

LTC Treatment Guidance

Empiric Recommendations for Common Infections for Adult LTC Patients

- 1. Acute Bacterial Sinusitis
- 2. Acute Bacterial Pharyngitis
- 3. Non-purulent Cellulitis
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 Tube Considerations

Disclaimer: This guidance is intended for educational purposes only. We do not provide direct medical care

treatment planning, or medical treatment services to individuals. The information provided through the service is not a replacement for local expertise. Information is offered as clinical decision support, is advisory in nature and is not intended to replace local healthcare decision-making or provision. Final clinical decisions are the sole responsibility of the healthcare provider.

Acute Bacterial Rhinosinusitis (S. pneumoniae, H. influenzae, M. catarrhalis)

Mild: Afebrile, no purulent nasal discharge, no facial pain longer than 3-4 days. Severe: Fever, purulent nasal discharge, facial pain longer than 3-4 consecutive days, or worsening symptoms after 5-6 days ("double sickening")

		Treatment	Renal Dose Adjustment	Feeding Tube Compatible	Duration
No antimicrobial treatment warranted empirically . "Watchful waiting" with symptomatic management is recommended. Most sinusitis is viral and will spontaneously improve. <i>If no improvement after 10 Days of symptomatic treatment/supportive care:</i>					
Mild	Preferred	Amoxicillin/Clavulanate 2000/125 mg BID*	Yes	No	5 Days
IVIIIU	Alternative	Amoxicillin/Clavulanate 875/125 mg BID OR Doxycycline 100 mg BID OR	Yes No	Yes Yes	5 Days
		Cefpodoxime 200 mg BID	Yes	Yes	3 Days
	Treat empirically wi	th antibiotics, "watchful waiting" NOT indicated			
Severe	Preferred	Amoxicillin/Clavulanate 2000/125 mg BID*	Yes	No	
		Amoxicillin/Clavulanate 875/125 mg BID OR	Yes	Yes	5-7 Days
	Alternative	Doxycycline 100 mg BID OR	No	Yes	0 1 20.,0
		Cefpodoxime 200 mg BID	Yes	Yes	

^{*}High-dose amoxicillin-clavulanate is preferred in those with a risk of a poor outcome include patients ≥65 years, recently hospitalized, antibiotic use in 30 days, immunocompromised or in areas with >10% resistance to *S. pneumoniae*. When amoxicillin-clavulanate (Augmentin XR™ 1000/62.5 mg) is not available or accessible, amoxicillin/clavulanate 875/125 mg BID is preferred over alternative options.

Acute Bacterial Pharyngitis (Group A Strep, Streptococcus spp.)

Treatment is not recommended for patients with viral pharyngitis

	Antibiotic	Renal Dose Adjustment	Feeding Tube Compatible	Duration
Preferred	Penicillin V 500 mg BID OR Amoxicillin 500 mg BID	Yes	Yes	10 Days
Alternative	Cephalexin 500 mg BID OR Clindamycin 300 mg TID	Yes No	Yes Yes	10 Days

Non-purulent Cellulitis (Streptococcus pyogenes, also known as Group A Strep)					
	Antibiotic	Renal Dose Adjustment	Feeding Tube Compatible	Duration	
Preferred	Cephalexin 1,000 mg TID OR Amoxicillin 875 mg BID	Yes Yes	Yes Yes	5 Days	
Alternative	Sulfamethoxazole/Trimethoprim 1600/320 mg (2 DS) BID OR Dicloxacillin 500 mg four times daily	Yes Yes	Yes Yes	5 Days	
Alternative, IV	Cefazolin 1 g IV Q8H*	Yes	N/A	5 Days	
*For patients ≥120 kg consider use of 2 g cefazolin					

Abscess or Purulent Cellulitis (Staphylococcus aureus, including MRSA and MSSA) Prioritize incision and drainage for primary treatment of abscess. Antimicrobials not always recommended if small and drained **Feeding Tube Renal Dose Antibiotic** Compatible Duration **Adjustment** Preferred Doxycycline 100 mg BID 7 Days No Yes Sulfamethoxazole/Trimethoprim 1600/320 mg (2 DS) BID OR Yes Yes Alternative 7 Days Clindamycin 450 mg TID No Yes Vancomycin (Follow facility specific dosing and monitoring) OR N/A Yes Alternative, IV 7 Days Clindamycin 600 mg IV Q8H N/A No

