pursuing as high as 10 to 12 mg/kg every 24 hours (Hoff 2020)									
Doxycycline (IV/PO) ^{1,2}	Load: 200 mg x 1 for severe infections 100 mg IV/PO q12h	No change	No change	No change	No change								
Ertapenem (IV/IM) ^{1,2,46-48}	1 g IV q24h	<u>CrCl <30:</u> 500 mg IV q24h	500 mg IV q24h	500 mg IV q24h <i>Dose daily, but after HD on HD days</i> <u>alt:</u> 500 - 1000 mg IV post-HD (low vs. high-flux HD, degree of renal failure, residual UOP)	1 g IV q24h								
Ethambutol (PO) ^{1,5,49,50} (Use lean BW for obesity) (See footnote for lean BW equation)	<u>Dose range:</u> 15 – 25 mg/kg/day (max dose: 1,600 mg/day) <table border="1"> <tr> <th>Lean body weight</th> <th>Dose</th> </tr> <tr> <td>40 – 55 kg</td> <td>800 mg</td> </tr> <tr> <td>56 – 75 kg</td> <td>1,200 mg</td> </tr> <tr> <td>76 – 90 kg</td> <td>1,600 mg</td> </tr> </table>	Lean body weight	Dose	40 – 55 kg	800 mg	56 – 75 kg	1,200 mg	76 – 90 kg	1,600 mg	<u>CrCl 10 – 50:</u> 15 – 25 mg/kg PO q24–36h	<u>CrCl < 10:</u> 15 – 25 mg/kg PO q48h	15 – 25 mg/kg PO 3 times per week post-HD <i>Administer after HD only</i>	15 – 25 mg/kg PO q24–36h
Lean body weight	Dose												
40 – 55 kg	800 mg												
56 – 75 kg	1,200 mg												
76 – 90 kg	1,600 mg												
Fidaxomicin (PO) ^{1,2}	200 mg q12h x 10 days	No change	No change	No change	No change								

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD) <i>Assumes thrice weekly dialysis</i>	CRRT		
Fluconazole (IV/PO) ^{1-4,17,28,51-53} (See obesity dosing guide)	Indication	CrCl > 50	CrCl ≤ 50	HD	CRRT		
	Oropharyngeal candidiasis, Candida peritonitis	Load 200 mg x1, then 100-200mg Q24H	Load 200 mg x1, then 100mg Q24H	Load 200 mg x 1 dose, Then 200mg Q48H <i>Dose q48h, but after HD on HD days</i>	Load 400 mg x1, then 100 – 200mg Q24H‡		
	Esophageal, osteoarticular candidiasis, Candida pyelonephritis) See below for <i>C. glabrata</i>	400 mg (or 6 mg/kg) Q24H	400 mg (or 6 mg/kg) IV/PO x1, then 200 mg (or 3 mg/kg) Q24H	Load 400 mg (or 6 mg/kg) x 1 dose, then 400 mg (or 6 mg/kg) post-HD or 200mg (or 3 mg/kg) Q24H <i>Dose q24h, but after HD on HD days</i>	Load 800 mg (or 12mg/kg) x 1 dose, then 400mg (or 6mg/kg) Q24H‡		
	Severe Candidiasis: Candidemia/CNS/ endophthalmitis/endovascular/ intra-abdominal Strongly recommend ID consult for cryptococcosis, coccidioidomycosis, severe candidiasis, etc.	Load 800 mg (or 12 mg/kg) x 1 dose, then 400 – 800 mg* IV/PO Q24H <i>C. glabrata (SDD)†</i> : 800 mg (or 12 mg/kg) Q24H	Load 800 mg (or 12 mg/kg) x 1 dose, then 200 – 400 mg* IV/PO (or 3 – 6 mg/kg) Q24H <i>C. glabrata (SDD)†</i> : Load 800 mg (or 12 mg/kg) x 1 dose, then 400 mg (or 6 mg/kg) Q24H	Load 800 mg (or 12 mg/kg) x 1 dose, then 400 – 800 mg* post-HD or 200 – 400 mg* (3 – 6 mg/kg) Q24H. <i>Dose q24h, but after HD on HD days</i> <i>C. glabrata (SDD)†</i> : Load 800 mg (or 12 mg/kg) x 1, then 800 mg (or 12mg/kg) post-HD or 400 mg (or 6 mg/kg) Q24H <i>Dose q24h, but after HD on HD days</i>	Load 800 mg (or 12 mg/kg) - 1200 mg* x 1 dose, then 400 – 800 mg (or 6 – 12 mg/kg) IV/PO daily‡‡; 1200 mg given in 2 divided doses <i>C. glabrata (SDD)†‡</i> : Load 800 mg (or 12 mg/kg) - 1200 mg* x 1 dose, then 800 mg (or 12 mg/kg) daily‡; 1200mg given in 2 divided doses		
* Consider higher end of dose range for MIC = 4 (SDD) for <i>C. albicans</i> , <i>C. tropicalis</i> , <i>C. parapsilosis</i> , CrCl > 130 ml/min. Endophthalmitis, endocarditis/endovascular- consult ID. †SDD = susceptible-dose dependent; all <i>C. glabrata</i> isolates are considered SDD or resistant. Limited data on isolates with MIC ≥ 16, consider consultation with ID ‡Some experts recommend doubling the dose (up to max dose 1200mg) for CRRT based on high-flux dialyzers and effluent flow rates of 20 to 25 mL/kg/hour Weight-based dosing may be used for extremes of weight and in consideration of dose-related ADRs e.g. QTc prolongation.							
Foscarnet (IV) ^{1,2,54-56} (Use adjusted BW for obesity) Adj CrCl (mL/min/kg) $\left(\frac{140 - \text{age}}{\text{SCR} \times 72}\right) \times (0.85 \text{ if female})$	CrCl (mL/min/kg)	CMV induction		CMV maintenance		HSV	
	> 1.4	60 mg/kg IV q8h	90 mg/kg IV q12h	90 mg/kg IV q24h	120 mg/kg IV q24h	40 mg/kg IV q12h	40 mg/kg IV q8h
	> 1.0 – 1.4	45 mg/kg IV q8h	70 mg/kg IV q12h	70 mg/kg IV q24h	90 mg/kg IV q24h	30 mg/kg IV q12h	30 mg/kg IV q8h
	> 0.8 – 1.0	50 mg/kg IV q12h	50 mg/kg IV q12h	50 mg/kg IV q24h	65 mg/kg IV q24h	20 mg/kg IV q12h	35 mg/kg IV q12h
	> 0.6 – 0.8	40 mg/kg IV q12h	80 mg/kg IV q24h	80 mg/kg IV q48h	105 mg/kg IV q48h	35 mg/kg IV q24h	25 mg/kg IV q12h
	> 0.5 – 0.6	60 mg/kg IV q24h	60 mg/kg IV q24h	60 mg/kg IV q48h	80 mg/kg IV q48h	25 mg/kg IV q24h	40 mg/kg IV q24h
	≥ 0.4 – 0.5	50 mg/kg IV q24h	50 mg/kg IV q24h	50 mg/kg IV q48h	65 mg/kg IV q48h	20 mg/kg IV q24h	35 mg/kg IV q24h
	< 0.4	Not recommended		Not recommended		Not recommended	
IHD	45 – 60 mg/kg/dose IV post-HD only		No data		No data		
CRRT	No data						
Ganciclovir (IV) ^{1,2} (Use adjusted BW for obesity)	CMV	CrCl >70*	CrCl >50	CrCl >25	CrCl >10	CrCl <10	I: 1.25 mg/kg IV post HD only M: 0.625 mg/kg IV post HD only I: 2.5 mg/kg IV q12h M: 2.5 mg/kg IV q24h
	Induction (I)	5 mg/kg IV q12h	2.5 mg/kg IV q12h	2.5 mg/kg IV q24h	1.25 mg/kg IV q24h	1.25 mg/kg IV 3x/week	
	Maintenance (M)	5 mg/kg IV q24h	2.5 mg/kg IV q24h	1.25 mg/kg IV q24h	0.625 mg/kg IV q24h	0.625 mg/kg IV 3x/week	
*Manufacturer's CrCl cutoffs. Please refer to BMT protocols if applicable							
Gentamicin (IV) ^{1,3,57} (Use adjusted BW for obesity)	Refer to Aminoglycoside Dosing Guide						
Imipenem/Cilastatin (IV) ¹ (SHC Restriction)		CrCL >60	CrCL 30 – 59	CrCL 15 – 29	CrCL < 10		250 – 500 mg IV q12h 1g load, then 500 mg IV q6h
	General	500 mg IV q6H or 1g IV q8h	500 mg IV q8h	500 mg IV q12h	Not recommended unless dialysis initiated within 48 hrs		
	NTM	1,000 mg IV q12H	750 mg IV q12H	500 mg IV q12H			
Isavuconazole (IV/PO) ^{1,2}	Initial: 372 mg IV/PO q8h x 6 doses Maintenance: 372 mg IV/PO q24h	No change	No change	No change	No change	No change	

