# African Antibiotic Treatment Guidelines for Common Bacterial Infections and Syndromes

For Adult Patients First Edition (English) Pending final publication in 2021

Draft for external review.

Not for circulation, reprint, or clinical use.

Prepared by: Africa Centres for Disease Control and Prevention The Center for Disease Dynamics, Economics & Policy





THE CENTER FOR Disease Dynamics, Economics & Policy WASHINGTON DC • NEW DELHI

### Table of Contents

Acronyms 2
List of Acronyms
Recommended Antibiotic Treatment for Common Bacterial Infections & Syndromes
Central Nervous System
Acute Bacterial Meningitis (Community-Acquired)
Acute Otitis Media
Dental Abscess including Gingivitis
Cardiac7
Infective Endocarditis - Native valve endocarditis7
Prosthetic valve or pacemaker infection
Upper & Lower Respiratory
Acute Bronchitis
Acute Exacerbation of Chronic Obstructive Pulmonary Diseases (COPD)
Mild to Moderate Community-Acquired Pneumonia (CAP) in Ambulatory Outpatients10
Severe Community-Acquired Pneumonia for Hospitalized Patients
Hospital-Acquired (Nosocomial) Pneumonia (HAP)11
Gastrointestinal13
Acute Invasive Diarrheal Disease (Dysentery)
Complicated Intra-Abdominal Infections (cIAI)
Typhoid (Enteric) Fever15
Genitourinary
Mild to Moderate Acute Uncomplicated Prostatitis
Uncomplicated Urinary Tract Infection (UTI)
Acute Pyelonephritis
Skin, Soft Tissue & Joints19
Skin & Soft Tissue Infections (SSTI)19
Acute Osteomyelitis & Septic Arthritis 20
Bloodstream21
Sepsis (Septicemia) & Septic Shock
References





#### Acronyms

# List of Acronyms

List of Actollyllis	
Pathogens	
	Acinetobacter baumannii
C. difficile	Clostridioides difficile
C. diphtheriae	Corynebacterium diphtheriae
C. trachomatis	Chlamydia trachomatis
E. coli	Escherichia coli
H. influenzae	Haemophilus influenzae
K. pneumoniae	Klebsiella pneumoniae
L. monocytogenes	Listeria monocytogenes
L. pneumophilia	Legionella pneumophilia
M. catarrhalis	Moraxella catarrhalis
N. gonorrhea	Neisseria gonorrhoeae
N. meningitidis	Neisseria meningitidis
<i>P. aeruginosa</i>	Pseudomonas aeruginosa
S. Aureus	Staphylococcus aureus
S. enterica	Salmonella enterica
S. epidermidis	Staphylococcus epidermidis
S. marcescens	Serratia marcescens
S. pneumoniae	Streptococcus pneumoniae
S. pyogenes	Streptococcus pyogenes
S. saprophyticus	Staphylococcus saprophyticus
Clinical	Stuphylococcus suprophylicus
AST	Antimicrobial suspectibility testing
CAP	Community-acquired pneumonia
cIAI	
CMV	Complicated intrabdominal infection
	Cytomegalovirus Chronic obstructive pulmonary disease
	C-reactive protein
CSF	
	•
HAP	Hospital-acquired pneumonia
	Human Immunodeficiency Virus
	Intramuscular
	Intravenous
	Procalcitonin
PO	Oral/by mouth
SSTI	Skin and soft tissue infection
TB	Tuberculosis
UTI	Urinary tract infection
VP	1
XDR	Extensively drug-resistant
Units of Measure	
g	Gram
IU	International unit
kg	Kilogram
mg	
mL	
MU	Million units
	3





Recommended Antibiotic Treatment for Common Bacterial Infections & Syndromes

Central Nervous System

Acute Bacterial Me	eningitis (Community-Acquired	)	
Clinical definition:	: Inflammation of meninges and	l subarachnoid space. C	
include headache,	fever, stiff neck, reduced consci	ousness. Major causes	of bacterial meningitis
	itidis, S. pneumoniae, L. monoc	cytogenes.	
Preferred antibioti	ic choice(s)		
Drug	<b>Formulation</b> <sup>1</sup>	Dosage	Duration
	Powder for injection: 250		If culture negative:
Ceftriaxone (IV)	mg; 1 g (as sodium salt) in	2 g 12 hourly	10 days. In case of
	vial		proven S.
Cefotaxime (IV)	Powder for injection: 250 mg	2 g 6 hourly	pneumoniae
	per vial (as sodium salt	2 g 0 110011y	infection: 14 days
Alternative antibio	ptic choice(s)		
	Powder for injection: 500		10 days, or if
Ampicillin (IV)	mg; 1 g (as sodium salt) in	3 g 6 hourly	confirmed <i>L</i> .
	vial	JSOHOUH	monocytogenes: 3
			weeks
Benzylpenicillin	Powder for injection: 600		_
(IV)	mg; 3 g (sodium or	4 MU 4 hourly	10 days
	potassium salt) in vial		
Chloramphenicol	Powder for injection: 1 g	1 g 6 hourly	10 days
(IV) <sup>A</sup>	(sodium succinate) in vial	0 /	10 augs
	ere penicillin allergy		
Ceftriaxone	Powder for injection: 250	2 g 12 hourly	If culture negative:
	mg; 1 g (as sodium salt) in		10 days. In case of
	vial		proven S.
Cefotaxime (IV)	Powder for injection: 250 mg	2 g 6 hourly	pneumoniae
	per vial (as sodium salt	2 g 0 nourly	infection: 14 days
In case of severe Penicillin allergy			
			If culture negative:
Moxifloxacin	Tablet: 400 mg or 100 mg		10 days. In case of
(IV, PO)	(dispersible); Injectable	400 mg once daily	proven S.
(1,10)	solution: 400mg/250 mL <sup>3</sup>		pneumoniae
			infection: 14 days

Principles of Stewardship:

- A. Chloramphenicol is not preferred and should only be used if other listed antibiotics are not available.
- > Acute meningitis may be caused by a range of pathogens, some of which are not bacteria. Microbiologic diagnosis, including bacterial culture from CSF, should be obtained as soon as possible to confirm etiology.
- > In presentations of subacute or chronic nature, consider diagnostic tests for TB meningitis, particularly in HIV-endemic areas.