Mild Diabetic Foot Wound

Diabetic or non-healing wounds do not always require treatment with antibiotics. Diagnosis of soft tissue diabetes-related infections should be based on the presence of local or systemic signs/symptoms of inflammation. Local wound findings may include: penetrating wound, rapidly progressing cellulitis or signs of induration, crepitus, bullae, discoloration, necrosis, gangrene, ecchymosis, petechiae or new pain

		Antibiotic	Renal Dose Adjustment	Feeding Tube Compatible	Duration
	Low risk MRSA	Cephalexin 1,000 mg TID/four times daily	Yes	Yes	7-14 Days
Preferred	High risk MRSA	Sulfamethoxazole/Trimethoprim 1600/320 mg (2 DS) BID OR Doxycycline 100 mg BID	Yes No	Yes	7-14 Days
Alternative,	Low risk MRSA	Cefazolin 1,000 mg IV Q8H	Yes	N/A	7-14 Days
IV	High risk MRSA	Vancomycin IV (Follow facility specific dosing and monitoring)	Yes	N/A	7-14 Days

Evaluate severity and rule out deeper involvement (i.e. bone/joint). Longer treatment and surgical management may be necessary for resolution

Uncomplicated UTI (uUTI) (E. coli, Klebsiella spp., Proteus spp.)

Uncomplicated: men and women with local bladder signs/symptoms (dysuria, urgency, frequency or suprapubic pain). uUTI patients may have underlying urologic abnormalities, immunocompromising factors, and/or have diabetes. Recurrent UTI considered uncomplicated

	Antibiotic	Renal Dose Adjustment	Feeding Tube Compatible	Duration
Preferred	Nitrofurantoin 100 mg BID (only if CrCl >30 mL/min)	No	Yes	5 Days
Alternative	Cephalexin 500 mg BID OR Sulfamethoxazole/Trimethoprim 800/160 mg (1 DS) BID OR Fosfomycin trometamol 3,000 mg x1 dose	Yes Yes No	Yes Yes Yes	5 Days 3 Days 1 Day
Alternative, IV	Ceftriaxone 1 g IV once daily Gentamicin 5 mg/kg IV or IM x1 dose	No No	N/A N/A	3-5 Days 1 Day

*For gentamicin, use adjusted body weight in patients with total body weight >20% than ideal body weight. Single doses of gentamicin generally do not require renal dose adjustments

AdjBW = IBW + [0.4 x (TBW - IBW)]

IBW (male) = 50 kg + 2.3 kg for each inch over 5 feet IBW (female) = 45 kg + 2.3 kg for each inch over 5 feet

Complicated UTI (cUTI) (E. coli, Klebsiella spp., Proteus spp.)

Complicated: UTI with signs/symptoms of infection extending beyond the bladder such as fever, flank pain, and/or signs of systemic illness (chills, hemodynamic instability, and/or significant fatigue). Pyelonephritis, febrile UTI, catheter-associated UTI, and bacteremia are included

	Antibiotic	Renal Dose Adjustment	Feeding Tube Compatible	Duration
Preferred	Gentamicin 5 mg/kg IV/IM once OR Ceftriaxone 1 g IV/IM x1 dose <i>followed by</i> Sulfamethoxazole/Trimethoprim 800/160 mg (1 DS) BID	No Yes	N/A Yes	7 Days
	Gentamicin 5 mg/kg IV/IM once OR Ceftriaxone 1 g IV/IM x1 dose <i>followed by</i>	No	N/A	
Alternative	Amoxicillin/Clavulanate 875/125 mg BID OR	Yes	Yes	7 Days total
	Cefpodoxime 200 mg BID OR Ciprofloxacin 750 mg BID	Yes Yes	Yes Avoid*	
Alternative, IV	Ceftriaxone 1 g IV once daily	No	N/A	7 Days

Note: Catheterized patients will be colonized with bacteria within 24 hours of placement. Culturing/urinalysis ought to be considered only in patients with suprapubic tenderness, fever or signs of systemic infection. Changes in mental status are more suggestive of polypharmacy. Single doses of gentamicin generally do not require renal dose adjustments.*Poor absorption with quinolones observed when given via feeding tubes. See above uUTI table for recommendations on gentamicin dosing

Community Acquired Bacterial Pneumonia (S. pneumoniae, H. influenzae, M. catarrhalis)

