

# 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: September 2020

**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)

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD)	CRRT		
<b>Acyclovir (IV)</b> <sup>1-7</sup> (Use adjusted BW for obese)		CrCl > 50	CrCl 25 – 50	CrCl < 25	CrCl < 10	IHD	CRRT
	<b>Prophylaxis</b>						
	BMT	250 mg/m <sup>2</sup> q12h	125 mg/m <sup>2</sup> q12h	125 mg/m <sup>2</sup> q24h	62.5 mg/m <sup>2</sup> q24h	62.5 mg/m <sup>2</sup> q24h	125 mg/m <sup>2</sup> q12h
	Hematology/Oncology	2 mg/kg q12h	2 mg/kg q12h	2 mg/kg q24h	1 mg/kg q24h	1 mg/kg q24h	2 mg/kg q12h
	<b>Treatment</b>						
	General (e.g. mucocutaneous HSV)	5 mg/kg q8h	5 mg/kg q12h	5 mg/kg q24h	2.5 mg/kg q24h	2.5 mg/kg q24h	5 – 10 mg/kg q12h
Severe (e.g. CNS/ocular/disseminated HSV infections, Zoster)	10 mg/kg q8h	10 mg/kg q12h	10 mg/kg q24h	5 mg/kg q24h	5 mg/kg q24h	10 mg/kg q12h	
<b>Acyclovir (PO)</b> <sup>1,2,7</sup>		CrCl > 50	CrCl 25 – 50	CrCl < 25	CrCl < 10	IHD	CRRT
	<b>Prophylaxis</b>						
	BMT	800 mg BID	400 mg BID	200 mg BID	200 mg daily	200 mg daily	No data
	Hematology/Oncology	400 mg BID	400 mg BID	200 mg BID	200 mg daily	200 mg daily	No data
	<b>Treatment</b>						
	General (e.g. mucocutaneous HSV)	400 mg q8h Alt: 200 mg 5x daily		200 mg q8h	200 mg q12h	200 mg q12h	No data
Severe (e.g. CNS/ocular/disseminated HSV infections, Zoster)	800 mg q4h (or 5x daily) Consider valacyclovir for less frequent dosing		800 mg q8h	800 mg q12h	800 mg q12h	No data	
<b>Amikacin</b> <sup>1,2,5,8,9</sup> (Use adjusted BW in obese)  Refer to Aminoglycoside Dosing Guide		CrCl > 60	CrCl 40 – 60	CrCl 20 – 40	CrCl < 20		
	Conventional dosing	5 – 7.5 mg/kg q8h	5 – 7.5 mg/kg q12h	5 – 7.5 mg/kg q24h	5 mg/kg load, then by level	5 – 7.5 mg/kg post HD only consult pharmacist	
	High-dose extended-interval dosing	15 – 20 mg/kg q24h	15 mg/kg q36h	CrCl > 30: 15 mg/kg q48h CrCl < 30: Not recommended	alt: 7.5 mg/kg q48–72h		
<b>Timing of levels:</b> Draw trough 30 min prior to 4 <sup>th</sup> dose. Draw peak 30 min after infusion ends <b>Once daily dosing:</b> goal peak 35 – 60 mcg/mL; goal trough < 4 mcg/mL <b>Conventional dosing:</b> goal peak 25 – 35 mcg/mL for serious infections; 15 – 20 mcg/mL for UTI; goal trough < 4 – 8 mcg/mL							
<b>Amoxicillin (PO)</b> <sup>1,2</sup>	<b>Usual dose:</b> 500 – 1,000mg q8-12h or 875 mg q12h <b>CAP:</b> 1,000 mg q8h <b>H pylori:</b> 1,000 mg q12h <b>Procedural ppx:</b> 2,000 mg x 1	<b>CrCl 10–30:</b> 250 – 500 mg q12h	250 – 500 mg q24h	250 – 500 mg q24h	250 – 500 mg q24h; administer additional dose at the end of dialysis	No data	
	<b>Usual dose:</b> 250 – 500 mg q8h or 875 mg q12h	<b>CrCl &lt; 30:</b> Do not use 875 mg tablet <b>CrCl 10 – 30:</b> 250 – 500 mg q12h	250 – 500 mg q24h	250 – 500 mg q24h	250 – 500 mg q24h; administer additional dose at the end of dialysis	No data	
<b>Amphotericin B Liposomal</b> <sup>1,2</sup> (Consider adjusted BW in obese)	3 – 5 mg/kg/day	No change	No change	No change	No change	No change	
<b>Ampicillin (IV)</b> <sup>1-3</sup>	<b>Mild/uncomplicated:</b> 1 – 2 g q6h <b>Meningitis/endovascular/PJI:</b> 2 g q4h	<b>Mild/uncomplicated:</b> 1 g q6–8h <b>Meningitis/endovascular/PJI:</b> 2 g q6–12h	<b>Mild/uncomplicated:</b> 1 g q12h <b>Meningitis/endovascular/PJI:</b> 2 g q12–24h; or 1 g q8h	<b>Mild/uncomplicated:</b> 1 g q12h <b>Meningitis/endovascular/PJI:</b> 2 g q12–24h	<b>Mild/uncomplicated:</b> 1 g q12h <b>Meningitis/endovascular/PJI:</b> 2 g q12–24h	CVVH: 2 g q8–12H CVVHDF: 2 g q6–8h <b>Meningitis/endovascular/PJI:</b> 2 g q6h	
	1.5 – 3 g q6h	CrCl < 30: 1.5 – 3 g q12h	CrCl < 15: 1.5 – 3 g q24h	1.5 – 3 g q12–24h Dose daily, but after HD on HD days	3 g q6–8h		
<b>Azithromycin (IV/PO)</b> <sup>1,2</sup>	500 mg q24h	No change	No change	No change	No change	No change	
<b>Aztreonam</b> <sup>1-3,10</sup> Severe: pseudomonas, life-threatening infections	1 – 2 g q8h <b>Severe/Meningitis:</b> 2 g q6–8h	CrCl < 30: 1 g q8h <b>Severe/Meningitis:</b> 1 g q6–8h	500 mg q8h <b>Severe/Meningitis:</b> 1g q12h	1 g q24h <b>Severe/Meningitis:</b> 1 g q12h	2 g load, then 1 g q8h – or – 2 g q12h		
	70 mg x 1, then 50 mg q24h 70 mg q24h if on phenytoin, rifampin, other strong enzyme inducers <b>Endocarditis/Endovascular:</b> 150 mg q24h Dosage adjustments are not required for Child-Pugh B or C cirrhosis				No change	No change	
<b>Cefazolin</b> <sup>1-5,14</sup>	CrCl ≥ 35: <b>Mild/moderate:</b> 1 g q8h <b>Severe:</b> 2 g q8h	CrCl 10 – 34: <b>Mild/moderate:</b> 1 g q12h <b>Severe:</b> 2 g q12h	<b>Mild/moderate:</b> 1g q24h <b>Severe:</b> 2 g q24h	1 g q24h Dose daily, but after HD on HD days alt: 2g/2g/3g post-HD only	2 g q12h		