Other Notes:





- Add ampicillin in situations of confirmed Listeria outbreaks and for patients at high risk for Listeria including: >
  - •
  - Patients over 50 years of age Immunosuppressed patients cancer, transplantation etc. Patients with alcoholism, cirrhosis, etc.

  - Pregnant women

Head, Eyes, Ear, Nose & Throat

Acute Otitis Med	ia		
	n: Acute infection and inflammatio	n of the middle ear.	Common symptoms
	and difficulty hearing. Common ba		
	. catarrhalis, S. aureus, Group A S		
Preferred antibio	tic choice(s)		
Drug	Formulation <sup>1</sup>	Dosage	Duration
Amoxicillin (PO)	Powder for oral liquid: 125 mg (as trihydrate)/5 mL; 250 mg (as trihydrate)/5 mL; Solid oral dosage form: 250 mg; 500 mg (as trihydrate)	500 mg 8 hourly	5 days
Alternative antib			•
Amoxicillin + clavulanic acid (PO)	Oral liquid: 125 mg amoxicillin + 31.25 mg clavulanic acid/5 mL AND 250 mg amoxicillin + 62.5 mg clavulanic acid/5 mL; Tablet: 500 mg (as trihydrate) + 125 mg (as potassium salt)	500 mg of amoxicillin component 12 hourly	5 days
In case of confirm	ned drug allergy or medical contrai	ndication	
Azithromycin (PO)	Capsule: 250 mg; 500 mg (anhydrous); Oral liquid: 200 mg/5 mL	500 mg daily	3 days
Principles of Stev	vardship:		
	has received antibiotics in the past e to amoxicillin.	month, use amoxic	illin-clavulanic acid in
Other Notes:			
> None			





Dental Abscess including Gingivitis

Clinical definition: Tooth infections from cavities, gingivitis, and periodontitis. Common symptoms include severe pain, tooth sensitivity, and inflammation of the face and gums. Most infections are polymicrobial and include anaerobic bacteria.

Preferred antibio	Preferred antibiotic choice(s)			
Drug	Formulation <sup>1</sup>	Dosage	Duration	
Amoxicillin- clavulanic acid (PO)	Oral liquid: 125 mg amoxicillin + 31.25 mg clavulanic acid/5 mL AND 250 mg amoxicillin + 62.5 mg clavulanic acid/5 mL; Tablet: 500 mg (as trihydrate) + 125 mg (as potassium salt)	500 mg component of amoxicillin 8 hourly	3 days if adequate source control, or 5 days if not	
Phenoxymethyl- penicillin (penicillin V) (PO)	Powder for oral liquid: 250 mg (as potassium salt)/5 mL; Tablet: 250 mg (as potassium salt)	500 mg 6 hourly	3 days if adequate source control, or 5 days if not	
	ed drug allergy or medical contrai	ndication		
Combination therapy with:	Azithromycin- Capsule: 250 mg; 500 mg (anhydrous); Oral liquid: 200 mg/5 mL	500 mg 6 hourly		
Azithromycin (PO) PLUS Metronidazole	Metronidazole- Oral liquid: 200 mg (as benzoate)/ 5 mL; Tablet: 200 mg to 500 mg	400 mg 8 hourly	5 days	
(PO)				
<ul> <li>Principles of Stewardship:</li> <li>&gt; Dental abscess requires surgical drainage, not just antibiotics.</li> <li>&gt; If the abscess is drained and the patient is improving, consider stopping antibiotics after 3 days of treatment.</li> <li>&gt; Although gingivitis is a risk factor for dental abscess, only acute necrotizing gingivitis should be treated with antibiotics</li> <li>&gt; For gingivitis without necrosis or abscess, do not treat with antibiotics.</li> </ul>				
<ul> <li>&gt; For acute necrotizing gingivitis:</li> <li>Treat with clindamycin [Dosage: Capsule: 150 mg (as hydrochloride); Injection: 150 mg (as phosphate)/ mL; Oral liquid: 75 mg/5 mL (as palmitate)] for 3 days</li> <li>For cases of acute necrotizing gingivitis associated with malnutrition, treat with</li> </ul>				

• For cases of acute necrotizing gingivitis associated with malnutrition, treat with vitamins.





Bacterial Pharyngotonsillitis, including Streptococcal & Diphtheria

Clinical definition: Infection causing acute inflammation of the pharyngeal wall and tonsils caused by various classes of *S. pyogenes* or *C. diphtheriae* (diphtheria). Common symptoms include sore throat, low-grade fever, and inflammation of the tonsils, uvula, lymph nodes, submandibular region, and neck.