	Antibiotic	Renal Dose Adjustment	Feeding Tube Compatible	Duration
Preferred	Amoxicillin 1,000 mg TID	Yes	Yes	5 Days
Alternative	Doxycycline 100 mg BID OR Cefpodoxime 200 mg BID	No Yes	Yes Yes	5 Days
Patients with Comorbidities, Oral	Amoxicillin/Clavulanate 875 mg/125 BID Plus Azithromycin 500 mg x1 then 250 mg thereafter	Yes No	Yes Yes	5 Days
Patients with Comorbidities, IV	Ampicillin/Sulbactam 3 g IV Q6H <i>Plus</i> Azithromycin 500 mg x1 then 250 mg thereafter	Yes No	N/A N/A	5 Days

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Acute Exacerbation of COPD (C. pneumoniae, M. pneumoniae, S. pneumoniae, M. catarrhalis, H. influenzae)

AECOPD treatment involves steroids and bronchodilators, but does not routinely require antibiotics. Antibiotics are indicated for patients with the following 3 cardinal symptoms: increased dyspnea, sputum production, and sputum purulence. If increased sputum purulence is present, then treatment is indicated if only 2 symptoms are present.

	Antibiotic	Renal Dose Adjustment	Feeding Tube Compatible	Duration
Preferred	Doxycycline 100 mg BID OR Azithromycin 500 mg x1 dose then 250 mg OR Azithromycin 500 mg daily	No No No	Yes Yes Yes	5 Days 5 Days 3 Days
Recent Treatment, Oral	Amoxicillin/Clavulanate 875/125 mg BID OR Cefpodoxime 200 mg BID OR Cefuroxime 500 mg BID	Yes Yes Yes	Yes Yes Yes	5 Days 5 Days 5 Days
Recent Treatment, IV	Ampicillin/Sulbactam 3 g Q6H OR Ceftriaxone 1 g IV once daily	Yes No	N/A N/A	5 Days 5 Days
History of <i>P. aeruginosa</i>	Levofloxacin 750 mg once daily	Yes	Avoid*	5 Days

^{*}Poor absorption with quinolones observed when given via feeding tubes. See Appendix for more details. Intravenous formulations available

Thrush (Oropharyngeal Candidiasis) Candida albicans and other yeast					
	Antimicrobial	Renal Dose Adjustment	Feeding Tube Compatible	Duration	
Preferred	Clotrimazole Troche 10 mg five times daily	No	N/A	7-14 Days	
Alternative	Nystatin Suspension (100,000 units/mL) 4-6 mL four times daily	No	N/A	7-14 Days	
Moderate/Severe	Fluconazole 200 mg daily	Yes	Yes	7-14 Days	
Fluconazole, Refractory	Itraconazole 200 mg once daily	No	Yes	Up to 4 weeks	
•		1.1			

Denture-related candidiasis: disinfection of dentures recommended in addition to antifungal therapy

Coronavirus Disease 2019 (COVID-19)

High Risk of Progression: ≥65 years, asthma, BMI ≥30 kg/m², cardiovascular disease, CKD, diabetes, HIV or immunosuppressed, chronic lung disease, cancer, unvaccinated or prolonged duration since last vaccination

		Renal Dose	Feeding Tube	
	Antiviral	Adjustment	Compatible	Duration
High Risk of Progression to Severe Disease	Remdesivir 200 mg once followed by 100 mg IV daily OR Ritonavir-boosted nirmatrelvir (Paxlovid) 300/100 mg BID	No Yes	N/A See Below	3 Days 5 Days

Ritonavir-boosted Nirmatrelvir (Paxlovid) is interacts with many medications. See here for a list of common medications known to interact. No data exists on administering Ritonavir-boosted Nirmatrelvir (Paxlovid) via a feeding tube and therefore cannot be routinely recommended. Use of ritonavir-boosted Nirmatrelvir (Paxlovid) may have little benefit in vaccinated individuals and use should be used judiciously.

