

Bharati Vidyapeeth University Medical College Hospital & Research Centre, Pune Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026 Version - 11.0





Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Strilpa Acharya Quality Department	Dr. SK Nafwani Medical Director

Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier





Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026 Version - 11.0

S.NO	CONTENTS	PAGE NO
Prelude	List of amendments in Version 11	3
1.	Introduction	4
2.	Clinical Pathway	5
3.	Antimicrobial Stewardship: Goals workflow and Metrics	6-8
4.	a)List of Restricted antimicrobials	9
	b) Irrational Fixed dose antimicrobial combinations	9
	c)Irrational combination with double anaerobic coverage	9
	d)Common Types of antimicrobial resistant organisms	10
	Common isolates from various samples and their antibiogram for	>
	<u>the year 2024</u>	Y
5.	a) <u>Overall Hospital</u>	11-13
	b) Area wise OPD	14-16
	Critical care	17-20
	Medicine	21-23
	Surgery	24-26
	Neonatology	27-29
	Obst Gynae	30-31
	<u>Orthopedics</u>	32-33
	Pediatrics and PICU	34-36
6.	Antimicrobial Therapy in Hospitalized patients	37
7.	Empirical Antimicrobial Choice in various clinical conditions	38-62
	according to patient category, Definitive therapy	
8.	Surgical Antimicrobial prophylaxis/ Gastrointestinal endoscopy	63-65
	prophylaxis	
9.	Febrile Neutropenia	66-68
10.	Cardiac device implant prophylaxis	69
11.	Diagnostic Stewardship	70
12.	Appendix 1. Commonly used antimicrobials their spectrum of	71-73
_	activity and side effects	
13.	Appendix 2. Duration of therapy of common conditions	74
14.	Appendix 3. Antimicrobial Agent Form	75-76
15.	Appendix 4. Sample collection methods	77-79
	Appendix 5. List of BL-BLI combinations not recommended by WHO	80
16.	Appendix 3. List of BL-BLI combinations not recommended by WHO	80

Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK halwani Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

List of amendments in Version 11.0

- 1. Isolate listing and antibiogram for the year 2024
- 2. Updated Clinical Pathway (Pg.no 5)
- 3. Updated workflow of Antimicrobial Stewardship (Pg.no 7)
- 4. Restricted antimicrobials classified as per WHO watch and reserve class (Pg.no 9)
- 5. Added Irrational Fixed Dose Combination (Pg no.9)
- 6. Added list of Antibiotics with Overlapping anaerobic coverage (Pg no 9)
- 7. Added list of BL-BLI combinations not recommended by WHO (Appx 5)
- 8. Inj colistin changed to inj polymyxin B for treatment of XDR gram negative Pneumonia (Table 2)
- 9. Clarified timing of empiric dose of oseltamivir (Pg no 41)
- 10. Updated treatment guidelines for pelvic inflammatory disease (Table 4a)
- 11. Fluroquinolones removed from acute uncomplicated pyelonephritis (Table 4)
- 12. Updated empiric treatment of necrotizing fasciitis (Table 5)
- 13. Inj Vancomycin added to empiric treatment of, bone and joint infection in paediatrics (Table 10)
- 14. Loading doses of colistin, polymyxin and teicoplanin added (Table 21)
- 15. Added Diagnostic stewardship flowchart (Pg.no 69)
- 16. Table of duration of therapy updated. (Appx 2)
- 17. Added Antimicrobial agent form V3 (Appx 3)
- 18. Updated blood culture sample collection method (butterfly needle / adaptor) (Appx 4)

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

One Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

1. Introduction -

Over the last 75 years antimicrobials have been widely used to treat infectious diseases. Their indiscriminate use has led to resistance developing to almost all known antimicrobials. Antimicrobial resistance has become widespread not only in hospitals but also in the community.

A rational antimicrobial policy and antimicrobial stewardship is a must for all hospitals and is mandated by the Ministry of Health and Family Welfare through its document "National Policy for Containment of Antimicrobial Resistance, India". The purpose of this document is to provide a guide for rational antimicrobial use at Bharati Hospital based on local patterns of antimicrobial susceptibility.

DEFINITIONS

Antimicrobial agent (Antibiotic): Any agent, which has a potential for or is used with an intention of affecting microbial growth inside or on the human body. This includes antibacterial, antifungal, antiviral and anti-parasite agents.

Antimicrobial Stewardship: A set of coordinated activities that includes appropriate selection of antimicrobial agent, dosing, route and duration of antimicrobial therapy. The primary goal of antimicrobial stewardship is to optimize clinical outcomes while minimizing unintended consequences of antimicrobial use, including toxicity, the selection of pathogenic organisms and the emergence of resistance.