Preferred antibiotic choice(s) Formulation<sup>1</sup> Dosage Duration<sup>2</sup> Drug Phenoxymethyl-Powder for oral liquid: 250 mg 500 mg 6 penicillin (as potassium salt)/5 mL; Tablet: 5 days (penicillin V) hourly 250 mg (as potassium salt) (PO) Powder for oral liquid: 125 mg (as trihydrate)/5 mL; 250 mg (as Amoxicillin 500 mg 8 trihydrate)/5 mL; Solid oral 5 days hourly (PO)dosage form: 250 mg; 500 mg (as trihvdrate) In case of confirmed severe penicillin allergy or medical contraindication Capsule: 250 mg; 500 mg Azithromycin (anhydrous); Oral liquid: 200 500 mg daily 3 days (PO) mg/5 mL

Principles of Stewardship:

> 85% or more of pharyngotonsillitis is viral. Most cases of pharyngotonsillitis in adults should be managed with watchful waiting & symptomatic relief. Antibiotics should not be considered unless there is a confirmed diagnosis of group A *Streptococcus*.

Other Notes:

> If clinical findings or epidemiologic context suggest diphtheria, treat with diphtheria antitoxin in addition to penicillin or macrolide.

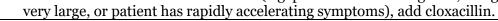
Cardiac

Infective Endocarditis - Native valve endocarditis					
Native valve endo	carditis				
		ole and non-specific. Commor	n etiologies include S.		
aureus <sup>A</sup> and strept	tococcal and enterococcal	species.			
Preferred antibiot	ic choice(s)				
Drug	<b>Formulation</b> <sup>1</sup>	Dosage	Duration		
Combination	Powder for injection:				
thereapy with:	600 mg; 3 g (sodium	5 MU 6 hourly	28 days		
	or potassium salt) in 5 MO 0 hourry 20 days				
Benzylpenicillin	vial				
(penicillin G, IV)	Gentamicin- Injection:				
	10 mg; 40 mg (as				
PLUS	sulfate)/mL in 2- mL	3 mg/kg daily	14 days		
	vial				
Gentamicin (IV)					
Alternative antibio	otic choice(s)				





			ſ
Combination	Ampicillin- Powder for		
therapy with:	injection: 500 mg; 1 g	2 g 4 hourly	28 days
Ampicillin (IV)	(as sodium salt) in vial		
	Gentamicin-Injection:		
PLUS	10 mg; 40 mg (as		14 days
	sulfate)/mL in 2mL	3 mg/kg daily	14 days
Gentamicin (IV)	vial		
In case of confirm	ed drug allergy or medica	l contraindication	
	Powder for injection:		
Vancomycin (IV)	250 mg (as	20 mg/kg 12 hourly	6 weeks
	hydrochloride) in vial		
	r pacemaker infection		
		n insertion or presence of pros	
		mon etiologies include S. aure	eus, S. epidermidis,
and other staphyle			
Preferred antibiot			
Drug	Formulation <sup>1</sup>	Dosage	Duration
Combination	Vancomycin- Powder	Loading dose: 25-30	
therapy with:	for injection: 250 mg	mg/kg followed by	
	(as hydrochloride) in	maintenance dose: 10-15	6 weeks
Vancomycin (IV)	vial	mg/kg	
DILIC	Gentamicin-Injection:		
PLUS	10 mg; 40 mg (as	- (1 1 1	- 1
Gentamicin (IV)	sulfate)/ mL in 2- mL	3 mg/kg daily	2 weeks
Gentamicin (1V)	vial		
PLUS	Rifampicin- Oral		
1 L05	liquid: 20 mg/mL;		
Rifampicin (PO)	Solid oral dosage form:	7.5 mg/kg 12 hourly	6 weeks
Kilampiciii (10)	150 mg; 300 mg		
Principles of Stew			
-	-	cases, 3 blood cultures should	l be obtained in
		tes within 6 hours before adm	
antibiotic		tes within o nours before adm	
		cases are culture-negative. The	n most common
		tics prior to the blood culture	
	-	tion by a fastidious organism,	-
		Fever), and <i>Brucella sp</i> , each	
		vestigation and treatment opti	
	laboratory.	construction and treatment opti	ons with your local
Other Notes:	100101013.		
	risk factors for S aurous	(e.g. patient is an IV drug use	or if vegetation is
		elerating symptoms) add clos	







Acute Bronchitis

Clinical definition: Inflammation of the upper airways due to viral infection or irritants.

Acute bronchitis is a viral infection and should **<u>NOT</u>** be treated with antibiotics.

Acute Exacerbatio	on of Chronic Obstructiv	e Pulmonary Diseases (C	OPD)	
			er than or equal to 5 on a	
			necessarily accompanied by	
	sputum volume, and/or			
	tic choice(s) – Mild-mod			
Drug	Formulation <sup>1</sup>	Dosage	Duration	
Amoxicillin	Powder for oral liquid: 125 mg (as trihydrate)/5 mL; 250 mg (as	500 mg 8 hourly	5 days	
(PO)	trihydrate)/5 mL; Solid oral dosage form: 250 mg; 500 mg (as trihydrate)	500 mg 0 nourly	5 days	
Doxycycline (PO)	Oral liquid: 25 mg/5 mL; 50 mg/5 mL (anhydrous); Solid oral dosage form: 50 mg; 100 mg (as hyclate)	200 mg STAT then 100 mg 12 hourly	5 days	
Preferred antibiotic choice(s) – Severe disease				
Amoxicillin + clavulanic acid (PO)	Oral liquid: 125 mg amoxicillin + 31.25 mg clavulanic acid/5 mL AND 250 mg amoxicillin + 62.5 mg clavulanic acid/5 mL; Tablet: 500 mg (as trihydrate) + 125 mg (as potassium salt)	500 mg of amoxicillin component 8 hourly	5 days	
In case of confirmed drug allergy or medical contraindication in severe disease				
Azithromycin	Capsule: 250 mg; 500 mg (anhydrous); Oral liquid: 200 mg/5 mL	500 mg daily	3 days	
Principles of Stewardship:				
> Up to 50% of infection-related acute exacerbations are viral. Biomarkers such as C-reactive protein (CRP) and procalcitonin (PCT) may play a role in differentiating,				

when available.