Influenza (Flu A, Flu B)

Vaccination: Inactivated Influenza Vaccine recommended for all residents. If ≥65 years, high-dose flu vaccines recommended annually before flu season

	Antiviral	Renal Dose Adjustment	Feeding Tube Compatible	Duration
Treatment	Oseltamivir 75 mg BID	Yes	Yes	5 Days
Prophylaxis, Post-exposure	Oseltamivir 75 mg once daily within 48 hours of contact Limit use to immunocompromised or severely ill patients within 48 hours of a person with suspected/confirmed influenza	Yes	Yes	1 Week (vaccinated) 2 Weeks (unvaccinated)
Prophylaxis, Outbreak	Oseltamivir 75 mg once daily for all residents in affected unit	Yes	Yes	2-3 Weeks*

^{*}CDC recommends prophylaxis in institutional outbreaks for a minimum of 2 weeks and continuing for at least 7 days after last known laboratory –confirmed case on affected units

Varicella Zoster Virus (VZV, Shingles)

Vaccination: Age 50 years or older: 2-dose series of Shingrix (RZV) given IM. First dose given at month 0 followed by a second dose 2-6 months later

	Antiviral	Renal Dose Adjustment	Feeding Tube Compatible	Duration**
Preferred	Acyclovir 800 mg five times daily	Yes	Yes	7 Days
Alternative	Valacyclovir 1 g TID	Yes	No*	7 Days
Ophthalmologic Disease/Severe	Acyclovir 10 mg/kg IV Q8H	Yes	N/A	7-14 Days

^{*}Consider acyclovir if given through a feeding tube.

	Antiviral	Renal Dose Adjustment	Feeding Tube Compatible	Duration
First Episode	Acyclovir 400 mg TID OR Valacyclovir 1 g BID	Yes Yes	Yes No*	7-10 Days 7-10 Days
Recurrent	Acyclovir 800 mg BID OR Valacyclovir 500 mg BID (Genital) OR Valacyclovir 2 g BID (Orolabial)	Yes Yes Yes	Yes No* No*	5 Days 3 Days 1 Day
Suppressive	Acyclovir 400 mg BID OR Valacyclovir 1 g daily	Yes Yes	Yes No*	Variable
Prophylaxis - Immunocompromised	Acyclovir 400 mg BID OR Valacyclovir 500 mg BID	Yes Yes	Yes No*	Variable
Special Population:	Persons with HIV			
Initial or Recurrent, Genital or Orolabial	400 mg TID Acyclovir OR 1 g BID Valacyclovir	Yes Yes	Yes No*	7-10 Days*

^{*}Consider acyclovir if given through a feeding tube. See appendix for additional dosing instructions. **Can be continued until resolution

^{**}Reasonable to continue treatment beyond 7-14 days if lesions resolve slowly

Trichomoniasis and Bacterial Vaginosis (Trichomonas vaginalis; Dysbiosis of vaginal flora)								
		Antibiotic	Renal Dose Adjustment	Feeding Tube Compatible	Duration			
Trichomoniasis	Female	Metronidazole 500 mg BID	No	Yes	7 Days			
Trictionioniasis	Male	Metronidazole 2,000 mg x 1 dose	No	Yes	1 Day			
	Oral	Metronidazole 500 mg BID	No	Yes	7 Days			
Bacterial Vaginosis	Tanical	Metronidazole 0.75% gel, 5 g vaginally once daily	No	N/A	5 Days			
	Topical	Clindamycin 2% cream, 5 g vaginally once daily	No	N/A	7 Days			

		Antibiotic	Renal Dose Adjustment	Feeding Tube Compatible	Duration		
Canarrhaa Drafarrad	<150 kg	Ceftriaxone 500 mg IM x1 dose	No	NI/A	1 Day		
Gonorrhea, Preferred	≥150 kg	Ceftriaxone 1,000 mg IM x1 dose	NO	N/A	1 Day		
Gonorrhea, Alternative		in 240 mg Intramuscular x1 dose <i>plus</i> vcin 2,000 mg x1 dose	No No	N/A Yes	1 Day		
Chlamydia, Preferred	Doxycycli	ne 100 mg BID	No	Yes	7 Days		
Chlamydia, Alternative	Azithromy	cin 1,000 mg x1 dose	No	Yes	1 Day		
Single doses of gentamicin	generally	do not require renal dose adjustments					
STI Additional Notes							
Re-testing	Any person who has a positive test for chlamydia or gonorrhea, along with women who have a positive test for trichomonas, should be rescreened 3 months after treatment						
Expedited Partner Therapy (EPT)	CDC supports issuing prescriptions to sex partners of those diagnosed with chlamydia or gonorrhea without the provider first examining the partner. EPT provides facilitation to treat partners with limited healthcare access.						