Surgical Antimicrobial Prophylaxis/Prophylactic anti-microbial agents: Administration of an antibiotic or antimicrobial agent prior to the commencement of a surgical procedure and appropriate readministration of the agent during prolonged surgery.

Empiric Antibiotic/Antimicrobial therapy: This is an early institution of antimicrobial therapy pending the results of culture and / or other relevant investigation and clinical response, in patients who have an illness and in whom there is an expectation of an infectious cause, the treatment being directed against the most likely microbial agent(s) in that particular episode.

Definitive Antimicrobial Therapy: Usage of antimicrobial agent against infection by specific microorganisms which have been confirmed by culture of appropriate samples.

Prepared by

Checked by

Verified by

Approved by

Dr. M Kumar

Prof & HoD

Prof Microbiology

Microbiology

Ochecked by

Verified by

Approved by

Dr. Sk Ialwani

Medical Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 4 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

2. Clinical Pathway

- 1. Postgraduate Resident of respective department will assess patient for symptoms and signs of infection, including laboratory evidence of infection.
- 2. He/she will document appropriately on the culture requisition form.
 - o suspected cause/site of infection,
 - o possibly community (CA)/hospital acquired(HA)
 - o patient type (types 1-3 described below)
- 3. Appropriate site cultures and blood cultures will be sent according to HICC protocol.
- 4. Antimicrobial will be chosen according to antimicrobial guide after informing lecturer on call and checking for allergy risks. Same will be endorsed in the case notes along with drug chart dose and indication
- 5. Any deviation from the policy will be documented along with the reason for deviation.
- 6. Some antimicrobials will be part of the restricted formulary and use of these "ALERT" antimicrobials will require infectious disease/ critical care (ICU/PICU/NICU) consult. These include:

Piperacillin/Tazobactam, Carbapenems, Ceftazidime avibactam, Aztreonam, Polymixins, Linezolid, Teicoplanin, Vancomycin, Echinocandins, Voriconazole, Amphotericin B

7. Clinical response will be followed up.

3harati

- 8. Once culture reports are available (Day 2 Day 4) antimicrobial is to be de-escalated (if possible) and duration of therapy is to be specified if not already done so.
- 9. Antimicrobial prescription should have a record of the day and expected duration of antimicrobials in the left-hand margin of the drug chart, eg D4/7
- 10. Antimicrobial Stewardship team will fill antimicrobial audit form and conduct regular department wise audits.

			/ .
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology		Dr. SK Nafwani Medical Director



2025-2026

Version - 11 (

3. Antimicrobial Stewardship



Antibiotic Stewardship



A set of coordinated strategies to improve the use of antimicrobials

50

Goal

- Enhancing patient health outcomes
- Reducing resistance to antibiotics
- Decreasing unnecessary costs

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

Ouality Department

Approved by

Dr. SK Malwani

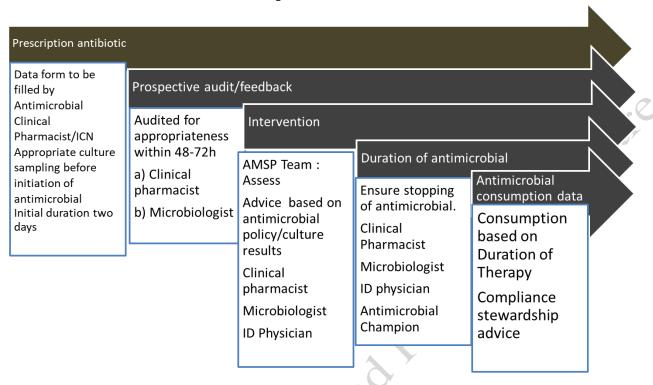
Medical Microbiology



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

4Workflow of Antimicrobial stewardship



Data of all patients who are prescribed antimicrobials are filled in the antimicrobial agent form and are audited for the following parameters:

- To ensure the antibiotics prescribed are appropriate as per their indication
- Streamlining or de-escalation will be done to ensure targeted therapy
- Dose optimization will be done based on body weight, site, renal and hepatic function and on drug pharmacokinetics and dynamics
- To ensure patient receives antibiotics for appropriate duration
- Surgical Prophylaxis will be monitored for correct timing and cessation at 24 hours

			/
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology		Dr. SK Nalwani Medical Nirector



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0



Metrics used in AMSP



- Days of therapy
- Cost metrics
- Average length of stay
- Mortality rate
- Acceptance of intervention
- Resistance pattern
- Comparison of HAI with ABX consumption rate