Other Notes:

> Exacerbations of COPD are commonly non-infectious and require optimization of non-antimicrobial therapeutic management.

Mild to Moderate Comr	Mild to Moderate Community-Acquired Pneumonia (CAP) in Ambulatory Outpatients			
Clinical definition: Pnet	umonia with onset in patier	nts not admitted to the h	nospital. Mild to	
	ity is treated in the outpatie	ent setting. (For severe C	CAP (CURB score	
>2), see below)	ina(a)			
Preferred antibiotic cho		Dogogo	Dunation	
Drug	Formulation <sup>1</sup> Amoxicillin- Powder for	Dosage	Duration	
	oral liquid: 125 mg (as			
	trihydrate)/5 mL; 250			
Amoxicillin (PO)	mg (as trihydrate)/5	1 g 8 hourly	5 days	
Autoxiciiiii (10)	mL; Solid oral dosage	1 g O Houriy	5 uays	
	form: 250 mg; 500 mg			
	(as trihydrate)			
Alternative antibiotic ch				
	Oral liquid: 25 mg/5			
	mL; 50 mg/5 mL			
Doxycycline (PO)	(anhydrous); Solid oral	100 mg 12 hourly	5 days	
	dosage form: 50	100 mg 12 nouny		
	mg;100 mg (as hyclate)			
	comorbidities – Alcoholisn		ılmonary disease,	
witnessed aspiration wh	nich is progressing after 24	-48 hours, etc.		
	Oral liquid: 125 mg			
	amoxicillin + 31.25 mg			
	clavulanic acid/5 mL			
Amoxicillin +	AND 250 mg	500 mg component	= dava	
clavulanic acid (PO)	amoxicillin + 62.5 mg clavulanic acid/5 mL;	of amoxicillin 8	5 days	
	Tablet: 500 mg (as	hourly		
	trihydrate) + $125 \text{ mg}$ (as			
	potassium salt)			
In case of confirmed dr	ug allergy or medical contra	aindication		
	Capsule: 250 mg; 500			
Azithromycin (PO) <sup>A</sup>	mg (anhydrous); Oral	500 mg daily	3 days	
	liquid: 200 mg/5 mL	0 0 7	0	
Principles of Stewardsh				
	Ild be avoided in countries			
pneumoniae, and should rather be reserved for treatment of patients with penicillin				
allergy.				
> Fluoroquinolones should be avoided, particularly in TB-endemic countries.				
Other Notes:				
> A blood culture is preferred to sputum culture if the patient is admitted to hospital.				
	or another macrolide is not	available, treat with a c	unolone such as	
moxifloxacin or	levofloxacin.			



Severe Community-Acquired Pneumonia for Hospitalized Patients Clinical definition: Severe disease is defined as CURB-65 score greater than two<sup>4</sup> and requires hospitalization.

Preferred antibiot	Preferred antibiotic choice(s)				
Drug	Formulation <sup>1</sup>	Dosage	Duration		
Combination	Ceftriaxone- Powder for				
therapy with:	injection: 250 mg; 1 g (as sodium salt) in vial	2 g daily	5 days		
Ceftriaxone	Cefotaxime - Powder for				
(IV/IM)	injection: 250 mg per vial	2 g 8 hourly	5 days		
OR	(as sodium salt				
	Clarithromycin-Solid oral				
Cefotaxime	dosage form: 500 mg;				
(IV/IM)	Powder for oral liquid: 125	500 mg by mouth	5 days		
DUID	mg/5 mL; 250 mg/5 mL;	12 hourly	Julys		
PLUS	Powder for injection: 500				
Clarithromyoin	mg in vial				
Clarithromycin	Azithromycin- Capsule:				
OR	250 mg; 500 mg				
	(anhydrous); Oral liquid:	500 mg daily	5 days		
Azithromycin	200  mg/5  mL				
(PO)	200 mg, 3 m2				
In case of confirm	ned drug allergy or medical con	ntraindication			
	Tablet: 400 mg; Tablet				
Moxifloxacin	(dispersible): 100 mg;	400 mg daily	5 days		
(IV/PO)	Injectable solution:	400 mg uany	5 days		
	400mg/250 mL <sup>3</sup>				
Principles of Stew	Principles of Stewardship:				

> Obtain a blood culture prior to starting antibiotic therapy.

If available, perform a legionella urinary antigen test – a positive result will allow > stopping of the b-lactam and extension of azithromycin to a minimum of 7 days to treat L. pneumophilia

In high TB-endemic areas, assess patients presenting with fever and  $cough \pm$ >

constitutional symptoms (anorexia, weight loss, night sweats) for active TB disease.

Doxycylcine may be used in place of a macrolide if unavailable. >

Other Notes:

None >

Drug

Hospital-Acquired (Nosocomial) Pneumonia (HAP)

Formulation<sup>1</sup>

Clinical definition: Pneumonia with onset at least 48 hours following hospital admission excluding ventilator-acquired pneumonia. Early onset HAP is defined as onset within 5 days of admission. Common etiologies of early onset HAP include S. Pneumoniae, S. aureus, H. *influenzae*, and enteric gram-negative bacilli. Late onset HAP is defined as onset after 5 days following admission; common etiologies include E. coli, S. marcescens, K. pneumoniae, A. baumannii, P. aeruginosa, and Enterobacter species.