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Vulvovaginal Candidiasis (Candida albicans)

Most agents/formulations available OTC. '*' denotes prescription only. Topicals recommended in pregnancy

		Antimicrobial	Duration		
Oral	Fluconazole*	150 mg x 1 dose May repeat 72 hours later for those with moderate symptoms	1+ Day(s)		
	Clatrimazala	1% Vaginal cream, 5 g once daily	7 Days		
Vacinal Crass	Clotrimazole	2% Vaginal cream, 5 g once daily	3 Days		
Vaginal Cream	NA:la	2% Vaginal cream, 5 g once daily	7 Days		
	Miconazole	4% Vaginal cream, 5 g once daily	3 Days		
		100 mg vaginally once daily	7 Days		
Vaginal Tablet	Clotrimazole	200 mg vaginally once daily	3 Days		
		500 mg vaginally x1 dose	1 Day		
Variant Conservation	N 4' l -	100 mg vaginal suppository once daily	7 Days		
Vaginal Suppository	Miconazole	200 mg vaginal suppository once daily	3 Days		
Refractory or Alternative	Nystatin supp	Nystatin suppository* 100,000 units vaginally once daily			

Pressure Injuries/Ulcers (Including decubitus ulcers)

Localized damage to skin and/or underlying soft tissue as a result of prolonged pressure, shear, immobility, poor nutrition, co-morbidities, perfusion, microclimate, and/or perfusion leading to tissue damage ranging from intact skin to open ulceration.

	Treatment	Duration
Preferred	Systemic antibiotics not routinely recommended for treatment in most pressure injuries. In the presence of a pressure injury, follow appropriate wound care recommendations for management and prevention of worsening injury. Ulceration and exposed bone does not necessarily correlate with acute infection. If superimposed infection is suspected (new cellulitis, abscess, osteomyelitis on imaging/pathology, and/or systemic signs of infection antimicrobials may be recommended alongside mechanical debridement/drainage to support proper wound healing and attain source control.	N/A

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Tracheitis (Ventilator-Associated Tracheobronchitis, VAT)

Fever with no other recognizable cause, with new or increased sputum production, positive endotracheal aspirate (>10⁶ CFU/mL) yielding a new bacteria, and no radiographic evidence of pneumonia.

	Treatment	Duration
Preferred	Treatment Not Recommended. Monitor off antimicrobials. IDSA guidelines recommend AGAINST treatment of VAT. Limited data exists and it is uncertain if treatment improves clinical outcomes. Treatment is met with increased risk of <i>C. difficile</i> , drug adverse events, and selection for multi-drug resistance organisms in an already at-risk population. Rule out new airway obstruction and in the setting of new purulent secretions, correlate with systemic signs of infection, clinical symptoms and imaging to guide a new diagnosis of pneumonia.	N/A

Clostridioides difficile (C. diff)

High Risk Patients: immunocompromised, ≥ 65 years, and/or severe episode

	Antibiotic	Renal Dose Adjustment	Feeding Tube Compatible	Duration
1 st Episode	Vancomycin 125 mg PO Q6H x 10 days	No	Yes	10 Days
Alternative	Fidaxomicin 200 mg PO BID	No	Yes	10 Days
Recurrence	Vancomycin PO taper: 125 mg PO four times daily x 14 days <i>followed by</i> 125 mg PO TID x 7 days <i>followed by</i> 125 mg PO BID x 7 days <i>followed by</i> 125 mg PO daily x 7 days <i>followed by</i> 125 mg PO once weekly x 7 weeks	No	Yes	12 Weeks

For high-risk 1st episodes, vancomycin taper may be used instead of traditional 10 day course alongside bezlotoxumab IV. Bezlotoxumab is also recommended for recurrent cases. Fidaxomicin and Bezlotoxumab may be cost prohibitive. If multiple recurrences occur consider fecal microbiota transplant (FMT)