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD)	CRRT		
<b>Cefepime</b> <sup>1-3,5,15-17</sup>	Extended Infusion (4-hour infusion)				0.5 – 1 g q24h Dose daily, but after HD on HD days  alt: 2 g post-HD only	2 g load, then 1 g q8h (4-hour infusion)	
		CrCl > 60	CrCl 30 – 60	CrCl < 11-29			CrCl < 10
	General	1 g q8h or 2 g q12h	1 g q12h or 2 g q24h	1 g q24h			0.5 g q24h
	Severe/CNS/FN/CF exacerbation/confirmed Pseudomonal infection	2 g q8h	2 g q12h	1 g q12h	1 g q24h		
<b>Cefpodoxime (PO)</b> <sup>1,2</sup>	<u>Uncomplicated cystitis:</u> 100 mg q12h <u>CAP/bronchitis:</u> 200 mg q12h <u>Skin/skin structure:</u> 400 mg q12h	CrCl < 30: same dose q24h		Same dose, post-HD only	No data		
<b>Ceftaroline</b> <sup>1,2,18</sup> <b>(SHC Restriction)</b>		CrCl > 50	CrCl 30 – 50	CrCl 15 – 30	CrCl < 15	200 mg q8-12h  <u>Endocarditis/S.aureus bacteremia/ SDD:</u> 200 mg q8-12h administered over 2-hr	
	General	600 mg q12h	400 mg q12h	300 mg q12h	200 mg q12h		
	Endocarditis/S.aureus bacteremia, Susceptible-dose dependent (SDD)	600 mg q8h administered over 2-hr	400 mg q8h administered over 2-hr	300 mg q8h administered over 2-hr	200 mg q8h administered over 2-hr		
<b>Ceftazidime (IV)</b> <sup>1-3,19</sup>	<u>Usual dose:</u> 1 – 2 g q8h <u>Severe:</u> 2 g q8h	CrCl 30 – 50: 1 – 2 g q12h CrCl 16 – 30: 1 – 2 g q24h CrCl 6 – 15: 0.5 – 1 g q24h	CrCl < 5: 0.5 g q24h		0.5 – 1 g q24h Dose daily, but after HD on HD days  alt: 1 – 2 g q48-72h or 1 g post-HD only TIW	2 g load, then 1 g q8h – or – 2 g q12h	
<b>Ceftazidime/avibactam</b> <sup>1,2,20-23</sup> <b>(SHC Restriction)</b>	2.5 g q8h	CrCl 31 – 50: 1.25 g q8h CrCl 16 – 30: 0.94 g q12h CrCl 6 – 15: 0.94 g q24h	CrCl < 5: 0.94 g q48h		0.94 g q24-48h Dose daily, but after HD on HD days	1.25g q8h 2.5g q8h if MIC > 4 mcg/mL or deep-seated	
<b>Ceftolozane/tazobactam</b> <sup>1,2,24-26</sup> <b>(SHC Restriction)</b>		CrCl > 50	CrCl 30 – 50	CrCl 15 – 29	CrCl < 15	General: 750 mg load, then 150 mg q8h  HAP/VAP: 2.25 g load, then 450 mg q8h	
	General	1.5 g q8h	750 mg q8h	375 mg q8h	750 mg q8h, then 150 mg q8h		
	Hospital-acquired/Ventilator-associated pneumonia	3 g q8h	1.5 g q8h	750 mg q8h	2.25 g load, then 450 mg q8h		
<b>Ceftriaxone (IV)</b> <sup>1,2,27</sup>	1 – 2 g q24h <u>Endovascular/osteomyelitis/PJ:</u> 2 g q24h <u>Meningitis, E. faecalis endocarditis:</u> 2 g q12h		No change		No change	No change	
<b>Cephalexin (PO)</b> <sup>1,2,28</sup>	250 – 1000 mg q6h <u>Uncomplicated cystitis:</u> 500 mg q12h <u>Cellulitis/SSTI:</u> 500 mg q6h	CrCl 15 – 29: 250 mg q8-12h CrCl 5 – 14: 250 mg q24h		500 mg q24h Dose daily, but after HD on HD days	No data		
<b>Ciprofloxacin (IV/PO)</b> <sup>1-4,22,29</sup>		CrCl > 50	CrCl 30 – 50	CrCl < 30		200 – 400 mg IV q24h 250 – 500 mg PO q24h Dose daily, but after HD on HD days	
	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			
<b>Clindamycin</b> <sup>1,2</sup>	600 – 900 mg IV q8h 150 – 450 mg PO q6h	No change		No change	No change	No change	
<b>Colistin (IV)</b> <sup>1-3,30-32</sup> <b>(SHC Restriction)</b> (Dosage expressed in terms of colistin base activity [CBA]; Use ideal BW in obese)	<b>U.S. FDA Package Insert</b>					Loading Dose: 300 mg CBA x 1  Maintenance Dose: 220 mg CBA q12h  alt: 100 mg CBA q8h  Note: These are general recommendations (based on dialysis flow/ultrafiltration rates of 1-2 L/hr and minimal residual renal function. Refer to ESCMID/EUCAST guidelines)	
		CrCl > 80	CrCl 50 – 79	CrCl 30 – 49	CrCl < 30		
	Loading Dose	5 mg/kg x 1 (max dose: 300 mg)					
	Maintenance Dose	1.25 – 2.5 mg/kg q12h	1.25 – 1.9 mg/kg q12h	2.5 mg/kg q24h	1.5 mg/kg q36h		
	<b>Preferred Dosing for Critically Ill Patients (Consult ID Pharmacist)</b>						
		CrCl		Dosing Regimen			
	Loading Dose			300 mg CBA x 1			
	Maintenance Dose	> 90 mL/min		180 mg q12h			
		80 – 89 mL/min		170 mg q12h			
		70 – 79 mL/min		150 mg q12h			
60 – 69 mL/min		138 mg q12h					
50 – 59 mL/min		122 mg q12h					
40 – 49 mL/min		110 mg q12h					
30 – 39 mL/min		98 mg q12h					
20 – 29 mL/min		88 mg q12h					
10 – 19 mL/min		80 mg q12h					
5 – 9 mL/min		72 mg q12h					
0 mL/min		65 mg q12h					
Suggested loading dose and daily doses of colistimethate for desired target colistin C <sub>ss</sub> , avg of 2 mg/L (CID 2017:64. Nation et al)							
<b>Daptomycin</b> <sup>1,2,33-39</sup> <b>(SHC Restriction)</b> (Use adjusted BW in obese)	<u>Skin/Soft tissue:</u> 4 – 6 mg/kg q24h <u>Bacteremia/Endovascular:</u> 8 mg/kg q24h <u>E. faecium infections:</u> 10 – 12 mg/kg q24h; consult ID	CrCl < 30: Same dose q48h		Same dose q48h		6 – 8 mg/kg q24h  Doses > 8mg/kg q24h increases the risk of CPK elevations and myopathy. Caution, clinical judgement and frequent monitoring for higher doses	