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD) <i>Assumes thrice weekly dialysis</i>	CRRT	
Isoniazid (PO) ^{1,2,49,50}	300 mg PO q24h (5 mg/kg/day)	No change	No change	No change	No change	
Levofloxacin (IV/PO) ¹⁻⁴		CrCl ≥ 50	CrCl 20 – 49	CrCl < 20	IHD	CRRT
	Cystitis	250 mg q24h	No change	No change	No change; give dose <i>after HD on HD days</i>	No change
	Mild to moderate diabetic foot infection (DFI)(non-Pseudomonal), prostatitis Severe/PNA/complicated UTI/osteomyelitis/PJI/ moderate to severe DFI /intra-abdominal/Pseudomonas/ Stenotrophomonas:	500 mg q24h 750 mg q24h	500mg x1 then 250 mg q24h 750 mg q48h	500 mg x1, then 250 mg q48h 750 mg x1, then 500 mg q48h	See CrCl < 20 ml/min <i>Dose q48h, but after HD on HD days</i> See CrCl < 20 ml/min <i>Dose q48h, but after HD on HD days</i>	500 mg x1, then 250 mg q24h or 500mg q48h 750 mg x1, then 500 mg q48h or 750mg q48h
Linezolid (IV/PO) ^{1,2} (SHC Restriction)	600 mg IV/PO q12h	No change	No change	No change	No change	
Meropenem (IV) ^{1-4,58} (See obesity dosing guide)		CrCl > 50	CrCl 26 – 50	CrCl 10 – 25	CrCl < 10	500 mg IV q24h <u>CF/CNS:</u> 1 g IV q24h <i>Dose daily, but after HD on HD days</i>
	Usual dose* (FN, PNA, Pseudomonas) CF/ CNS infections	1 g IV q8h 2 g IV q8h	1 g IV q12h 2 g IV q12h	0.5 g IV q12h 1 g IV q12h	0.5 g IV q24h 1 g IV q24h	
Metronidazole (IV/PO) ^{1,2}	CNS infections, <i>C.difficile</i> , SSTI/necrotizing infection: 500 mg IV/PO q8h Intra-abdominal: 500mg q 8 – 12 h Severe hepatic impairment: 500 mg IV/PO q12h Caution with accumulation if CrCl < 30 particularly if used > 1-2 weeks					
Moxifloxacin (IV/PO) ^{1,2}	400 mg IV/PO q24h	No change	No change	No change	No change	
Nafcillin (IV) ^{1,2}	2 g IV q4h Hepatic Impairment: No specific dose adjustment provided by manufacturer. Dosage adjustment may be necessary in the setting of concomitant renal impairment; nafcillin primarily undergoes hepatic metabolism.					
Oseltamivir (PO) ^{1,2,59}		CrCl ≥ 60	CrCl 30 – 60	CrCl 10 – 30	CrCl ≤ 10	Prophylaxis: 30 mg PO x 1, then 30 mg PO after every other HD session Treatment: 30 mg PO x 1, then 30 mg PO post-HD only
	Prophylaxis Treatment	75 mg PO q24h 75 mg PO q12h	30 mg PO q24h 75mg x1, then 30 mg PO q12h	30 mg PO q48h 30 mg PO q24h	Not recommended in ESRD not on dialysis Not recommended in ESRD not on dialysis	
Penicillin G (IV) ^{1-3,5}	2 – 4 mu IV q4h <u>Dose range:</u> 12 – 24 million units/day continuous infusion or in divided doses every 4 to 6 hours	2 – 3 mu IV q4h	1 – 2 mu IV q6h	Mild: 0.5 – 1 mu IV q4–6h; or 1 – 2 mu IV q8–12h Severe: 2 mu IV q4–6h; or 4 mu IV q8–12h		
Piperacillin/tazobactam (IV) ^{1-4,60,61}		CrCl > 40	CrCl 20 – 40	CrCl < 20	General: <u>2.25 g IV q12h</u> Severe infections: <u>3.375 g IV q12h over 4-hr</u> <u>alt: 2.25 g IV q8h</u>	
	Extended-Infusion Dosing (4-hr infusion)	Extended infusion for CrCl > 20: 3.375 – 4.5 g IV q8h over 4h*		3.375 g IV q12h over 4h		
	General, CF, Pseudomonas, nosocomial PNA	SDD: 4.5g IV q8h (4-hr infusion)				
	Intermittent Dosing (30-minutes)	General	3.375 g IV q6h	2.25 g IV q6h		2.25 g IV q8h
	Severe/sepsis/CF/nosocomial PNA	4.5 g IV q6h	3.375 g IV q6h	2.25 g IV q6h	3.375 g IV q6h over 30-minutes Extended infusion: 3.375 – 4.5 g IV q8h over 4-hr	
*In select cases, higher piperacillin/tazobactam dosing may be warranted, e.g. sepsis, critically ill patients with severe or deep-seated infections, infections with MIC = 16 mg/L, obesity with weight > 120kg or BMI > 40, CrCl > 120 mL/min, or enhanced drug clearance such as those with cystic fibrosis: consider doses of 4.5 g IV q8h (infused over 4 hours) or q6h.						
Polymyxin B (IV) ^{1,2,62,63} (SHC Restriction) (Use adjusted BW for obesity)	Dosing presented as units (10,000 units = 1 mg) 20,000 – 25,000 units/kg IV load x 1, then 12,500 – 15,000 units/kg IV q12h			No data	No change	