Antimicrobial	HD assumes thrice weekly. Time o		enal Function	/CrCl Adjuct	monts —				
Antimicrobiai		>25 mL/		11-25 mL/		≤10 mL/min	HD		
	Zoster	800 mg five ti		800 mg TID		800 mg BID	800 mg daily		
Acyclovir	HSV, 1 st Episode		400 mg TI				00 mg BID		
	HSV, Recurrent		800 mg BI				00 mg BID		
	HSV, Suppressive		400 mg BI				00 mg BID		
	Prophylaxis		400 mg BI				00 mg BID		
	Persons with HIV		400 mg TI				00 mg BID		
	Feeding Tu	be Additional Notes: Օլ	oen capsule or	use oral sus	pension p	referred over cru	shing tablet		
		>30 mL/min	10-30 mL/min <1		<10	0 mL/min	HD		
	Pneumonia	1 g TID	1 g B	SID	500 mg BID		500 mg BID		
Amoxicillin	Standard	500 mg BID/TID or 875 mg BID/TID	500 mg	g BID	500 mg daily		500 mg daily		
	Feeding Tube Additional Notes: Oral suspension preferred over crushing tablet/opening capsule								
	>30 mL/min	10-	10-30 mL/min		<10	mL/min	HD		
Amoxicillin/	875/125 mg BID	500	0/125 mg BID			500/125 mg	g daily		
Clavulanate	_	Feeding Tube Additional Notes: Oral suspension preferred over crushing tablet/opening capsule. Do not crush XR formulation and avoid Augmentin XR in renal insufficiency (use standard formulations).							
	>30 mL/min	15-	30 mL/min	<15		mL/min	HD		
Ampicillin/ Sulbactam	3 g every 6 hour	s 3 g e	very 12 hours	rs 3 g		3 g every 24	every 24 hours		
Juibactaiii	Feeding Tube Additional Notes: N/A								
0 -:+ laaaaia		No renal dose adjustment							
Azithromycin		Feeding Tub	e Additional N	lotes: Crush	tablet pre	eferred			
		>30 mL/min	11-30 ml	L/min	<1:	1 mL/min	HD		
	<120 kg	1 g every 8 hours	1 g every 1	.2 hours	1 g	every daily	1 g every daily		
Cefazolin	>120 kg	2 g every 8 hours	2 g every 1	y 12 hours 2 g		every daily	I g every daily		
	Feeding Tube Additiona	l Notes: N/A							

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Appendix (Note: HD	assumes thrice weekly. Time	doses <u>after</u> HI	D sessions wher	appropriate)				
	>30 mL/min		10-	30 mL/min		<10 mL/min		HD
Cefuroxime	500 mg BID	500 mg BID		500 mg every 24 hours		250 mg every 24 hours		
	Feeding Tube Additional Notes: Oral suspension preferred over crushing tablet/opening capsule (Bitter taste from tablet).							
	>30 mL/mii		10-30 mL/m	in		HD		
Cefpodoxime	200 mg BID)				200 mg daily		
	Feeding 1	ube Additio	onal Notes: O	ral suspension _ا	oreferred	over crushing table	t/openin	g capsule.
Ceftriaxone				No renal dose	adjustme	ent		
Certifiaxone			Fee	eding Tube Addi	tional No	tes: N/A		
		>30 m	nL/min	15-30 mL/	min	<15 mL/mir	า	HD
	Standard	1 g TI	D/QID	500 mg 7	ΓID	500 mg every oth		500 mg daily
Cephalexin	Uncomplicated UTI	500 mg BID			500 mg daily			
	Feeding Tube Additional Notes: Oral suspension preferred over crushing tablet/opening capsule							
		≥30 mL/min		n	<3	0 mL/min		HD
	Standard		750 mg BII)	500 mg BID			500 mg daily
Ciprofloxacin, PO	Uncomplicated UTI		250 mg BII)	250 mg daily			
	Feeding Tube Additional Notes: Diminished bioavailability. Avoid use. Give at least 2 hours before OR 6 hours after taking multivalent cations or antacids							
Clindamusia	No renal dose adjustment							
Clindamycin	Feeding Tube Additional Notes: Open capsule							
Dicloxacillin	No consen	No consensus on renal dose adjustment. Consider empiric dose adjustment in severe renal disease.						
			Feeding	Tube Additiona	l Notes: 0	Open capsule		