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD)	CRRT																																																																					
<b>Doxycycline (IV/PO)</b> <sup>1,2</sup>	(Load: 200 mg x 1 for severe infections) 100 mg q12h	No change	No change	No change	No change																																																																					
<b>Ertapenem (IV/IM)</b> <sup>1,2,40-42</sup>	1 g q24h	<u>CrCl &lt;30</u> : 500 mg q24h	500 mg q24h	500 mg q24h Dose daily, but after HD on HD days alt: 500 - 1000 mg post-HD (low vs. high-flux HD, degree of renal failure, residual UOP)	1 g q24h																																																																					
<b>Ethambutol (PO)</b> <sup>1,5,43,44</sup> (Use lean BW if obese) (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 q24–36h	<u>CrCl &lt; 10</u> : 15 – 25 mg/kg q48h	15 – 25 mg/kg 3 times per week post-HD Administer after HD only	15 – 25 mg/kg q24–36h																																																													
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<b>Fidaxomicin (PO)</b> <sup>1,2</sup> (SHC Restriction)	200 mg q12h x 10 days	No change	No change	No change	No change																																																																					
<b>Fluconazole (IV/PO)</b> <sup>1-4,13,22,45</sup> Dose by indication. Load 800 mg for candidemia	200 – 400 mg q24h <u>C.glabrata</u> 800 mg q24h <u>Severe/CNS/endovascular infections:</u> up to 800 mg q24h	(50% of normal dose) q24h		Dose by indication; 200 – 800 mg post HD only	If usual dose is 200mg daily, use 400 daily. If usual dose is 400mg daily, use 800 mg q24h in 1-2 divided doses If usual dose is 800mg daily, use 1200 mg q24h (in 2 divided doses)																																																																					
<b>Foscarnet (IV)</b> <sup>1,2,46-48</sup> (Consider adjusted BW in obese)  Adj CrCl (mL/min/kg) $\left(\frac{140 - \text{age}}{\text{SCr} \times 72}\right) \times (0.85 \text{ if female})$	<table border="1"> <thead> <tr> <th>CrCl (mL/min/kg)</th> <th colspan="2">CMV induction</th> <th colspan="2">CMV maintenance</th> <th colspan="2">HSV</th> </tr> </thead> <tbody> <tr> <td>&gt; 1.4</td> <td>60 mg/kg q8h</td> <td>90 mg/kg q12h</td> <td>90 mg/kg q24h</td> <td>120 mg/kg q24h</td> <td>40 mg/kg q12h</td> <td>40 mg/kg q8h</td> </tr> <tr> <td>&gt; 1.0 – 1.4</td> <td>45 mg/kg q8h</td> <td>70 mg/kg q12h</td> <td>70 mg/kg q24h</td> <td>90 mg/kg q24h</td> <td>30 mg/kg q12h</td> <td>30 mg/kg q8h</td> </tr> <tr> <td>&gt; 0.8 – 1.0</td> <td>50 mg/kg q12h</td> <td>50 mg/kg q12h</td> <td>50 mg/kg q24h</td> <td>65 mg/kg q24h</td> <td>20 mg/kg q12h</td> <td>35 mg/kg q12h</td> </tr> <tr> <td>&gt; 0.6 – 0.8</td> <td>40 mg/kg q12h</td> <td>80 mg/kg q24h</td> <td>80 mg/kg q48h</td> <td>105 mg/kg q48h</td> <td>35 mg/kg q24h</td> <td>25 mg/kg q12h</td> </tr> <tr> <td>&gt; 0.5 – 0.6</td> <td>60 mg/kg q24h</td> <td>60 mg/kg q24h</td> <td>60 mg/kg q48h</td> <td>80 mg/kg q48h</td> <td>25 mg/kg q24h</td> <td>40 mg/kg q24h</td> </tr> <tr> <td>≥ 0.4 – 0.5</td> <td>50 mg/kg q24h</td> <td>50 mg/kg q24h</td> <td>50 mg/kg q48h</td> <td>65 mg/kg q48h</td> <td>20 mg/kg q24h</td> <td>35 mg/kg q24h</td> </tr> <tr> <td>&lt; 0.4</td> <td colspan="2">Not recommended</td> <td colspan="2">Not recommended</td> <td colspan="2">Not recommended</td> </tr> <tr> <td>IHD</td> <td colspan="2">60 – 90 mg/kg loading dose, then 45 – 60 mg/kg/dose post-HD only</td> <td colspan="2">No data</td> <td colspan="2">No data</td> </tr> <tr> <td>CRRT</td> <td colspan="5">No data</td> </tr> </tbody> </table>					CrCl (mL/min/kg)	CMV induction		CMV maintenance		HSV		> 1.4	60 mg/kg q8h	90 mg/kg q12h	90 mg/kg q24h	120 mg/kg q24h	40 mg/kg q12h	40 mg/kg q8h	> 1.0 – 1.4	45 mg/kg q8h	70 mg/kg q12h	70 mg/kg q24h	90 mg/kg q24h	30 mg/kg q12h	30 mg/kg q8h	> 0.8 – 1.0	50 mg/kg q12h	50 mg/kg q12h	50 mg/kg q24h	65 mg/kg q24h	20 mg/kg q12h	35 mg/kg q12h	> 0.6 – 0.8	40 mg/kg q12h	80 mg/kg q24h	80 mg/kg q48h	105 mg/kg q48h	35 mg/kg q24h	25 mg/kg q12h	> 0.5 – 0.6	60 mg/kg q24h	60 mg/kg q24h	60 mg/kg q48h	80 mg/kg q48h	25 mg/kg q24h	40 mg/kg q24h	≥ 0.4 – 0.5	50 mg/kg q24h	50 mg/kg q24h	50 mg/kg q48h	65 mg/kg q48h	20 mg/kg q24h	35 mg/kg q24h	< 0.4	Not recommended		Not recommended		Not recommended		IHD	60 – 90 mg/kg loading dose, then 45 – 60 mg/kg/dose post-HD only		No data		No data		CRRT	No data				