Preferred antibiotic choice(s)



Safeguarding Africa's Health



Duration

Dosage

For facilities with low-level antibiotic resistance or where resistance is unknown and/or for				
	patients not transferred from facilities with high resistance:			
Ceftriaxone	Powder for injection: 250			
(IV)	mg; 1 g (as sodium salt) in vial	2 g daily	8 days	
Cofetarime (IV)	Powder for injection: 250	o a 9 hour	9 dava	
Cefotaxime (IV)	mg per vial (as sodium salt	2 g 8 hourly	8 days	
	Powder for injection: 500 mg (as sodium) + 100 mg			
Amoxicillin +	(as potassium salt); 1000	1 g of amoxicillin		
clavulanic acid (IV)	mg (as sodium) + 200	component 8 hourly	8 days	
	mg (as potassium salt) in vial.			
For facilities with	high Gram-negative resista	nce and/or for patients wit	h risk factors for	
resistance:		· · ·		
	Powder for injection: 2 g			
Piperacillin-	(as sodium salt) + 250 mg (as sodium salt); 4 g			
tazobactam	(as sodium salt) + 500	4.5 g 8 hourly	7 – 14 day	
(IV)	mg (as sodium salt) in			
vial       Alternative antibiotic choice(s)				
Alternative antibi				
Ertapenem	Powder for injection: 1g/vial <sup>3</sup>	1 g daily	7–14 days	
	ned drug allergy or medical of	contraindication		
Moxifloxacin (PO)	Tablet: 400 mg; Tablet (dispersible): 100 mg	400 mg daily	7 – 14 days	
Principles of Stev				
-	hoice of antibiotics for HAP	should be informed by the	local resistance	
	profiles in your hospital/unit.			
antibiotics.				
Switching from IV antibiotics to oral when patient can tolerate oral medication and as soon as signs and symptoms of infection are improving (e.g. clinical and laboratory				
white blood cell count improvement).				
Other Notes:				
> If risk factors for Pseudomonas infection exist, increase dosing frequency of				
	piperacillin-tazolbactam to 6-hourly, and use a 2 <sup>nd</sup> generation carbapenem (e.g.,			
meropene	meropenem or imipenem) in place of ertapenem.			



Ĵ



#### Gastrointestinal

Acute Invasive Diarrheal Disease (Dysentery)

Clinical definition: Acute infection commonly caused by bacteria resulting in bloody diarrhea, often with associated fever and abdominal pain. Bacterial etiologies include *Shigella flexneri*, *Campylobacter jejuni*, enteroinvasive and enterohaemorrhagic *E. coli*, and non-typhoidal *Salmonella* species. Dysentery may also be caused by the protozoan pathogen, *Entamoeba histolytica*.

Preferred antibiotic choice(s)				
Drug	<b>Formulation</b> <sup>1</sup>	Dosage	Duration	
Ciprofloxacin (PO)	Oral liquid: 250 mg/5 mL (anhydrous); Tablet: 250 mg (as hydrochloride)	500 mg 12 hourly	3 days	
Alternative antibiot	ic choice(s)			
Ceftriaxone (IV)	Powder for injection: 250 mg; 1 g (as sodium salt) in vial	1 g 12 hourly	5 days	
Azithromycin (PO)	Capsule: 250 mg; 500 mg (anhydrous); Oral liquid: 200 mg/5 mL	500 mg daily	3 days	
For severe cases or	those progressing despite cipro	floxacin, add Entan	noeba cover:	
Metronidazole (PO) Injection: 500 mg in 100- mL vial; Oral liquid: 200 mg (as benzoate)/5 mL; Suppository: 500 mg; 1 g; Tablet: 200 mg to 500 mg 7 days				
Principles of Stewar	Principles of Stewardship:			
<ul> <li>Non-bloody infectious diarrhea is generally caused by viruses and should not be treated empirically with antibiotics, but rather with supportive care and rehydration.</li> <li>Send stool sample for culture and sensitivity prior to starting antibiotics.</li> </ul>				
Other Notes:				
<ul> <li>In patients with advanced HIV and CD4 count &lt;100 cells/mm3, consider cytomegalovirus (CMV) colitis</li> </ul>				