Bhatati Hospital and Res

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

Dr. Skilpa Acharya

Quality Department

Medical Pirector



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

4 a) List of Restricted Antimicrobials

These will not be prescribed without obtaining concurrence of HoD/HoU;

Antibiotics (as per WHO AWaRE Classification)	
WATCH	 Piperacillin/tazobactam Carbapenem Vancomycin Teicoplanin
RESERVE	Linezolid Daptomycin Tigogyeline
RESERVE	 Tigecycline Polymyxin B Colistin Ceftazidime avibactam
2	Aztreonam
Antifungals	
	Echinocandins : Caspofungin, Micafungin

4 b) Irrational Fixed Dose Combination

- 1) Ofloxacin+ Ornidazole
- 2) Norfloxacin+ Tinidazole
- 3) Ciprofloxacin+ Tinidazole
- 4) Ofloxacin + Tinidazole
- 5) Norfloxacin + Metronidazole

c) Antibiotics with overlapping anaerobic coverage

- 1) Piperacillin/tazobactam+ Metronidazole
- 2) Amoxicillin/clavulanate+ Metronidazole
- 3) Meropenem + Metronidazole
- 4) Clindamycin + Metronidazole
- 5) Piperacillin/tazobactam + Clindamycin
- 6) Meropenem + Clindamycin
- 7) Amoxicillin/clavulanate + Clindamycin

	1	1	1 . /	
Prepared by	Checked by	Verified by	Approved by	_
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Skrilpa Acharya Quality Department	Dr. SK halwani Medical Director	



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

(Detailed list of irrational BL-BLI antibiotic combinations is mentioned in Appendix 5)

4 d) Common antimicrobial resistant organisms:

Extended spectrum beta-lactamase producers (ESBL)

These are Gram negative organisms (GNB) like E coli & Klebsiella, which are resistant to the penicillin's; first-, second and third-generation cephalosporins; In addition, the plasmids bearing genesencoding ESBLs frequently also carry genesencoding resistance to other antimicrobial agents, such as aminoglycosides, trimethoprim, sulphonamides, tetracyclines and chloramphenicol. They remain susceptible to beta lactam- beta lactamase inhibitor combinations and carbapenems.

Amp C beta lactamases

These are inducible beta lactamases produced by certain organisms after exposure to cephalosporins. The organisms are resistant to the penicillins; first-, second- and third-generation cephalosporins and beta-lactam-beta lactamase inhibitor combinations. They may remain susceptible to cefepime and carbapenems. Seen in *Serratia*, *Pseudomonas*, *Proteus*, *Citrobacter and Enterobacter* spp.

Carbapenemase producers:

These are Gram negative organisms resistant to the Carbapenems and almost all beta-lactam antimicrobials except monobactams. Colistin and polymyxins are currently used for these organisms.

Methicillin Resistant Staphylococcus aureus (MRSA)

These are resistant to all beta lactam antimicrobials (Penicillins, BL-BLI, Cephalosporins, monobactams and Carbapenems.)

Vancomycin Resistant Enterococcus (VRE):

These isolates are resistant to Vancomycin, Teicoplanin but susceptible to linezolid.

MDR (Multi-drug resistance):

Isolates resistant to representatives of three or more classes of antimicrobial agents,

XDR (Extensive drug resistance):

Isolates resistant to all but one or two classes

PDR (Pan drug resistance):

Isolates resistant to all classes of antimicrobial agents available

			/
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology		Dr. SK Nalwani Medical Nirector

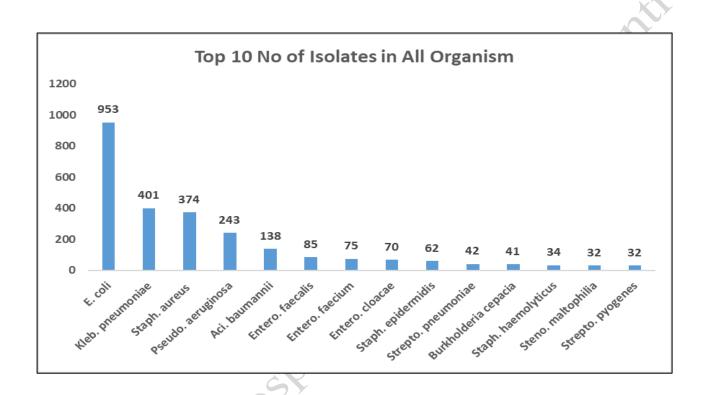


2025-2026 Version - 11.0

5. Organisms Commonly Isolated and Antibiogram:

Overall Hospital

All organism: Top 10 Isolates



Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar	Dr. M S Modak Prof & HoD	Dr. Stilpa Acharya	Dr. SK Nalwani	
Dr. M Kumar	Prof & HoD	Dr. Shilpa Acharya Quality Department	Dr. SK Laiwani Medical Director	