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD) Assumes thrice weekly dialysis	CRRT												
Posaconazole (IV/PO) ^{1,2} (SHC Restriction [IV])	<table border="1"> <thead> <tr> <th colspan="2">Formulation</th> <th>Dose</th> </tr> </thead> <tbody> <tr> <td colspan="2">Oral Suspension (NF) <i>Suspension and Delayed-release tablets are not interchangeable</i></td> <td><u>Prophylaxis</u>: 200 mg PO q8h <u>Treatment</u>: 200 mg PO q6–8h</td> </tr> <tr> <td colspan="2">Delayed-release tablet <i>Suspension and Delayed-release tablets are not interchangeable</i></td> <td>300 mg PO q12h x 2 doses, then 300 mg PO q24h</td> </tr> <tr> <td colspan="2">Intravenous solution</td> <td>300 mg IV q12h x 2 doses, then 300 mg IV q24h</td> </tr> </tbody> </table>		Formulation		Dose	Oral Suspension (NF) <i>Suspension and Delayed-release tablets are not interchangeable</i>		<u>Prophylaxis</u> : 200 mg PO q8h <u>Treatment</u> : 200 mg PO q6–8h	Delayed-release tablet <i>Suspension and Delayed-release tablets are not interchangeable</i>		300 mg PO q12h x 2 doses, then 300 mg PO q24h	Intravenous solution		300 mg IV q12h x 2 doses, then 300 mg IV q24h		No change	No change
	Formulation		Dose														
Oral Suspension (NF) <i>Suspension and Delayed-release tablets are not interchangeable</i>		<u>Prophylaxis</u> : 200 mg PO q8h <u>Treatment</u> : 200 mg PO q6–8h															
Delayed-release tablet <i>Suspension and Delayed-release tablets are not interchangeable</i>		300 mg PO q12h x 2 doses, then 300 mg PO q24h															
Intravenous solution		300 mg IV q12h x 2 doses, then 300 mg IV q24h															
Refer to Antifungal TDM Guide																	
Pyrazinamide (PO) ^{1,2,49,50} (Use lean BW for obesity) (See footnote for lean BW equation)	Usual Dose: 25 mg/kg PO q24h		CrCl < 30: 25 mg/kg PO 3 times per week	25 mg/kg PO 3 times per week Administer after HD only	No data												
	<table border="1"> <thead> <tr> <th>Lean body weight</th> <th>Dose</th> </tr> </thead> <tbody> <tr> <td>40 – 55 kg</td> <td>1,000 mg</td> </tr> <tr> <td>56 – 75 kg</td> <td>1,500 mg</td> </tr> <tr> <td>76 – 90 kg</td> <td>2,000 mg</td> </tr> </tbody> </table> (max dose: 2,000 mg/day)	Lean body weight				Dose	40 – 55 kg	1,000 mg	56 – 75 kg	1,500 mg	76 – 90 kg	2,000 mg					
Lean body weight	Dose																
40 – 55 kg	1,000 mg																
56 – 75 kg	1,500 mg																
76 – 90 kg	2,000 mg																
Rifampin (IV/PO) ^{1,2,49,50,64–66} Capsule size: 150mg, 300mg	<u>TB</u> : 600 mg IV/PO q24h (≤ 45 kg: 10 mg/kg q24h)		No change	No change	No change												
	<u>Endocarditis</u> : 300 mg IV/PO q8h <u>PJI</u> : 300 – 450 mg IV/PO q12h <u>Vertebral Osteomyelitis</u> : 600 mg IV/PO q24h																
Tedizolid (IV/PO) ^{1,2,67} (SHC Restriction)	200 mg IV/PO q24h	No change	No change	No change	No change												
Tobramycin (IV) ^{1,2,57}	Refer to Gentamicin for dosing. See appendix for complete guidelines.																
Trimethoprim (TMP)/ Sulfamethoxazole (IV/PO) ^{1,2,4,68} (Use adjusted BW for obesity) SS = 80 mg TMP = 10 ml po soln DS = 160 mg TMP = 20ml po soln	<u>Uncomplicated cystitis</u> : 1 DS tab PO BID <u>SSTI</u> : 1 – 2 DS tab PO BID <u>S. aureus (Bone/Joint)</u> : 8-10 mg/kg/day TMP in divided doses (2 DS tabs PO BID) <u>Gram-negative bacteremia</u> : 8-10 mg/kg/day TMP in divided doses (2 DS tab PO BID) <u>Stenotrophomonas</u> : 10-15 mg/kg/day TMP divided q8-12h <u>PCP</u> : 15 mg/kg/day TMP divided q8h (~2 DS tab TID)	CrCl 15 – 30: Administer 50% of recommended dose	CrCl < 15: Use is not recommended, but if needed for PCP: 5 – 7.5 mg/kg TMP q24h (25-50% of usual dose)	25-50% of usual dose 2.5 – 5 mg/kg TMP q24h <u>PCP, Stenotrophomonas</u> 5 – 7.5 mg/kg TMP q24h Dose daily, but after HD on HD days <u>alt</u> : 5 – 15 mg/kg TMP post-HD only	5 – 10 mg/kg/day TMP divided q12h <u>Stenotrophomonas</u> 10-15 mg/kg/day TMP divided q8-12h <u>PCP</u> 15 mg/kg/day TMP divided q8h (~2 DS tab TID)												