CrCl (mL/min/kg)	CMV induction		CMV maintenance		HSV																																																																					
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<b>Ganciclovir (IV)</b> <sup>1,2</sup> (Consider adjusted BW in obese)	<table border="1"> <thead> <tr> <th>CMV</th> <th>CrCl &gt;70*</th> <th>CrCl &gt;50</th> <th>CrCl &gt;25</th> <th>CrCl &gt;10</th> <th>CrCl &lt;10</th> <th>IHD</th> <th>CRRT</th> </tr> </thead> <tbody> <tr> <td>Induction (I)</td> <td>5 mg/kg q12h</td> <td>2.5 mg/kg q12h</td> <td>2.5 mg/kg q24h</td> <td>1.25 mg/kg q24h</td> <td>1.25 mg/kg 3x/week</td> <td>! 1.25 mg/kg post HD only M: 0.625 mg/kg post HD only</td> <td>! 2.5 mg/kg q12–24h M: 1.25 – 2.5 mg/kg q24h</td> </tr> <tr> <td>Maintenance (M)</td> <td>5 mg/kg q24h</td> <td>2.5 mg/kg q24h</td> <td>1.25 mg/kg q24h</td> <td>0.625 mg/kg q24h</td> <td>0.625 mg/kg 3x/week</td> <td></td> <td></td> </tr> </tbody> </table> <p>*Manufacturer's CrCl cutoffs. Please refer to BMT protocols if applicable</p>					CMV	CrCl >70*	CrCl >50	CrCl >25	CrCl >10	CrCl <10	IHD	CRRT	Induction (I)	5 mg/kg q12h	2.5 mg/kg q12h	2.5 mg/kg q24h	1.25 mg/kg q24h	1.25 mg/kg 3x/week	! 1.25 mg/kg post HD only M: 0.625 mg/kg post HD only	! 2.5 mg/kg q12–24h M: 1.25 – 2.5 mg/kg q24h	Maintenance (M)	5 mg/kg q24h	2.5 mg/kg q24h	1.25 mg/kg q24h	0.625 mg/kg q24h	0.625 mg/kg 3x/week																																															
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Induction (I)	5 mg/kg q12h	2.5 mg/kg q12h	2.5 mg/kg q24h	1.25 mg/kg q24h	1.25 mg/kg 3x/week	! 1.25 mg/kg post HD only M: 0.625 mg/kg post HD only	! 2.5 mg/kg q12–24h M: 1.25 – 2.5 mg/kg q24h																																																																			
Maintenance (M)	5 mg/kg q24h	2.5 mg/kg q24h	1.25 mg/kg q24h	0.625 mg/kg q24h	0.625 mg/kg 3x/week																																																																					
<b>Gentamicin</b> <sup>1,3,49</sup> (Use adjusted BW in obese)  See appendix for complete guidelines	<table border="1"> <thead> <tr> <th></th> <th>CrCl &gt; 60</th> <th>CrCl 40 – 59</th> <th>CrCl 20 – 39</th> <th>CrCl &lt; 20</th> <th>IHD</th> <th>CRRT</th> </tr> </thead> <tbody> <tr> <td><b>Gram negative</b></td> <td>1.7 mg/kg q8h or 5 – 7 mg/kg q24h (high-dose extended-interval)</td> <td>1.7 mg/kg q12h or 5 – 7 mg/kg q36h (high-dose extended-interval)</td> <td>1.7 mg/kg q24h or CrCl &gt; 30: 5 – 7 mg/kg q48h CrCl &lt; 30: Not recommended (high-dose extended-interval)</td> <td>2 mg/kg loading dose, then per level</td> <td>2 mg/kg loading dose, then 1.5 mg/kg post HD</td> <td>1.5 – 2.5 mg/kg q24–48h</td> </tr> <tr> <td><b>Gram positive synergy</b></td> <td>1 mg/kg q8h**</td> <td>1 mg/kg q12h</td> <td>1 mg/kg q24h</td> <td>1 mg/kg load, then by level</td> <td>1 mg/kg q48–72h; consider redosing when level &lt; 1 mcg/L</td> <td>1 mg/kg q24h, then per level</td> </tr> </tbody> </table> <p><b>Goal levels:</b> Gram-negative infections: Goal peak for traditional dosing 4 – 8 mcg/mL; goal trough &lt; 1 – 2 mcg/mL Gram-positive synergy: Goal peak 3 – 4 mcg/mL; goal trough &lt; 1 mcg/mL <b>Timing of levels:</b> Draw peak 30 minutes after completion of 3<sup>rd</sup> dose. Draw trough 30 minutes prior to 4<sup>th</sup> dose (For CrCl &lt; 20 mL/min, may check levels sooner than 3<sup>rd</sup>/4<sup>th</sup> dose) For 7 mg/kg once-daily dosing, draw a single random level 8 – 12 hours after dose administration. Adjust based on Hartford nomogram For HD, draw trough pre-HD (alternative: draw trough level 4-hr post-HD); and peak 30 minutes after end of each infusion ** Streptococci, <i>Streptococcus gallolyticus (bovis)</i>, <i>Streptococcus viridans</i> endocarditis: optional dosing 3 mg/kg q24h for CrCl &gt; 60 mL/min ** Staphylococci; Enterococcus spp (strains susceptible to PCN and