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Susceptibility Percentage of All Organism 2024

Organism	Number of patients	Nitrofurantoin	Penicillin	Ampicillin	Oxacillin	Levofloxacin	Ciprofloxacin	Norfloxacin	Ofloxacin	Cotrimoxazole	Amox/Clav	Cefuroxime	Cefuroxime axetil	Ceftazidime	Ceftriaxone	Cefotaxime	Cefepime	Cef/Sul	Piptaz	Amikacin	Gentamicin	Gentamicin high leve	Clindamycin	Erythromycin	Tetracycline	Ertapenem	Meropenem	Imipenem	Tigecycline	Daptomycin	Linezolid	Teicoplanin	vancomycin	Fosfomycin	Minocycline	Colistin
Escherichia coli	953	84	0	22		0	11	45	46	49	50	15	15	54	22	6	29	62	71	87	76				88	83	85	84	99					99	100	100
Klebsiella pneumoniae	401	28	0			0	37	78	78	51	48	35	36	69	40	29	43	51	51	65	66	0		67	100	60	58	56	91	0				0	100	100
Pseudomonas aeruginosa	243					64	65	100	100	67				77			78		84	74	71						72	71	0						100	100
Acinetobacter baumannii	138					100	15			30				0	15		12		13	13	19					0	11	10							100	99
Enterobacter cloacae	70	0					69	75	50	79	3	42	33	80	63	0	79	84	83	91	80			100	100	85	89	87	93					0	0	
Burkholderia cepacia	41					62	0			80				92	0		4	37	5	12	8						92								76	
Staphylococcus aureus	374	100	7		45	8	8		0	46											74	100	90	44	93	0			100	97	100	100	99			
Enterococcus faecalis	85	95	93			28	28															0	0	4	11				100	68	99	99	99			
Enterococcus faecium	75	8	6			5	5			0														3	9				0		95	81	89			
Staphylococcus epidermidis	62	100	13		22	48	49			48											80	100	73	25	80				0	100	100	88	100			
Streptococcus pneumoniae	42		100	0		92	0			56	0	0			92	100	0	0	100	100	100		47	11	31		100		97		97		97			
Staphylococcus haemolyticus	34	100	12		10	4	9			35											27		18	0	71				0	100	100	100	100			
Stenotrophomonas maltophilia	32					87				65				0																					67	
Streptococcus pyogenes	32		100	90		57	0							100	96	100							96	39	65				100		100		100			

Reserved/ Restricted Drugs:Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Dr. M Kumar

Prof & HoD

Prof Microbiology

Microbiology

Quality Department

Approved by

Dr. SK halwani

Medical Pirector



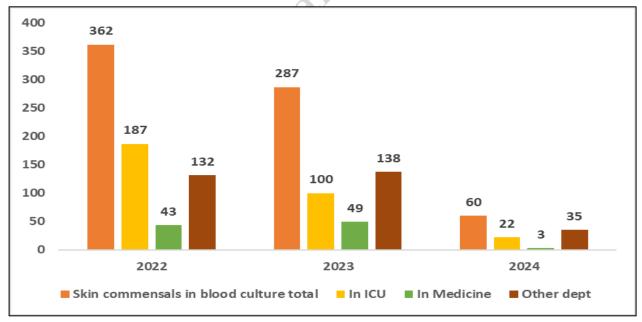
2025-2026 Version - 11.0

Candida Isolated in all location 2024

Specimen type	Organism	Number of patients	Fluconazole	Voriconazole	Ampho B	Caspofungin	Micafungin
	Candida albicans	12	91	100	33	100	92
	Candida auris	13	0	0	100	100	100
Blood	Candida glabrata	3		0	67	33	100
	Candida parapsilosis	21	76	95	85	95	95
	Candida tropicalis	8	88	100	100	100	88
Fluid	Candida parapsilosis	1	100	100	100	100	100
	Candida albicans	2	100	50		100	100
Pus	Candida parapsilosis	2	100	100	100	100	100
	Candida tropicalis	2	100	100	100	100	100
	Candida albicans	5	100	100	20	100	100
Urine	Candida parapsilosis	2	50	100	100	100	100
	Candida tropicalis	3	67	100	100	100	100

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Skin flora/Collection contamination in blood culture:



There has been a reduction in the number of collection contamination or skin

Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Naiwani Medical Pirector

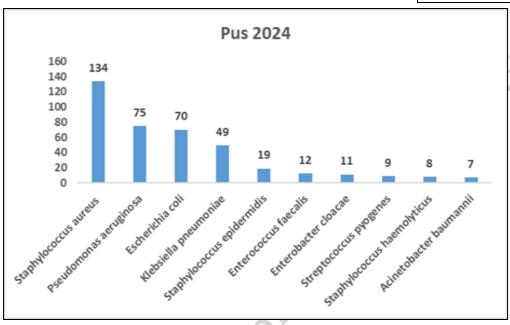


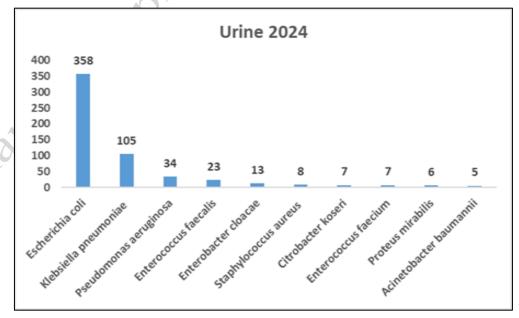
2025-2026 Version - 11.0

flora reported in blood culture during the year 2024 Organism Commonly Isolated and Antibiogram: Area wise

OPD: Top 10 Isolates Sample wise

Specimen	Overall No. of
Туре	Isolates
Pus	451
Urine	601





Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Naiwani Medical Nirector



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

OPD Antibiogram:

Percentage Susceptibility Gram Negative Organism 2024

Location	Specimen type	Organism	Number of patients	Nitrofurantoin	Levofloxacin	Ciprofloxacin	Cotriamoxazole	Amox/Clav	Cefuroxime	Cefuroxime axetil	Ceftazidime	Ceftriaxone	Cefotaxime	Cefepime	Cef/Sul	Piptaz	Amikacin	Gentamicin	Ertapenem	Meropenem	Imipenem	Tigecycline	Fosfomycin	Minocycline	Colistin
		Escherichia coli	39			7.7	49	54	18	21		26		28	74	82	81	74	90	86	85	100			100
	Pus	Klebsiella pneumoniae	26			39	54	46	46	47		50		50	58	56	58	68	62	59	65	100			100
		Pseudomonas aeruginosa	51		63	67	50		100		90			94	79	89	90	80		89	84			100	100
OPD		Escherichia coli	358	85	0	15	50	52	20	18	59	28	14	35	69	75	86	75	90	88	88	99	99	100	100
	Urine	Klebsiella pneumoniae	105			38	48	47	33	32	73	40	43	42	46	51	67	64	63	54	50	88			100
	orine	Pseudomonas aeruginosa	34		26	33					46			47	35	80	58	46		37	40	0			100
		Enterobacter cloacae	13			85	85	8	60	50	67	62		100	100	85	92	92	85	100	100	100			100

Reserved/ Restricted Drugs:Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Prepared by

Checked by

Verified by

Approved by

Dr. M Kumar

Prof & HoD

Prof Microbiology

Microbiology

Ochecked by

Verified by

Approved by

Dr. Sk Ialwani

Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 V

Version - 11.0

Percentage susceptibility Gram Positive organism 2024

Locatio	Specimen type	Organism	Number of patients	Nitrofurantoin	Oxacillin	Penicillin	Ciprofloxacin	levofloxacin	Clindamycin	Gentamicin	Gentamicin high level	Erythromycin	Tetracycline	Sulfa/Trimeth		Tigecycline	Daptomycin	Linezolid	Teicoplanin	Vancomycin
	Dura	Staphylococcus aureus	125		45	6	9	10	89	73	100	30	91	45	:	100	99	100	100	100
OPD	Pus	Staphylococcus epidermidis	12		27		75	78	67	82		33	73	58		0	100	100	100	100
	Urine	Enterococcus faecalis	23	86		100	17	11										100	100	100

Reserved/ Restricted Drugs:Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Dr. M Kumar