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD) Assumes thrice weekly dialysis	CRRT		
Valacyclovir (PO) ^{1,2} Please refer to transplant protocols if applicable	VZV	CrCl > 30 CrCl >50: 1 g PO q8h CrCl 30-50: 1 g q12h	CrCl 10 – 30 1 g PO q24h	< 10 500 mg PO q24h	500 mg PO q24h Dose daily, but after HD on HD days	No data	
	Genital herpes	Initial episode: 1 g PO q12h Recurrent episode: 500 mg PO q12h	Initial episode: 1 g PO q24h Recurrent: 500 mg PO q24h	Initial/recurrent episode: 500 mg PO q24h			
	Herpes labialis	CrCl >50: 2 g PO q12h x 2 doses CrCl 30 – 50: 1 g PO q12h x 2 doses	500 mg PO q12h x 2 doses	500 mg PO x 1 dose			
Valganciclovir (PO) ^{1,2} Please refer to transplant protocols if applicable	Induction (14-21 days)	CrCl > 60 900 mg PO q12h	CrCl 40 – 59 450 mg PO q12h	CrCl 25 – 39 450 mg PO q24h	CrCl 10 – 24 450 mg PO q48h	CrCl < 10; IHD 200 mg PO 3x/week after HD only	CRRT No data
	Maintenance/prophylaxis	900 mg PO q24h	450 mg PO q24h	450 mg PO q48h	450 mg twice/week	100 mg PO 3x/week after HD only	No data
Vancomycin (IV) ^{1,2,69,70}	See Vancomycin Dosing Protocol						
Vancomycin PO ^{1,2,71}	Poor systemic absorption- used for the treatment of <i>Clostridium difficile</i> -associated diarrhea Mild/moderate/severe: 125 mg PO q6h Severe complicated (CDI-related septic shock, ileus, toxic megacolon): 500 mg PO q6h			No change	No change		
Voriconazole (IV/PO) ^{1,2,72,73} (Use adjusted BW for obesity)	IV: 6 mg/kg IV q12h x 2, then 4 mg/kg IV q12h PO: 400 mg PO q12h x 2, then 200 mg PO q12h	<ul style="list-style-type: none"> IV→PO conversion 1:1 (round to nearest tablet size- available in 200 mg and 50 mg tablets) Caution with IV: accumulation of IV vehicle cyclodextrin occurs. Consider PO if CrCl < 50 mL/min unless benefits justify risks of IV use. Please refer to Antifungal TDM Guide 					

Abbreviations: CAP = community acquired pneumonia; CRRT = continuous renal replacement therapy; FN = febrile neutropenia; HD = hemodialysis; LD = loading dose; MU = million units; PCP = pneumocystis jiroveci pneumonia; PNA = pneumonia; SCR = serum creatinine; TB = tuberculosis; TMP = trimethoprim; UF = ultrafiltration

CRRT dosing: doses listed are for CVVHDF and CVVHD modalities, which are the most common modes at SHC. Note that these are generally higher than doses used in CVVH.