gentamicin) endocarditis: optional dosing 3 mg/kg in 2 or 3 equally divided doses</p>						CrCl > 60	CrCl 40 – 59	CrCl 20 – 39	CrCl < 20	IHD	CRRT	<b>Gram negative</b>	1.7 mg/kg q8h or 5 – 7 mg/kg q24h (high-dose extended-interval)	1.7 mg/kg q12h or 5 – 7 mg/kg q36h (high-dose extended-interval)	1.7 mg/kg q24h or CrCl > 30: 5 – 7 mg/kg q48h CrCl < 30: Not recommended (high-dose extended-interval)	2 mg/kg loading dose, then per level	2 mg/kg loading dose, then 1.5 mg/kg post HD	1.5 – 2.5 mg/kg q24–48h	<b>Gram positive synergy</b>	1 mg/kg q8h**	1 mg/kg q12h	1 mg/kg q24h	1 mg/kg load, then by level	1 mg/kg q48–72h; consider redosing when level < 1 mcg/L	1 mg/kg q24h, then per level																																																
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<b>Imipenem/Cilastatin (IV)</b> <sup>1</sup> (SHC Restriction)	<table border="1"> <thead> <tr> <th></th> <th>CrCL &gt;60</th> <th>CrCL 30 – 59</th> <th>CrCL 15 – 29</th> <th>CrCL &lt; 10</th> <th>IHD</th> <th>CRRT</th> </tr> </thead> <tbody> <tr> <td>General</td> <td>500 mg q6H or 1g q8h</td> <td>500 mg q8h</td> <td>500 mg q12h</td> <td>Not recommended unless dialysis initiated within 48-hrs</td> <td>250 – 500 mg q12h</td> <td>1g load, then 500 mg q6h</td> </tr> <tr> <td>NTM</td> <td>1,000 mg q12H</td> <td>750 mg q12H</td> <td>500 mg q12H</td> <td></td> <td></td> <td></td> </tr> </tbody> </table>						CrCL >60	CrCL 30 – 59	CrCL 15 – 29	CrCL < 10	IHD	CRRT	General	500 mg q6H or 1g q8h	500 mg q8h	500 mg q12h	Not recommended unless dialysis initiated within 48-hrs	250 – 500 mg q12h	1g load, then 500 mg q6h	NTM	1,000 mg q12H	750 mg q12H	500 mg q12H																																																			
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<b>Isavuconazole (IV/PO)</b> <sup>1,2</sup>	<u>Initial:</u> 372 mg q8h x 6 doses <u>Maintenance:</u> 372 mg q24h	No change	No change	No change	No change																																																																					
<b>Isoniazid (PO)</b> <sup>1,2,43,44</sup>	300 mg q24h (5 mg/kg/day)	No change	No change	No change	No change																																																																					
<b>Levofloxacin (IV/PO)</b> <sup>1-4</sup>	<table border="1"> <thead> <tr> <th></th> <th>CrCl ≥ 50</th> <th>CrCl 20 – 49</th> <th>CrCl &lt; 20</th> <th>IHD</th> <th>CRRT</th> </tr> </thead> <tbody> <tr> <td>General</td> <td>250 – 500 mg q24h</td> <td>250 mg q24h - or - 500 mg q48h</td> <td>500 mg x1, then 250 mg q48h</td> <td>See CrCl &lt; 20 ml/min Dose q48h, but after HD on HD days</td> <td>750 mg load, then 250 – 500 mg q24h</td> </tr> <tr> <td>Severe/PNA/Pseudomonas/Stenotrophomonas:</td> <td>750 mg q24h</td> <td>750 mg q48h</td> <td>750 mg x1, then 500 mg q48h</td> <td></td> <td></td> </tr> </tbody> </table>						CrCl ≥ 50	CrCl 20 – 49	CrCl < 20	IHD	CRRT	General	250 – 500 mg q24h	250 mg q24h - or - 500 mg q48h	500 mg x1, then 250 mg q48h	See CrCl < 20 ml/min Dose q48h, but after HD on HD days	750 mg load, then 250 – 500 mg q24h	Severe/PNA/Pseudomonas/Stenotrophomonas:	750 mg q24h	750 mg q48h	750 mg x1, then 500 mg q48h																																																					
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Severe/PNA/Pseudomonas/Stenotrophomonas:	750 mg q24h	750 mg q48h	750 mg x1, then 500 mg q48h																																																																							
<b>Linezolid (IV/PO)</b> <sup>1,2</sup> (SHC Restriction)	600 mg q12h	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)	CRRT		
<b>Meropenem</b> <sup>1-4,50</sup> (SHC Restriction) 3-hr extended infusion		CrCl > 50	CrCl 26 – 50	CrCl 10 – 25	CrCl < 10	500 mg q24h <u>CF/CNS:</u> 1 g q24h <i>Dose daily, but after HD on HD days</i>	1 g q8h <u>CF/CNS:</u> 2 g q12h
	Usual dose (FN, PNA, Pseudomonas)	1 g q8h	1 g q12h	0.5 g q12h	0.5 g q24h		
	CF/Meningitis	2 g q8h	2 g q12h	1 g q12h	1 g q24h		
<b>Metronidazole (IV/PO)</b> <sup>1,2</sup>	500 mg q6–8h	No change Severe hepatic impairment: can consider 500 mg q12h		500 mg q8h	500 mg q6–8h		
<b>Moxifloxacin (IV/PO)</b> <sup>1,2</sup>	400 mg IV/PO q24h	No change	No change	No change	No change		