Complicated Intra-Abdominal Infections (cIAI)				
	Intramural inflammation of the	gastrointestinal trac	t extending into the	
peritoneal space				
Preferred antibioti	c choice(s)			
Drug	Formulation <sup>1</sup>	Dosage	Duration <sup>5</sup>	
If mild to moderate	2:			
Amoxicillin + clavulanic acid (IV/PO)	Powder for injection: 500 mg (as sodium) + 100 mg (as potassium salt); 1000 mg (as sodium) + 200 mg (as potassium salt) in vial	875 mg of amoxicillin component 8 hourly		
If severe:				
Combination therapy with: Cefotaxime (IV)	Cefotaxime- Powder for injection: 250 mg per vial (as sodium salt	2 g 8 hourly	4 days if source	
PLUS Metronidazole (IV)	Metronidazole- Injection: 500 mg in 100- mL vial	500 mg 6 hourly	control has been achieved and clinical condition is improving. If not, duration will depend	
Combination therapy with:	Ampicillin- Powder for injection: 500 mg; 1 g (as sodium salt) in vial	200 mg/kg 4 hourly	on clinical and radiological progress, jointly managed with	
Ampicillin (IV) PLUS	Gentamicin- Injection: 10 mg; 40 mg (as sulfate)/ mL in 2- mL vial.	1 mg/kg 8 hourly	surgeons.	
Gentamicin (IV)	Maharaha Inistian			
PLUS	Metronidazole- Injection: 500 mg in 100- mL vial	500 mg 6 hourly		
Metronidazole (IV)				
If hospital-acquire	d in a facility where resistance h	as been documented		
Piperacillin- tazobactam	Powder for injection: 2 g (as sodium salt) + 250 mg (as sodium salt); 4 g (as sodium salt) + 500 mg (as sodium salt) in vial	4.5 g 6 hourly	4 days if source control has been achieved and clinical condition is improving. If not, duration will depend on clinical and radiological progress, jointly managed with surgeons.	
Alternative antibiotic choice				
Meropenem	Powder for injection: 500 mg (as trihydrate); 1 g (as trihydrate) in vial	1 g 8 hourly	4 days if source control has been achieved and clinical condition is	





			improving. If not, duration will depend on clinical and radiological progress, jointly managed with surgeons.
In case of confirme	d penicillin allergy or medical c	ontraindication	
Combination therapy with:	Clindamycin- Injection: 150 mg (as phosphate)/mL	20 mg/kg/day divided every 6 to 8 hours	4 days if source control has been
Clindamycin (IV) PLUS	Gentamicin- Injection: 10 mg; 40 mg (as sulfate)/ mL in 2- mL vial.	1 mg/kg 8 hourly	achieved and clinical condition is improving. If not,
Gentamicin (IV) OR	Ciprofloxacin- Solution for IV infusion: 2 mg/ mL (as hyclate)	500 mg 12 hourly	duration will depend on clinical and radiological progress, jointly managed with
Ciprofloxacin (IV)	ilyclate)		surgeons.
Principles of Stewardship:			
> Obtain a blood culture prior to starting any new antibiotic therapy.			
> Breach of the gastrointestinal tract mucosa is a risk factor for candida infection, which should be considered if source control and antibiotic treatment are not inducing a response.			
> Investigate for TB in endemic areas.			

Other Notes:

cIAI is often a difficult infection to treat and requires close collaboration with surgical colleagues to manage, as source control is a key aspect of management. >

Typhoid (Enteric) Fever				
Clinical definition: Systemic illness due to <i>S. enterica</i> serotype Typhi or Paratyphi, commonly acquired from ingestion of contaminated food or water. High fever and diarrhea or constipation are common presenting symptoms.				
Preferred antibiotic	c choice(s)			
Drug	<b>Formulation</b> <sup>1</sup>	Dosage	Duration	
For uncomplicated cases from outside of South Asia or Pakistan (low levels quinolone resistance):				
Ciprofloxacin (PO)	Oral liquid: 250 mg/5 mL (anhydrous); Tablet: 250 mg (as hydrochloride)	500 mg 12 hourly	For mild cases: 7 days For severe cases: 10 days	
For uncomplicated cases from South Asia or Pakistan (high levels quinolone resistance):				
Azithromycin (PO)	Capsule: 250 mg; 500 mg (anhydrous); Oral liquid: 200 mg/5 mL	500 mg daily	3 days	
For complicated cases, if patient is unable to take oral medication, or in case of confirmed				





drug allergy or medical contraindication:				
Ceftriaxone (IV, with de- escalation to ciprofloxacin or azithromycin depending on fluoroquinolone resistance)	Powder for injection: 250 mg; 1 g (as sodium salt) in vial	2 g daily	For mild cases: 7 days For severe cases: 10 days	
Alternative antibio	tic choice(s)			
Cefixime (PO)	Capsule or tablet: 200 mg; 400 mg (as trihydrate); Powder for oral liquid: 100 mg /5 mL	100 mg 12 hourly	For mild cases: 7 days For severe cases: 10 days	
Principles of Stewa	rdship:			
> Obtain a blood culture prior to starting antibiotic therapy.				
Other Notes:				
> Patients who aquire S. Typhi from Pakistan who have complicated, severe infection should be considered for empirical meropenem due to ongoing outbreak of XDR-S. <i>Typhi</i> .				
> Median time to fever reduction is 5 days.				





## Genitourinary

Mild to Moderate Acute Uncomplicated Prostatitis				
Clinical definition: Common symptoms include fever, chills, malaise, myalgia, pelvic pain,				
dysuria, and cloudy urine. In younger patients, common etiologies include <i>N. gonorrhea and C. trachomatis</i> . In older patients, common etiologies include <i>Enterobacteriaceae</i> species.				
Preferred antibiotic choic		s include Enterobaci	teriaceae species.	
	Formulation <sup>1</sup>	Docogo	Duration	
Drug		Dosage	Duration	
Ciprofloxacin (PO)	Oral liquid: 250 mg/5 mL (anhydrous); Tablet:	500 mg 12 hourly	10 – 14 days	
cipionoxaciii (10)	250 mg (as hydrochloride)		10 14 days	
Alternative antibiotic choice(s)				
	Capsule: 250 mg; 500			
Azithromycin	mg (anhydrous); Oral liquid: 200 mg/5 mL	500 mg daily	3 days	
Principles of Stewardship:				
> None				
Other Notes:				
> In sexually active men, syndromic treatment for gonorrhoea and chlamydia should be				

added, as per national protocol.