LBW (men) = (1.10 x Weight(kg)) - 128 x (Weight²/(100 x Height(m))²)

LBW (women) = (1.07 x Weight(kg)) - 148 x (Weight²/(100 x Height(m))²)

LBW online calculator: <http://www.empr.com/medical-calculators/lean-body-weight-calculator/article/170219/>

References:

- Lexicomp Online. Accessed April 9, 2017. <http://online.lexi.com>
- MICROMEDEX®. Accessed April 9, 2017. <http://www.micromedexsolutions.com.laneproxy.stanford.edu/micromedex2/librarian>
- Heintz BH, Matzke GR, Dager WE. Antimicrobial Dosing Concepts and Recommendations for Critically Ill Adult Patients Receiving Continuous Renal Replacement Therapy or Intermittent Hemodialysis. *Pharmacother J Hum Pharmacol Drug Ther.* 2009;29. doi:10.1592/phco.29.5.562
- Trotman RL, Williamson JC, Shoemaker DM, Salzer WL. Antibiotic Dosing in Critically Ill Adult Patients Receiving Continuous Renal Replacement Therapy. *Clin Infect Dis.* 2005;41(8):1159-1166. doi:10.1086/444500
- Aronoff G, Bennett W, Berns J, et al. *Drug Prescribing in Renal Failure*. 5th ed. American College of Physicians; 2007.
- Turner RB, Cumpston A, Sweet M, et al. Prospective, Controlled Study of Acyclovir Pharmacokinetics in Obese Patients. *Antimicrob Agents Chemother.* 2016;60. doi:10.1128/aac.02010-15
- Tomblyn M, Chiller T, Einsele H, et al. Guidelines for Preventing Infectious Complications among Hematopoietic Cell Transplantation Recipients: A Global Perspective. *Biol Blood Marrow Transplant.* 2009;15(10):1143-1238. doi:10.1016/j.bbmt.2009.06.019
- Roger C, Wallis SC, Muller L, et al. Influence of Renal Replacement Modalities on Amikacin Population Pharmacokinetics in Critically Ill Patients on Continuous Renal Replacement Therapy. *Antimicrob Agents Chemother.* 2016;60. doi:10.1128/aac.00828-16
- Taccone FS, Backer D de, Laterre PF, et al. Pharmacokinetics of a loading dose of amikacin in septic patients undergoing continuous renal replacement therapy. *Int J Antimicrob Agents.* 2011;37. doi:10.1016/j.ijantimicag.2011.01.026