<b>Nafcillin</b> <sup>1,2</sup>	2 g q4h Mild infections: 1 g q4h	No change for renal impairment. <u>Hepatic Impairment:</u> No specific dose adjustment provided by manufacturer. Dosage adjustment may be necessary in the setting of concomitant renal impairment; nafcillin primarily undergoes hepatic metabolism.					
<b>Osetamivir (PO)</b> <sup>1,2,51</sup>		CrCl ≥ 60	CrCl 30 – 60	CrCl 10 – 30	Prophylaxis: 30 mg x 1, then 30 mg after every other HD session <u>Treatment:</u> 30 mg x 1, then 30 mg post-HD only	Prophylaxis: 75 mg q24h <u>Treatment:</u> 75 mg q12h	
	Prophylaxis	75 mg q24h	30 mg q24h	30 mg q48h			
	Treatment	75 mg q12h	30 mg q12h	30 mg q24h			
<b>Penicillin G (IV)</b> <sup>1-3,5</sup>	2 – 4 mu q4h <u>Dose range:</u> 12 – 24 million units/day continuous infusion or in divided doses every 4 to 6 hours	2 – 3 mu q4h	1 – 2 mu q6h	Mild: 0.5 – 1 mu q4–6h; or 1 – 2 mu q8–12h Severe: 2 mu q4–6h; or 4 mu q8–12h	4 mu q4–6h		
<b>Piperacillin/tazobactam</b> <sup>1-4,52,53</sup>		CrCl > 40	CrCl 20 – 40	CrCl < 20	General: 2.25 g q12h  Severe infections: 3.375 g q12h over 4-hr <u>alt:</u> 2.25 g q8h	3.375 g q6h  Extended infusion: 3.375 – 4.5 g q8h over 4-hr	
	<u>Intermittent Dosing</u>						
	General	3.375 g q6h	2.25 g q6h	2.25 g q8h			
	Severe/sepsis/CF/nosocomial PNA	4.5 g q6h	3.375 g q6h	2.25 g q6h			
<u>Extended-Infusion Dosing (4-hr infusion)</u>							
General, CF Pseudomonas, nosocomial PNA:	<u>Extended infusion for CrCl &gt; 20:</u> 3.375 – 4.5 g q8h over 4h*		3.375 g q12h over 4h				
*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 q8h (infused over 4 hours) or q6h.							
<b>Polymyxin B</b> <sup>1,2,54,55</sup> (SHC Restriction) (Use adjusted BW if obese)	Dosing presented as units (10,000 units = 1 mg) 20,000 – 25,000 units/kg load x 1, then 12,500 – 15,000 units/kg q12h (maximum: 25,000 units/kg/day)			No data	No change		
<b>Posaconazole (PO/IV)</b> <sup>1,2</sup> (SHC Restriction [IV])		Oral Suspension		Delayed-release tablet / Intravenous solution			
	Prophylaxis	200 mg q8h		300 mg q12h x 1 day, then 300 mg q24h			
	Treatment	<u>Usual dose:</u> 200 mg q6–8h or 400 mg q12h					
No renal adjustment							
<ul style="list-style-type: none"> <li>Delayed-release tablet and oral suspension are not interchangeable</li> <li>Posaconazole levels shown to have great degree of interpatient variability. Consider drawing a trough 4 – 7 days after initiating dose.</li> <li>Please refer to <a href="#">Antifungal TDM Guide</a></li> </ul>							
<b>Pyrazinamide (PO)</b> <sup>1,2,43,44</sup> (Use lean BW if obese) (See footnote for lean BW equation)	<u>Usual Dose:</u> 25 mg/kg q24h (max dose: 2,000 mg/day)		CrCl < 30: 25 mg/kg 3 times per week		25 mg/kg 3 times per week Administer after HD only	No data	
	Lean body weight	Dose					
	40 – 55 kg	1,000 mg					
	56 – 75 kg	1,500 mg					
	76 – 90 kg	2,000 mg					
<b>Rifampin (IV/PO)</b> <sup>1,2,43,44,56–58</sup> Capsule size: 150mg, 300mg	TB: 600 mg q24h (≤ 45 kg: 10 mg/kg q24h) Endocarditis: 300 mg q8h PJI: 300 – 450 mg q12h Vertebral Osteomyelitis: 600 mg q24h		No change	No change	No change		
<b>Tedizolid (IV/PO)</b> <sup>1,2,59</sup> (SHC Restriction)	200 mg q24h	No change	No change	No change	No change		
<b>Tobramycin</b> <sup>1,2,49</sup>	Refer to Gentamicin for dosing. See appendix for complete guidelines.						
<b>Trimethoprim (TMP)/ Sulfamethoxazole (IV/PO)</b> <sup>1,2,4,60</sup>  (Dose by adjusted BW in obese) SS = 80 mg TMP = 10 ml po soln DS = 160 mg TMP = 20ml po soln	<u>Uncomplicated cystitis:</u> 1 DS tab PO BID		CrCl 15 – 30: Administer 50% of recommended dose	CrCl < 15: Use is not recommended, but if needed for PCP: 5 – 8 mg/kg TMP q24h	2.5 – 5 mg/kg TMP q24h  PCP: 5 – 8 mg/kg TMP q24h <i>Dose daily, but after HD on HD days</i>  <u>alt:</u> 5 – 15 mg/kg TMP post-HD only		
	SSTI: 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 (2DS 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)							
					<u>PCP/ Stenotrophomonas:</u> 15 mg/kg/day TMP divided q8–12h		