Uncomplicated Urinary Tract Infection (UTI)				
Clinical definition	: Infection of the bladder and	lower urinary tract. Sy	mptoms include	
urgency, dysuria,	and frequency of micturition.	UTIs are more commo	n in women than men.	
Commonly caused	l by the enterobacteriales, E. c	oli and K. pneumoniae	2	
Preferred antibiot	ic choice(s)			
Drug	Formulation <sup>1</sup>	Dosage	Duration	
Nitrofurantoin	Oral liquid: 25 mg/5 mL;	50 mg 6 hourly	5 days	
(PO)	Tablet: 100 mg.	50 mg 0 nourly	5 days	
Amoxicillin + clavulanic acid (PO)	Oral liquid: 125 mg amoxicillin + 31.25 mg clavulanic acid/5 mL AND 250 mg amoxicillin + 62.5 mg clavulanic acid/5 mL; Tablet: 500 mg (as trihydrate) + 125 mg (as potassium salt).	500 mg of amoxicillin component 12 hourly	5 days	
Principles of Stewardship:				
> Treatment with quinolones should be avoided.				
> Do not tro	> Do not treat notionts with asymptomatic bacteriumia execut in programmy, and consider			

> Do not treat patients with asymptomatic bacteriuria except in pregnancy, and consider in those persons undergoing genitourinary tract biopsy.

Other Notes:

> None



	Acute Pyelonephritis				
	Clinical definition: Bacterial infection of the kidney commonly presenting in women ages 18-				
	n symptoms include high feve				
	mmon etiologies include the		<i>li, K. pneumoniae,</i> and		
	<i>ruginosa</i> and <i>Enterococci</i> are				
Preferred antibioti	c choice(s)for mild-moderate	e cases			
Drug	Formulation <sup>1</sup>	Dosage	Duration		
Ciprofloxacin (PO)	Oral liquid: 250 mg/5 mL (anhydrous); Tablet: 250 mg (as hydrochloride)	500 mg 12 hourly	7 days		
For severe cases co	onsider:				
	Injection: 10 mg; 40 mg				
Gentamicin (IV)	(as sulfate)/ mL in 2- mL vial	5 mg/kg daily	7 days		
Amikacin (IV)	Injection: 250 mg (as sulfate)/mL in 2- mL vial	15 mg/kg daily	7 days		
Ceftriaxone (IV)	Powder for injection: 250 mg; 1 g (as sodium salt) in vial	1 g daily	7 days		
	Cefotaxime- Powder for		7 days		
Cefotaxime (IV)	injection: 250 mg per vial	1 g 8 hourly			
	(as sodium salt				
Principles of Stewardship:					
> Obtain urine and blood cultures for bacterial identification and condct antimicrobial					
suspectibility testing (AST) prior to starting antibiotic therapy.					
> If treating <i>Pseudomonas</i> infection with ciprofloxacin, increase dose to 750 mg and					
treat 12 hourly.					
Other Notes:					
> Avoid treatment with aminoglycosides in patients with renal impairment.					





# Skin, Soft Tissue & Joints

Skin & Soft Tissue	Skin & Soft Tissue Infections (SSTI)				
Clinical definition: Bacterial infections of skin and underlying soft tissue including cellulitis					
and abscess.			_		
Preferred antibiot	ic choice(s)				
Drug	<b>Formulation</b> <sup>1</sup>	Dosage	Duration		
Cloxacillin (PO)	Capsule: 500 mg; 1 g (as sodium salt); Powder for oral liquid: 125 mg (as sodium salt)/5 mL.	250 mg 6 hourly	5 days		
Alternative antibio	otic choices:	·			
Amoxicillin + clavulanic acid (PO) Used in animal bites <sup>A</sup>	Oral liquid: 125 mg amoxicillin + 31.25 mg clavulanic acid/5 mL AND 250 mg amoxicillin + 62.5 mg clavulanic acid/5 mL; Tablet: 500 mg (as trihydrate) + 125 mg (as potassium salt).	500 mg of amoxicillin component 8 hourly	5 days		
Cefalexin (PO)	Powder for reconstitution with water: 125 mg/5 mL; 250 mg/5 mL (anhydrous); Solid oral dosage form: 250 mg (as monohydrate)	500 mg 6 hourly	5 days		
In case of confirm	ed drug allergy or medical cont	traindication			
Clindamycin (PO)	Capsule: 150 mg (as hydrochloride); Oral liquid: 75 mg/5 mL (as palmitate)	300 mg 8 hourly	5 days		
Principles of Stewardship:					
<ul> <li>&gt; Withhold antibiotics for standard, uncomplicated abscess in an otherwise well person if the abscess can be incised and drained.</li> <li>&gt; If IV antibiotic therapy is clinically indicated, review patient progress at day 3 of treatment to consider switch from IV to oral therapy.</li> </ul>					
Other Notes:					
A. For patients with suspected animal bite, assess for rabies risk and consider administering a tetanus booster.					