10. Mora Lopez L, Serra Pla S, Serra-Aracil X, Ballesteros E, Navarro S. Application of a modified Neff classification to patients with uncomplicated diverticulitis. *Colorectal Dis.* 2013;15(11):1442-1447. doi:10.1111/codi.12449
11. Biondo S, Golda T, Kreisler E, et al. Outpatient Versus Hospitalization Management for Uncomplicated Diverticulitis: A Prospective, Multicenter Randomized Clinical Trial (DIVER Trial). *Ann Surg.* 2014;259(1):38-44. doi:10.1097/SLA.0b013e3182965a11
12. Mora-López L, Ruiz-Edo N, Estrada-Ferrer O, et al. Efficacy and Safety of Nonantibiotic Outpatient Treatment in Mild Acute Diverticulitis (DINAMO-study): A Multicentre, Randomised, Open-label, Noninferiority Trial. *Ann Surg.* 2021;274(5):e435. doi:10.1097/SLA.0000000000005031
13. Tamma PD, Aitken SL, Bonomo RA, Mathers AJ, van Duin D, Clancy CJ. Infectious Diseases Society of America Guidance on the Treatment of AmpC β -Lactamase-Producing Enterobacterales, Carbapenem-Resistant *Acinetobacter baumannii*, and *Stenotrophomonas maltophilia* Infections. *Clin Infect Dis Off Publ Infect Dis Soc Am.* 2022;74(12):2089-2114. doi:10.1093/cid/ciab1013
14. Gerig JS, Bolton ND, Swabb EA, Scheld WM, Bolton WK. Effect of hemodialysis and peritoneal dialysis on aztreonam pharmacokinetics. *Kidney Int.* 1984;26. doi:10.1038/ki.1984.174
15. Gustot T, ter Heine R, Brauns E, Cotton F, Jacobs F, Brüggemann RJ. Caspofungin dosage adjustments are not required for patients with Child–Pugh B or C cirrhosis. *J Antimicrob Chemother.* 2018;73(9):2493-2496. doi:10.1093/jac/dky189
16. Roger C, Wallis SC, Muller L, et al. Caspofungin Population Pharmacokinetics in Critically Ill Patients Undergoing Continuous Venovenous Hemofiltration or Hemodiafiltration. *Clin Pharmacokinet.* 2017;56(9):1057-1068. doi:10.1007/s40262-016-0495-z
17. Pappas PG, Kauffman CA, Andes DR, et al. Clinical Practice Guideline for the Management of Candidiasis: 2016 Update by the Infectious Diseases Society of America. *Clin Infect Dis Off Publ Infect Dis Soc Am.* 2016;62(4):e1-50. doi:10.1093/cid/civ933
18. Stryjewski ME, Szczech LA, Benjamin DK, et al. Use of Vancomycin or First-Generation Cephalosporins for the Treatment of Hemodialysis-Dependent Patients with Methicillin-Susceptible *Staphylococcus aureus* Bacteremia. *Clin Infect Dis.* 2007;44. doi:10.1086/510386
19. Wong G, Briscoe S, McWhinney B, et al. Therapeutic drug monitoring of β -lactam antibiotics in the critically ill: direct measurement of unbound drug concentrations to achieve appropriate drug exposures. *J Antimicrob Chemother.* 2018;73(11):3087-3094. doi:10.1093/jac/dky314
20. Roberts JA, Udy AA, Jarrett P, et al. Plasma and target-site subcutaneous tissue population pharmacokinetics and dosing simulations of ceftazolin in post-trauma critically ill patients. *J Antimicrob Chemother.* 2015;70(5):1495-1502. doi:10.1093/jac/dku564
21. Crandon JL, Bulik CC, Kuti JL, Nicolau DP. Clinical Pharmacodynamics of Cefepime in Patients Infected with *Pseudomonas aeruginosa*. *Antimicrob Agents Chemother.* 2010;54. doi:10.1128/AAC.01183-09
22. Bauer KA, West JE, O'Brien JM, Goff DA. Extended-infusion ceftazidime reduces mortality in patients with *Pseudomonas aeruginosa* infections. *Antimicrob Agents Chemother.* 2013;57. doi:10.1128/AAC.02365-12
23. Hoff BM, Maker JH, Dager WE, Heintz BH. Antibiotic Dosing for Critically Ill Adult Patients Receiving Intermittent Hemodialysis, Prolonged Intermittent Renal Replacement Therapy, and Continuous Renal Replacement Therapy: An Update. *Ann Pharmacother.* 2020;54(1):43-55. doi:10.1177/1060028019865873
24. Vidaillet C, Leonard SN, Rybak MJ. In vitro activity of ceftaroline against methicillin-resistant *Staphylococcus aureus* and heterogeneous vancomycin-intermediate *S. aureus* in a hollow fiber model. *Antimicrob Agents Chemother.* 2009;53(11):4712-4717. doi:10.1128/AAC.00636-09
25. Loo AS, Neely M, Anderson EJ, Ghossein C, McLaughlin MM, Scheetz MH. Pharmacodynamic target attainment for various ceftazidime dosing schemes in high-flux hemodialysis. *Antimicrob Agents Chemother.* 2013;57(12):5854-5859. doi:10.1128/AAC.00474-13
26. Wenzler E, Bunnell KL, Bleasdale SC, Benken S, Danziger LH, Rodvold KA. Pharmacokinetics and Dialytic Clearance of Ceftazidime-Avibactam in a Critically Ill Patient on Continuous Venovenous Hemofiltration. *Antimicrob Agents Chemother.* 2017;61(7). doi:10.1128/AAC.00464-17
27. Soukup P, Faust AC, Edpuganti V, Putnam WC, McKinnell JA. Steady-State Ceftazidime-Avibactam Serum Concentrations and Dosing Recommendations in a Critically Ill Patient Being Treated for *Pseudomonas aeruginosa* Pneumonia and Undergoing Continuous Venovenous Hemodiafiltration. *Pharmacother J Hum Pharmacol Drug Ther.* 2019;39(12):1216-1222. doi:10.1002/phar.2338
28. Pistolesi V, Morabito S, Di Mario F, Regolisti G, Cantarelli C, Fiaccadori E. A Guide to Understanding Antimicrobial Drug Dosing in Critically Ill Patients on Renal Replacement Therapy. *Antimicrob Agents Chemother.* 2019;63(8). doi:10.1128/AAC.00583-19
29. Li L, Li X, Xia Y, et al. Recommendation of Antimicrobial Dosing Optimization During Continuous Renal Replacement Therapy. *Front Pharmacol.* 2020;11. doi:10.3389/fphar.2020.00786
30. Bremmer DN, Nicolau DP, Burcham P, Chunduri A, Shidham G, Bauer KA. Ceftolozane/Tazobactam Pharmacokinetics in a Critically Ill Adult Receiving Continuous Renal Replacement Therapy. *Pharmacotherapy.* 2016;36(5):e30-e33. doi:10.1002/phar.1744
31. Oliver WD, Heil EL, Gonzales JP, et al. Ceftolozane-Tazobactam Pharmacokinetics in a Critically Ill Patient on Continuous Venovenous Hemofiltration. *Antimicrob Agents Chemother.* 2016;60. doi:10.1128/aac.02608-15
32. Aguilar G, Ferriols R, Martínez-Castro S, et al. Optimizing ceftolozane-tazobactam dosage in critically ill patients during continuous venovenous hemodiafiltration. *Crit Care.* 2019;23. doi:10.1186/s13054-019-2434-5
33. Tamma PD, Aitken SL, Bonomo RA, Mathers AJ, van Duin D, Clancy CJ. Infectious Diseases Society of America 2022 Guidance on the Treatment of Extended-Spectrum β -lactamase Producing Enterobacterales (ESBL-E), Carbapenem-Resistant Enterobacterales (CRE), and *Pseudomonas aeruginosa* with Difficult-to-Treat Resistance (DTR-P. *aeruginosa*). *Clin Infect Dis Off Publ Infect Dis Soc Am.* 2022;75(2):187-212. doi:10.1093/cid/ciac268
34. Laville M, Mercatello A, Freney J, et al. Pharmacokinetics of ceftriaxone in hemodialysis. *Pathol Biol (Paris).* 1987;35(5 Pt 2):719-723.
35. Stevens DL, Bisno AL, Chambers HF, et al. Practice Guidelines for the Diagnosis and Management of Skin and Soft Tissue Infections: 2014 Update by the Infectious Diseases Society of America. *Clin Infect Dis.* 2014;59. doi:10.1093/cid/ciu296
36. Roger C, Wallis SC, Louart B, et al. Comparison of equal doses of continuous venovenous haemofiltration and haemodiafiltration on ciprofloxacin population pharmacokinetics in critically ill patients. *J Antimicrob Chemother.* 2016;71(6):1643-1650. doi:10.1093/jac/dkw043
37. Marbury T, Dowell JA, Seltzer E, Buckwalter M. Pharmacokinetics of dalbavancin in patients with renal or hepatic impairment. *J Clin Pharmacol.* 2009;49(4):465-476. doi:10.1177/0091270008330162
38. Dvorchik BH, Dampousse D. The pharmacokinetics of daptomycin in moderately obese, morbidly obese, and matched nonobese subjects. *J Clin Pharmacol.* 2005;45(1):48-56. doi:10.1177/0091270004269562
39. Pai MP, Norenberg JP, Anderson T, et al. Influence of morbid obesity on the single-dose pharmacokinetics of daptomycin. *Antimicrob Agents Chemother.* 2007;51(8):2741-2747. doi:10.1128/AAC.00059-07
40. Haselden M, Leach M, Bohm N. Daptomycin dosing strategies in patients receiving thrice-weekly intermittent hemodialysis. *Ann Pharmacother.* 2013;47(10):1342-1347. doi:10.1177/1060028013503110
41. Patel N, Cardone K, Grabe DW, et al. Use of pharmacokinetic and pharmacodynamic principles to determine optimal administration of daptomycin in patients receiving standardized thrice-weekly hemodialysis. *Antimicrob Agents Chemother.* 2011;55(4):1677-1683. doi:10.1128/AAC.01224-10
42. Falcone M, Russo A, Cassetta MI, et al. Daptomycin serum levels in critical patients undergoing continuous renal replacement. *J Chemother Florence Italy.* 2012;24(5):253-256. doi:10.1179/1973947812Y.0000000033
43. Preiswerk B, Rudiger A, Fehr J, Corti N. Experience with daptomycin daily dosing in ICU patients undergoing continuous renal replacement therapy. *Infection.* 2013;41(2):553-557. doi:10.1007/s15010-012-0300-3
44. Xu X, Khadzhynov D, Peters H, et al. Population pharmacokinetics of daptomycin in adult patients undergoing continuous renal replacement therapy. *Br J Clin Pharmacol.* 2017;83(3):498-509. doi:10.1111/bcp.13131
45. Dirolez J, Venisse N, Belmouaz S, Bauwens MA, Bridoux F, Beraud G. Pilot Pharmacokinetic Study of High-Dose Daptomycin in Hemodialysis Patients With Infected Medical Devices. *Am J Kidney Dis Off J Natl Kidney Found.* 2017;70(5):732-734. doi:10.1053/j.ajkd.2017.05.011
46. Geerlings CJC, de Man P, Rietveld AP, Touw DJ, Cohen Tervaert JW. A practical thrice weekly Ertapenem dosage regime for chronic hemodialysis patients? *Clin Nephrol.* 2013;80(4):312. doi:10.5414/cn108071