Prof & HoD

Prof Microbiology

Microbiology

Quality Department

Approved by

Dr. SK Italwani

Medical Director



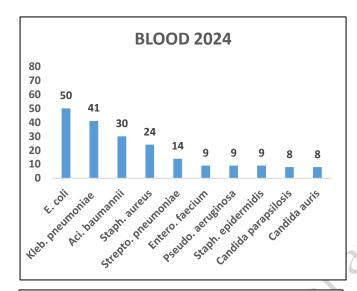


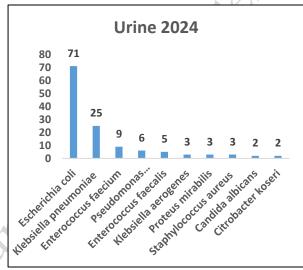
2025-2026

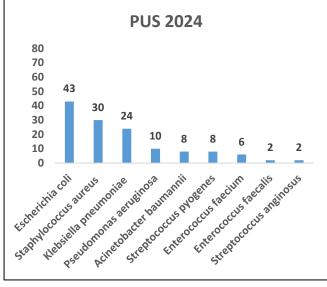
Version - 11.0

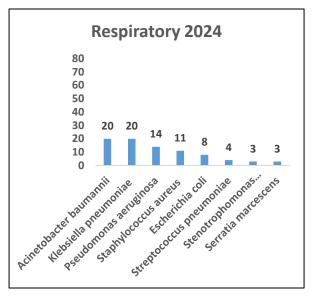
INTENSIVE CARE UNIT: TOP 10 ISOLATES BY SAMPLE:

	OVERALL
SPECIMEN	NO. OF
TYPE	ISOLATES
Blood	290
Pus	142
Respiratory	89
Urine	140









			/ .
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shipa Acharya Quality Department	Dr. SK Nalwani Medical Nirector





2025-2026

Version - 11.0

INTENSIVE CARE UNIT ANTIBIOGRAM:

Percentage Susceptibility Gram Negative Organism 2024

Location	Specimen type	Organism	Number of patients	Levofloxacin	Ciprofloxacin	Cotrimoxazole	Amox/Clav	Cefuroxime	Cefuroxime axetil	Ceftazidime	Ceftriaxone	Cefepime	Cef/Sul	Piptaz	Amikacin	Gentamicin	Ertapenem	Meropenem	Imipenem	Tigecycline	Fosfomycin	Colistin
		Escherichia coli	50	0	6	47	5 9	14	14		22	32	69	80	83	80	84	84	86	100		100
	Blood	Klebsiella pneumoniae	41		27	48	36	24	25		29	32	32	39	51	61	35	42	38	90		100
		Acinetobacter baumannii	30		7	37	0	0	0		8	7	7	7	3	11		3	3			100
	Dura	Escherichia coli	17		12	47	41	12	13		12	24	59	59	71	53	69	81	69	94		100
	Pus	Klebsiella pneumoniae	12		17	42	18	18	11		17	17	17	17	25	50	20	27	33	80		100
ICU	Tissue	Escherichia coli	15		13	53	31	14	18		13	27	40	40	62	47	58	64	69	100		100
	11 of a s	Escherichia coli	71		9	48	47	12	14	100	12	26	55	60	89	80	71	76	76	100	99	100
	Urine	Klebsiella pneumoniae	25		25	28	36	16	17		16	21	36	33	60	56	46	44	39	83		100
		Acinetobacter baumannii	20		5	20					6	5	0	5	5	10		5	5			100
	Respiratory	Klebsiella pneumoniae	20		11	32	29	5	0		5	11	0	26	40	45	33	35	31	87		100
		Pseudomonas aeruginosa	14	85	79	0	100			85		85	0	90	86	100		58	58			100

Reserved/ Restricted Drugs:Not to be used emperically unless justified Will be useful clinically
Will be useful clinically in about 2/3 cases
Will not be useful clinically
Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

			, .	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Stilpa Acharya Quality Department	Dr. SK halwani Medical Director	



2025-2026

Version - 11.0

Percentage susceptibility gram positive organism 2024

I	ocation	Specimen type	Organism	Number of patients	Oxacillin	Penicillin	Ciprofloxacin	Levofloxacin	Moxifloxacin	Clindamycin	Gentamicin	Erythromycin	Tetracycline	Sulfa/Trimeth	Tigecycline	Daptomycin	Linezolid	Teicoplanin	Vancomycin
		Dland	Staphylococcus aureus	24	57	17	17	21		95	48	48	96	48	100	95	100	100	100
	ICII	Blood	Streptococcus pneumoniae	14		100		82	90	67		9	23	31	100		100		100
	ICU	Pus	Staphylococcus aureus	30	48	14	3	3		94	76	44	94	64	100	96	100	100	97
		Respiratory	Staphylococcus aureus	11	30	0	10	10		100	73	50	91	46	100	89	100	100	100

Reserved/ Restricted Drugs:Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Dr. M Kumar