Drug	CrCl > 50 mL/min	CrCl 10 – 50 mL/min	CrCl < 10 mL/min	Intermittent Hemodialysis (IHD)	CRRT			
<b>Valacyclovir (PO)</b> <sup>1,2</sup>  Please refer to transplant protocols if applicable	CrCl > 30		CrCl 10 – 30	< 10	500 mg q24h Dose daily, but after HD on HD days	No data		
	VZV	CrCl >50: 1 g q8h CrCl 30-50: 1 g q12h	1 g q24h	500 mg q24h				
	Genital herpes	Initial episode: 1 g q12h Recurrent episode: 500 mg q12h	Initial episode: 1 g q24h Recurrent: 500 mg q24h	Initial/recurrent episode: 500 mg q24h				
Herpes labialis	CrCl >50: 2 g q12h x 2 doses CrCl 30 – 50: 1 g q12h x 2 doses	500 mg q12h x 2 doses	500 mg x 1 dose					
<b>Valganciclovir (PO)</b> <sup>1,2</sup>  Please refer to transplant protocols if applicable	CrCl > 60		CrCl 40 – 59	CrCl 25 – 39	CrCl 10 – 24	CrCl < 10; IHD	CRRT	
	Induction (14-21 days)		900 mg q12h	450 mg q12h	450 mg q24h	450 mg q48h	200 mg 3x/week after HD only	No data
	Maintenance/ prophylaxis		900 mg q24h	450 mg q24h	450 mg q48h	450 mg twice/week	100 mg 3x/week after HD only	No data
<b>Vancomycin (IV)</b> <sup>1,2,61,62</sup>  (Use actual body weight; Excel AUC dose calculator encouraged, especially for obese.	<b>Consider loading dose of 20 – 35 mg/kg (max 3 g) for severe infections</b>							
	CrCl (mL/min)	Dose & Frequency		Total daily dose range				
	> 90	15 mg/kg q8h – 12h; use AUC calculator		30 – 45 mg/kg/day				
	51 – 89	10 – 20 mg/kg q12h; use AUC calculator		20 – 40 mg/kg/day				
	30 – 50	10 – 15 mg/kg q12h to 20 mg/kg q24h; use AUC calculator		20 – 30 mg/kg/day				
10 – 29	10 – 15 mg/kg q24h to 15 mg/kg q48h		7.5 – 15 mg/kg/day					
< 10 or AKI	15 mg/kg x 1, then dose by level		N/A					
<b>Goal AUC/MIC ratio: 400-600 (most indications)</b> <b>Refer to Vancomycin Guide for complete guidelines</b>								
<b>Vancomycin PO</b> <sup>1,2,63</sup>	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		
	<b>IV→PO conversion 1:1 (round to nearest tablet size- available in 200 mg and 50 mg tablets)</b> Caution with IV: accumulation of IV vehicle cyclodextrin occurs. Consider PO if CrCl < 50 mL/min unless benefits justify risks of IV use. Levels shown to have great degree of interpatient variability. Consider drawing a trough 4 – 7 days after new dose. Please refer to <a href="#">Antifungal TDM Guide</a>							
<b>Voriconazole (IV/PO)</b> <sup>1,2,64,65</sup>  (Dose by adjusted BW in obese)	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							
	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. Levels shown to have great degree of interpatient variability. Consider drawing a trough 4 – 7 days after new dose. Please refer to <a href="#">Antifungal TDM Guide</a>							