47. Hsaiky LM, Salinitri FD, Wong J, et al. Pharmacokinetics and investigation of optimal dose ertapenem in intermittent hemodialysis patients. *Nephrol Dial Transplant*. 2019;34(10):1766-1772. doi:10.1093/ndt/gfy166
48. Ueng YF, Wang HJ, Wu SC, Ng YY. A Thrice-Weekly Ertapenem Regimen Is Practical for Hemodialysis Patients. *Antimicrob Agents Chemother*. 2019;63(12). doi:10.1128/AAC.01427-19
49. Drug-Resistant Tuberculosis: A Survival Guide for Clinicians, 3rd edition | Curry International Tuberculosis Center. Accessed April 10, 2017. <http://www.currytbccenter.ucsf.edu/products/cover-pages/drug-resistant-tuberculosis-survival-guide-clinicians-3rd-edition>
50. Nahid P, Dorman SE, Alipanah N, et al. Official American Thoracic Society/Centers for Disease Control and Prevention/Infectious Diseases Society of America Clinical Practice Guidelines: Treatment of Drug-Susceptible Tuberculosis. *Clin Infect Dis Off Publ Infect Dis Soc Am*. 2016;63(7):e147-195. doi:10.1093/cid/ciw376
51. Muilwijk EW, Lange DW de, Schouten JA, et al. Suboptimal Dosing of Fluconazole in Critically Ill Patients: Time To Rethink Dosing. *Antimicrob Agents Chemother*. 2020;64(10). doi:10.1128/AAC.00984-20
52. Pea F, Lewis RE. Overview of antifungal dosing in invasive candidiasis. *J Antimicrob Chemother*. 2018;73(suppl_1):i33-i43. doi:10.1093/jac/dkx447
53. EM100 Connect - CLSI M27M44S ED3:2022. Accessed June 17, 2024. <https://em100.edaptivedocs.net/GetDoc.aspx?doc=CLSI%20M27M44S%20ED3:2022&scope=user>
54. Aweeka FT, Jacobson MA, Martin-Munley S, et al. Effect of renal disease and hemodialysis on fosfarnet pharmacokinetics and dosing recommendations. *J Acquir Immune Defic Syndr Hum Retrovirology Off Publ Int Retrovirology Assoc*. 1999;20(4):350-357.
55. Jayasekara D, Aweeka FT, Rodriguez R, Kalayjian RC, Humphreys MH, Gambertoglio JG. Antiviral therapy for HIV patients with renal insufficiency. *J Acquir Immune Defic Syndr* 1999. 1999;21(5):384-395.
56. MacGregor RR, Graziani AL, Weiss R, Grunwald JE, Gambertoglio JG. Successful fosfarnet therapy for cytomegalovirus retinitis in an AIDS patient undergoing hemodialysis: rationale for empiric dosing and plasma level monitoring. *J Infect Dis*. 1991;164(4):785-787.
57. Nicolau DP, Freeman CD, Belliveau PP, Nightingale CH, Ross JW, Quintiliani R. Experience with a once-daily aminoglycoside program administered to 2,184 adult patients. *Antimicrob Agents Chemother*. 1995;39(3):650-655.
58. Kuti JL, Dandekar PK, Nightingale CH, Nicolau DP. Use of Monte Carlo simulation to design an optimized pharmacodynamic dosing strategy for meropenem. *J Clin Pharmacol*. 2003;43(10):1116-1123. doi:10.1177/0091270003257225
59. Robson R, Buttimore A, Lynn K, Brewster M, Ward P. The pharmacokinetics and tolerability of oseltamivir suspension in patients on haemodialysis and continuous ambulatory peritoneal dialysis. *Nephrol Dial Transplant Off Publ Eur Dial Transpl Assoc - Eur Ren Assoc*. 2006;21(9):2556-2562. doi:10.1093/ndt/gfl267
60. Lodise TP, Lomaestro B, Drusano GL. Piperacillin-tazobactam for Pseudomonas aeruginosa infection: clinical implications of an extended-infusion dosing strategy. *Clin Infect Dis Off Publ Infect Dis Soc Am*. 2007;44(3):357-363. doi:10.1086/510590
61. Patel N, Scheetz MH, Drusano GL, Lodise TP. Identification of optimal renal dosage adjustments for traditional and extended-infusion piperacillin-tazobactam dosing regimens in hospitalized patients. *Antimicrob Agents Chemother*. 2010;54(1):460-465. doi:10.1128/AAC.00296-09
62. Sandri AM, Landersdorfer CB, Jacob J, et al. Population pharmacokinetics of intravenous polymyxin B in critically ill patients: implications for selection of dosage regimens. *Clin Infect Dis Off Publ Infect Dis Soc Am*. 2013;57(4):524-531. doi:10.1093/cid/cit334
63. Tsuji BT, Pogue JM, Zavascki AP, et al. International Consensus Guidelines for the Optimal Use of the Polymyxins: Endorsed by the American College of Clinical Pharmacy (ACCP), European Society of Clinical Microbiology and Infectious Diseases (ESCMID), Infectious Diseases Society of America (IDSA), International Society for Anti-infective Pharmacology (ISAP), Society of Critical Care Medicine (SCCM), and Society of Infectious Diseases Pharmacists (SIDP). *Pharmacother J Hum Pharmacol Drug Ther*. 2019;39(1):10-39. doi:10.1002/phar.2209
64. Baddour LM, Wilson WR, Bayer AS, et al. Infective Endocarditis in Adults: Diagnosis, Antimicrobial Therapy, and Management of Complications: A Scientific Statement for Healthcare Professionals From the American Heart Association. *Circulation*. 2015;132(15):1435-1486. doi:10.1161/CIR.0000000000000296
65. Osmon DR, Berbari EF, Berendt AR, et al. Diagnosis and management of prosthetic joint infection: clinical practice guidelines by the Infectious Diseases Society of America. *Clin Infect Dis Off Publ Infect Dis Soc Am*. 2013;56(1):e1-e25. doi:10.1093/cid/cis803
66. Berbari EF, Kanj SS, Kowalski TJ, et al. 2015 Infectious Diseases Society of America (IDSA) Clinical Practice Guidelines for the Diagnosis and Treatment of Native Vertebral Osteomyelitis in Adults. *Clin Infect Dis Off Publ Infect Dis Soc Am*. 2015;61(6):e26-46. doi:10.1093/cid/civ482
67. Flanagan S, Minassian SL, Morris D, et al. Pharmacokinetics of tedizolid in subjects with renal or hepatic impairment. *Antimicrob Agents Chemother*. 2014;58(11):6471-6476. doi:10.1128/AAC.03431-14
68. Nahata MC. Dosage regimens of trimethoprim/sulfamethoxazole (TPM/SMX) in patients with renal dysfunction. *Ann Pharmacother*. 1995;29(12):1300.
69. Rybak M, Lomaestro B, Rotschafer JC, et al. Therapeutic monitoring of vancomycin in adult patients: a consensus review of the American Society of Health-System Pharmacists, the Infectious Diseases Society of America, and the Society of Infectious Diseases Pharmacists. *Am J Health-Syst Pharm AJHP Off J Am Soc Health-Syst Pharm*. 2009;66(1):82-98. doi:10.2146/ajhp080434
70. Liu C, Bayer A, Cosgrove SE, et al. Clinical practice guidelines by the infectious diseases society of america for the treatment of methicillin-resistant Staphylococcus aureus infections in adults and children. *Clin Infect Dis Off Publ Infect Dis Soc Am*. 2011;52(3):e18-55. doi:10.1093/cid/ciq146
71. Cohen SH, Gerding DN, Johnson S, et al. Clinical Practice Guidelines for Clostridium difficile Infection in Adults: 2010 Update by the Society for Healthcare Epidemiology of America (SHEA) and the Infectious Diseases Society of America (IDSA). *Infect Control Hosp Epidemiol*. 2010;31(5):431-455. doi:10.1086/651706
72. Koselke E, Kraft S, Smith J, Nagel J. Evaluation of the effect of obesity on voriconazole serum concentrations. *J Antimicrob Chemother*. 2012;67(12):2957-2962. doi:10.1093/jac/dks312
73. Davies-Vorbrodt S, Ito JI, Tegtmeier BR, Dadwal SS, Kriengkauykiat J. Voriconazole serum concentrations in obese and overweight immunocompromised patients: a retrospective review. *Pharmacotherapy*. 2013;33(1):22-30. doi:10.1002/phar.1156