Acute Osteomyelitis & Septic Arthritis Clinical definition: Acute osteomyelitis is an infection of bone infection with symptoms lasting days or a few weeks, commonly caused by methicillin-susceptible or resistant S. aureus. Common etiologies of septic arthritis include N. gonorrhea, S. aureus, Streptococcus species, and Gram-negative bacilli. Preferred antibiotic choice(s) Formulation<sup>1</sup> Dosage Drug Duration For the empiric treatment of acute osteomyelitis or septic arthritis: Powder for injection: 500 mg Cloxacillin (IV) 2 g 6 hourly 4 - 6 weeks (as sodium salt) in vial. Alternative antibiotic choice(s) Powder for injection: 250 mg; 1 g (as sodium salt) in Ceftriaxone (IV) 1 g daily vial Powder for injection: 250 mg Cefotaxime (IV) 2 g 8 hourly per vial (as sodium salt 4 - 6 weeks Powder for injection: 500 mg Amoxicillin + (as sodium) + 100 mg (as1 g Amoxicillin potassium salt); 1000 mg (as clavulanic acid componenent 8 (IV) sodium) + 200 mg (ashourly potassium salt) in vial For the treatment of monoarticular septic arthritis with STD risk Powder for injection: 250 Ceftriaxone (IV) mg; 1 g (as sodium salt) in 1 g daily 2 weeks vial In case of confirmed drug allergy or medical contraindication Clindamycin-Injection: 150 mg (as phosphate)/mL; Clindamycin 600 mg 8 hourly 2 weeks (IV) Oral liquid: 75 mg/5 mL (as palmitate) Principles of Stewardship: Do not give empirical antibiotics for chronic bone and joint infections. Instead, conduct bone and tissue biopsies, and treat with directed therapy. For septic arthritis, conduct a joint culture before administering antibiotic therapy and > refer to an orthopedic surgeon for assessment. If patient cannot take oral antibiotics, start with IV antibiotics and switch to oral > therapy as soon as patient is able to take antibiotics orally. Other Notes:

> Adequate drainage of purulent joint fluid is needed in addition to antibiotic therapy for septic arthritis.





#### Bloodstream

Sepsis (Septicemia) & Septic Shock

Clinical definition: Sepsis is life-threatening organ dysfunction caused by a dysregulated host response to infection. Septic shock is sepsis that requires vasopressor therapy to maintain blood pressure. **The choice of empiric antibiotic(s) will depend on the likely source of infection (see individual infections).** The guidance given here, relates to sepsis/septic shock where no infection source is immediately identifiable.

Preferred antibiotic choice(s) when no source is identified and/or is community-acquired with low risk of drug-resistant bacteria

Drug	Formulation <sup>1</sup>	Dosage	Duration	
Combination	Ampicillin- Powder for	u u	Durution	
therapy with:	injection: 500 mg; 1 g (as	200 mg/kg 4	10 days	
<b>T</b> J · ·	sodium salt) in vial	hourly		
Ampicillin (IV)	Amoxicillin-clavulanic acid-			
OR	Powder for injection: 500 mg	1 a amoviaillia		
	(as sodium) + 100 mg (as	1 g amoxicillin component 8	10 days	
Amoxicillin-	potassium salt); 1000 mg (as	hourly	10 days	
clavulanic acid	sodium) + 200 mg (as	nourry		
(IV)	potassium salt) in vial			
DLUG				
PLUS	Gentamicin- Injection: 10		,	
Contomioin	mg; 40 mg (as sulfate)/ mL	2 mg/kg 12 hourly	5 days	
Gentamicin (IV)	in 2- mL vial.			
Alternative antib	intic choice(s)			
Combination	Powder for injection: 250	2 g daily	10 days	
therapy with;	mg; 1 g (as sodium salt) in	2 g daily	10 days	
dierupy with,	vial			
Ceftriaxone				
(IV)				
	Gentamicin- Injection: 10			
PLUS	mg; 40 mg (as sulfate)/mL in	2 mg/kg 12 hourly	5 days	
	2-mL vial.			
Gentamicin				
(IV)				
	tic choice(s) when no source is i	dentified and hospital	acquired with high	
risk of drug-resistant bacteria				
Drug	Formulation <sup>1</sup>	Dosage	Duration	
Combination	Piperacillin-tazobactam			
therapy with	powder for injection: 2 g (as			
Piperacillin- tazobactam	sodium salt) + 250 mg (as	4.5 g 6 hourly	10 days	
(IV)	sodium salt); 4 g (as sodium salt) + 500 mg (as sodium			
	salt) in vial			
PLUS				
	Amikacin - Injection: 250 mg	15 mg/kg daily	5 days	
Amikacin (IV)	(as sulfate)/mL in 2- mL vial	10 mg/ ng uuny	Juujo	
	l	l	1]	





Principles of Stewardship:

> If the primary source of sepsis is defined, amend treatment duration according to the suggested duration for individual infections.

Other Notes:

- > Early administration of broad-spectrum antibiotics is critical in patients presenting with sepsis.
- > Amikacin has better coverage for extended-spectrum betalactamases than gentamicin.





#### References

- 1. World Health Organization Model List of Essential Medicines 21st List. 2019.
- Casey, J. R., Pichichero, M.E. (2005). Metaanalysis of Short Course Antibiotic Treatment for Group A Streptococcal Tonsillopharyngitis. The Pediatric Infectious Disease Journal; 24(10): 909-917.
- 3. Medscape. Drugs and Diseases. Retrieved 27 Jan 2021. Retrieved from: https://reference.medscape.com/
- 4. Jones, B.E., Jones, J.J., Bewick, T., Lim, W.S., Aronsky, D., Brown, S.M., Boersma, W.G., et al. (2011). CURB-65 Pneumonia Severity Assessment Adapted for Electronic Decision Support. CHEST; 140(1):150-63.
- 5. Sawyer, R. G., Claridge, J. A., Nathens, A. B., Rotstein, O. D., Duane, T. M., Evans, H.L., et al. (2015). Trial of Short-Course Antimicrobial Therapy for Intraabdominal Infection. *The New England Journal of Medicine*. 372: 1996-2005.