**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 (Weight<sup>2</sup>/(100 x Height(m)<sup>2</sup>)<sup>2</sup>)  
 LBW (women) = (1.07 x Weight(kg)) - 148 x (Weight<sup>2</sup>/(100 x Height(m)<sup>2</sup>)<sup>2</sup>)  
 LBW online calculator: <http://www.empr.com/medical-calculators/lean-body-weight-calculator/article/170219/>

**References:**

1. Lexicomp Online. Accessed April 9, 2017. <http://online.lexi.com>
2. MICROMEDEX®. Accessed April 9, 2017. <http://www.micromedexsolutions.com.laneproxy.stanford.edu/micromedex2/librarian>
3. 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
4. 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
5. Aronoff G, Bennett W, Berns J, et al. *Drug Prescribing in Renal Failure.* 5th ed. American College of Physicians; 2007.
6. 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
7. Tomblin 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
8. 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
9. Taccone FS, Backer D de, Laterre P-F, 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. 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
11. 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
12. Roger C, Wallis SC, Muller L, et al. Caspofungin Population Pharmacokinetics in Critically Ill Patients Undergoing Continuous Venovenous Hemofiltration or Haemodiafiltration. *Clin Pharmacokinet.* 2017;56(9):1057-1068. doi:10.1007/s40262-016-0495-z
13. 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
14. Strykowski 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
15. 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
16. Bauer KA, West JE, O'Brien JM, Goff DA. Extended-infusion cefepime reduces mortality in patients with Pseudomonas aeruginosa infections. *Antimicrob Agents Chemother.* 2013;57. doi:10.1128/AAC.02365-12
17. 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
18. Vaidialac 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
19. 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
20. 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
21. Soukup P, Faust AC, Edruganti 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. *Pharmacoher J Hum Pharmacol Drug Ther.* 2019;39(12):1216-1222. doi:10.1002/phar.2338
22. 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



23. 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
24. 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. *Pharmacother J Hum Pharmacol Drug Ther.* 2016;36(5):e30-e33. doi:10.1002/phar.1744
25. 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
26. 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
27. Laville M, Mercatello A, Frenay J, et al. Pharmacokinetics of ceftriaxone in hemodialysis. *Pathol Biol (Paris).* 1987;35(5 Pt 2):719-723.
28. 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
29. 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
30. Nation RL, Garonzik SM, Thamlikitkul V, et al. Dosing Guidance for Intravenous Colistin in Critically Ill Patients. *Clin Infect Dis.* 2017;64(5):565-571. doi:10.1093/cid/ciw839
31. Plachouras D, Karvanen M, Friberg LE, et al. Population pharmacokinetic analysis of colistin methanesulfonate and colistin after intravenous administration in critically ill patients with infections caused by gram-negative bacteria. *Antimicrob Agents Chemother.* 2009;53(8):3430-3436. doi:10.1128/AAC.01361-08
32. Dalfino L, Puntillo F, Mosca A, et al. High-dose, extended-interval colistin administration in critically ill patients: is this the right dosing strategy? A preliminary study. *Clin Infect Dis Off Publ Infect Dis Soc Am.* 2012;54(12):1720-1726. doi:10.1093/cid/cis286
33. 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
34. 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
35. Haselden M, Leach M, Bohm N, Bohm N. Daptomycin dosing strategies in patients receiving thrice-weekly intermittent hemodialysis. *Ann Pharmacother.* 2013;47(10):1342-1347. doi:10.1177/1060028013503110
36. 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
37. 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
38. 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
39. Xu X, Khadzhyrov 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
40. 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
41. Hsaiky LM, Salintri 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
42. Ueng Y-F, Wang H-J, Wu S-C, Ng Y-Y. A Thrice-Weekly Ertapenem Regimen Is Practical for Hemodialysis Patients. *Antimicrob Agents Chemother.* 2019;63(12). doi:10.1128/AAC.01427-19
43. Drug-Resistant Tuberculosis: A Survival Guide for Clinicians, 3rd edition | Curry International Tuberculosis Center. Accessed April 10, 2017. <http://www.currytcenter.ucsf.edu/products/cover-pages/drug-resistant-tuberculosis-survival-guide-clinicians-3rd-edition>
44. 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
45. Muijlwijk 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
46. Aweeka FT, Jacobson MA, Martin-Munley S, et al. Effect of renal disease and hemodialysis on foscarnet pharmacokinetics and dosing recommendations. *J Acquir Immune Defic Syndr Hum Retrovirology Off Publ Int Retrovirology Assoc.* 1999;20(4):350-357.
47. 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;21(5):384-395.
48. MacGregor RR, Graziani AL, Weiss R, Grunwald JE, Gambertoglio JG. Successful foscarnet 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.
49. 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.
50. 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
51. Robson R, Buttmore 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
52. 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
53. 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
54. 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
55. 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
56. 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
57. Osmon DR, Barbari 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
58. Barbari 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
59. 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
60. Nahata MC. Dosage regimens of trimethoprim-sulfamethoxazole (TPM/SMX) in patients with renal dysfunction. *Ann Pharmacother.* 1995;29(12):1300.
61. 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
62. 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
63. 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
64. 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
65. Davies-Vorbrodt S, Ito JI, Tegtmeyer 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

**A. Original Author/Date**

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

Lina Meng, PharmD, BCPS, BCCCP; 11/2010, 03/2011, 03/2017, 08/2019, 10/2019, 05/2020, 09/2020

Marisa Holubar, MD; 03/2017

Stan Deresinski, MD; 03/2017

Will Alegria, PharmD; 08/2019, 01/2020, 05/2020, 09/2020

David Ha, PharmD; 01/2020, 05/2020, 09/2020

**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

**Pharmacy & Therapeutics Committee** 04/2007, 02/2009, 04/2010, 05/2011, 08/2012, 09/2012, 08/2013, 02/2014, 04/2017, 11/2019; 02/2020, 10/2020