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Department of Pharmacy; 07/1998

B. Gatekeeper

Stanford Antimicrobial Stewardship Safety and Sustainability Program

C. Review and Renewal Requirement

This document will be reviewed every three years and as required by change of law or practice

D. Revision/Review History

Deepak Sisodiya, PharmD; 04/2005

Maggie Cudny, PharmD, BCOP; 04/2007, 01/2009
Katherine Miller, PharmD; 01/2009
Sean Carlton, PharmD, BCPS; 03/2010
Emily Mui, PharmD, BCIDP; 11/2010, 03/2011, 05/2012, 05/2013, 01/2014, 03/2017, 02/2019, 07/2019, 08/2019, 10/2019, 01/2020, 05/2020, 09/2020, 12/2021, 06/2022, 12/2022
Lina Meng, PharmD, BCPS, BCCCP; 11/2010, 03/2011, 03/2017, 08/2019, 10/2019, 05/2020, 09/2020, 12/2021, 06/2022, 1/2025 (fluconazole-oseltamivir)
Marisa Holubar, MD; 03/2017
Stan Deresinski, MD; 03/2017
Will Alegria, PharmD; 08/2019, 01/2020, 05/2020, 09/2020, 12/2021, 06/2022, 12/2022
David Ha, PharmD; 01/2020, 05/2020, 09/2020, 12/2021, 06/2022, 12/2022
David Epstein, MD; 06/2022
Dora Ho, MD, PhD; 06/2022
Brian Lu, PharmD; 12/2022

E. Approvals

Antimicrobial Subcommittee 09/2004, 04/2007, 01/2009, 11/2010, 03/2011, 05/2012, 05/2013, 01/2014, 03/2017, 07/2019, 10/2019; 01/2020, 09/2020, 08/2021, 12/2021, 06/2022, 12/2022, 1/2025
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