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Appendix (Note: H	HD assumes thrice weekly. Time	doses after HD sessions wh	nen appropriate							
••				dose adjustme	ent					
Doxycycline	Feeding Tube Addition hours before OR 4 hou	•			tablet or slow r	elease dosage f	orm. Give at least 2			
Fidenesials			No renal	dose adjustme	ent					
Fidaxomicin		Feeding Tube Additional Notes: Oral suspension preferred over crushing tablet								
		≥50 mL/n	nin	<50	mL/min		HD			
Fluconazole	Thrush	200 mg da	aily		10	00 mg daily				
		Feeding Tube Addition	al Notes: Ora	l suspension p	referred over c	rushing tablet				
Gentamicin		Variable dosing strategies exist								
Gentamicin				Additional Not						
		>50 mL/min	<20-50	mL/min	10-19 ml	_/min	HD			
	Standard	750 mg daily	750 mg eve	750 mg every other day 750		mg x1 then 500 mg every other day				
Levofloxacin	Uncomplicated UTI	250	mg daily 250 mg every other day				her day			
		Feeding Tube Additional Notes: Preferred over ciprofloxacin if an alternative cannot be used. Avoid multivalent cations and antacids within 2 hours of administration								
	>30 mL		10-30 mL/min			HD				
Nitrofurantoin	100 mg	g BID	Avoid use if CrCL <30 mL/min							
	Feeding Tube Additional Notes: Use oral suspension. Difficult to crush tablet									
		>60 mL/m	in 30-6	60 mL/min	10-29 mL/min		HD			
Oseltamivir	Treatment	75 mg Bll	75	mg daily		75 mg every ot	her day			
	Prophylaxis	75 mg dai	ly 75	mg every othe	ier day 75 mg ev		other HD session			
	Feeding Tube Additional Notes: Oral suspension preferred over opening capsule									
	>30 m	<u> </u>		10-30 mL/min			HD			
Remdesivir	200 mg x1 the			Caution use if CrCL <30 mL/min						
		Feeding Tube Additional Notes: N/A								

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Appendix (Note: HD	assumes thrice weekly. Tir	ne doses <u>after</u> HD session	s when appropriat	e)						
	>60 mL/m	in 30-	60 mL/min	<30 m/min	Н	D				
Ritonavir-boosted	300/100 mg	BID 150,	100 mg BID	See note*	300/100 mg x1 the	n 150/100 mg daily				
Nirmatrelvir	consider 150/100 mg	*Use is not recommended for CrCl <30 mL/min, but low risk of toxicity expected with a 5-day course. It is reasonable to consider 150/100 mg BID for 5 days if use is desired. Ritonavir is high-risk for drug-drug interactions. Recommend running interactions prior to prescribing. Feeding Tube Additional Notes: Avoid crushing tablet if possible due to limited data.								
		>30 mL/min		10-30 mL/min	<10 mL/min	HD				
Sulfamethoxazole/	Standard	1600/320 mg (2 DS) E	SID 1600/32	20 mg (2 DS) once daily	Avoid	Use				
Trimethoprim	UTI	800/160 mg (1 DS) B	D 800/16	0 mg (1 DS) once daily	Avoid	Use				
		Feeding Tube Addi	ional Notes: Or	al suspension preferred o	over crushing tablet					
		≥50 mL/min	30-49 mL/r	nin 10-29 mL/min	<10 mL/min	HD				
	Zoster/Shingles	1 g TID	1 g BID	1 g daily	500 r	ng daily				
	Orolabial HSV, Initial/Recurrent (Given for 1 day)	2 g BID	1 g BID	500 mg BID	500 mլ	g x1 dose				
	Genital HSV, Initial	1 {	g BID	1 g daily	500 mg daily					
Valacyclovir	Genital HSV, Recurrent	500	mg BID		500 mg daily					
	HSV, Suppressive	500	mg BID		500 mg daily					
	Prophylaxis	500	mg BID		500 mg daily					
	HIV, Initial/Recurrent (Genital/Orolabial)	1 1	g BID	1000 mg daily	500 r	ng daily				
	Feeding Tube Additional Notes: Consider acyclovir over Valacyclovir in setting of feeding tube									
			Variable d	osing strategies exist						
Vancomycin, IV	Additional Notes: IV vancomycin is for systemic infections only, NOT for treatment of <i>C. difficile</i> (see below)									
			No rena	l dose adjustment						
Vancomycin, PO	Feeding Tube Additional Notes: Oral suspension preferred over crushing tablet PO vancomycin is for infections caused by <i>C. difficile</i> only. Oral vancomycin formulations have minimal systemic absorption.									

Page | 15 Antibiotic preferences incorporate guideline recommendations and Kentucky resistance patterns. See Appendix for suggested renal dose adjustments

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