Prof & HoD

Prof Microbiology

Microbiology

Quality Department

Approved by

Dr. SK Italwani

Medical Director



2025-2026

Version - 11.0

Candida Isolated in ICU

Location	Specimen type	Organism	Number of patients	Fluconazole	Voriconazole	Ampho B	Caspofungin	Micafungin
		Candida albicans	6	100	100	0	100	100
		Candida glabrata	2		0	50	50	100
		Meyerozyma guilliermondii	2	0	0	50	50	50
	Blood	Issatchenkia orientalis	1	0	100	0	100	100
		Candida parapsilosis	8	63	100	75	100	100
		Candida auris	8	0	0	100	100	100
1011		Candida tropicalis	4	75	100	100	100	75
ICU	Fluid	Issatchenkia orientalis	1	0	100	0	100	100
	Pleural fluid	Candida glabrata	1		0	100	100	100
	Pus	Candida tropicalis	1	100	100	100	100	100
		Candida albicans	2	100	100	0	100	100
	Urine	Meyerozyma guilliermondii	1	0	0		100	100
	Offile	Candida parapsilosis	2	50	100	100	100	100
		Candida tropicalis	1	100	100	100	100	100

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

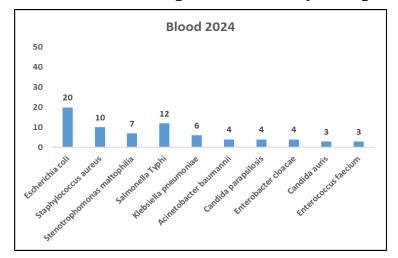
Dr. Ski laiwani

Medical Director

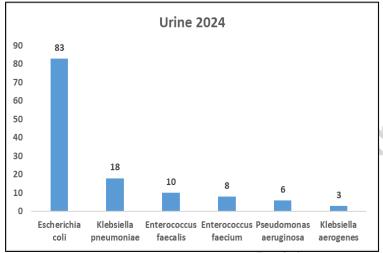


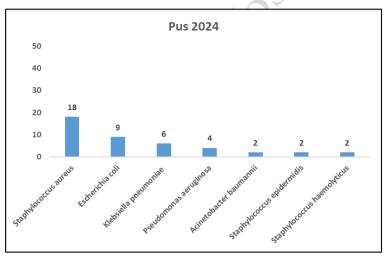
2025-2026 Version - 11.0

Medicine: Top 10 Isolates by Sample



Specimen	Overall No. of
Туре	Isolates
Blood	93
Pus	56
Urine	142





Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Naiwani Medical Pirector



2025-2026

Version - 11.0

Medicine Antibiogram Gram Negative:

Percentage Susceptibility Gram Negative Organism 2024

L	ocation.	Specimen type	Organism	Number of patients	Ciprofloxacin	Norfloxacin	Cotrimoxazole	Amox/clav	Cefuroxime	Cefuroxime axetil	Ceftazidime	Ceftriaxone	Cefotaxime	Cefepime	Cef/Sul	Piptaz	Amikacin	Gentamicin	Ertapenem	Meropenem	Imipenem	Tigecycline	Fosfomycin	Colistin
		Dland	Escherichia coli	20	5		60	35	10	11		10	0	26	65	75	90	90	95	95	95	100		100
	MED	Blood	Salmonella Typhi	12	17		92					100							100	100	100			
	IVIED	Huina	Escherichia coli	83	7	0	39	49	13	13	0	15	0	22	57	63	83	68	75	78	81	99	99	100
		Urine	Klebsiella pneumoniae	18	41	100	61	67	50	50	0	44	0	53	65	71	78	83	72	77	77	94		100

Reserved/ Restricted Drugs:Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

[D	Checked by	Verified by	Approved by	
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Smipa Acharya Quality Department	Dr. SK Laiwani Medical Director	



2025-2026 Version - 11.0

Medicine Antibiogram Gram Positive:

Percentage susceptibility gram positive organism 2024

Location	Specimen type	Organism	Number of patients	Nitrofurantoin	Oxacillin	Penicillin	Ciprofloxacin	Levofloxacin	Clindamycin	Gentamicin	Erythromycin	Tetracycline	Sulfa/Trimeth	Tigecycline	Daptomycin	Linezolid	Teicoplanin	Vancomycin
MED	Blood	Staphylococcus aureus	10		56	25	11	0	100	90	44	100	22	100	100	100	100	100
MED	Urine	Enterococcus faecalis	10	90		100	20	20			0	0		100	78	100	100	100

Reserved/ Restricted Drugs: Not to be used emperically unless justified Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Candida Isolated in Medicine:

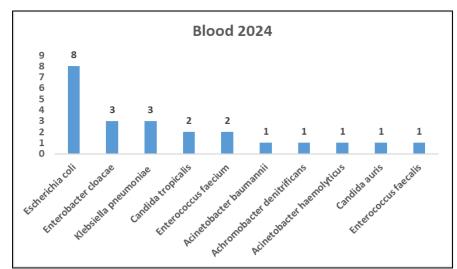
Location	Specime n type	Organism	Number of patients	Fluconazole	Voriconazole	Ampho B	Caspofungin	Micafungin
		Candida albicans	1			100	100	100
		Candida glabrata	1		0	100	0	100
	Blood	Candida parapsilosis	4	75	100	100	100	100
Med		Candida auris	3	0	0	100	100	100
		Candida tropicalis	1	100	100	100	100	100
	Pus	Candida parapsilosis	1	100	100	100	100	100
	Urine	Candida parapsilosis	1	100	100	100	100	100

Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Naiwani Medical Pirector

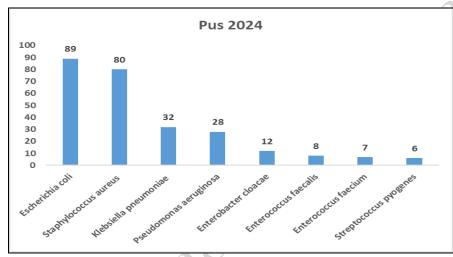


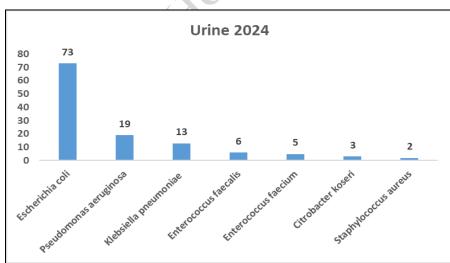
2025-2026 Version - 11.0

Surgery: Top 10 Isolates by Sample



Specimen	Overall No.
Туре	of Isolates
Blood	25
Pus	307
Urine	126





			/ .
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shipa Acharya Quality Department	Dr. SK Nalwani Medical Nirector



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Surgery Antibiogram:

Percentage Susceptibility Gram Negative Organism 2024

Location	Specimen type	Organism	Number of patients	Levofloxacin	Ciprofloxacin	Cotrimoxazole	Amox/Clav	Cefuroxime	Cefuroxime axetil	Ceftazidime	Ceftriaxone	Cefepime	Cef/Sul	Piptaz	Amikacin	Gentamicin	Ertapenem	Meropenem	Imipenem	Tigecycline	Fosfomycin	Colistin
	Pus	Escherichia coli	3 9		8	41	46	14	17		18	24	58	65	94	77	79	79	82	100		100
	ru3	Klebsiella pneumoniae	12		75	83	80	50	44		58	50	64	82	83	92	78	78	78	100		100
		Escherichia coli	14		7	50	50	7	8		7	21	64	69	93	57	92	92	92	100		100
Sura	Tissue	Klebsiella pneumoniae	11		36	46	30	36	22		46	44	50	50	55	55	44	44	33	90		100
Surg		Pseudomonas aeruginosa	10	67	67					88		78	70	88	100	100		80	78			100
		Escherichia coli	73		8	48	41	10	8		12	25	58	70	85	59	81	84	84	98	99	100
	Urine	Pseudomonas aeruginosa	19	28	32					33		37	22	63	50	43		39	39			100
		Klebsiella pneumoniae	13		33	46	36	39	36		39	33	46	33	54	46	55	50	55	82		100

Reserved/ Restricted Drugs: Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

One Medical Director



2025-2026 Version - 11.0

Percentage susceptibility gram positive organism 2024

Location	Specimen type	Organism	Number of patients	Oxacillin	Penicillin	Ciprofloxacin	levofloxacin	Ofloxacin	Clindamycin	Gentamicin	Gentamicin high level	Erythromycin	Tetracycline	Sulfa/Trimeth	Tigecycline	Daptomycin	Linezolid	Teicoplanin	Vancomycin
Surg	Pus	Staphylococcus aureus	80	45	5	8	4	0	92	75	100	51	98	47	100	97	100	100	100

Reserved/ Restricted Drugs: Not to be used emperically unless justified Will be useful clinically Will be useful clinically in about 2/3 cases Will not be useful clinically Not tested/not appropriate antibiotic

Candida Isolates:

Candida Isolated in Surgery

Location	Specimen type	Organism	Number of patients	Fluconazole	Voriconazole	Ampho B	Caspofungin	Micafungin
	Blood	Candida auris	1	0	0	100	100	100
	Blood	Candida tropicalis	2	100	100	100	100	100
	Fluid	Candida parapsilosis	1	100	100	100	100	100
Surg		Candida parapsilosis	2	50	100	100	100	100
	Pus	Candida albicans	2	50	50	0	100	100
		Candida tropicalis	1	100	100	100	100	100
	Urine	Candida albicans	1	100	100	100	100	100

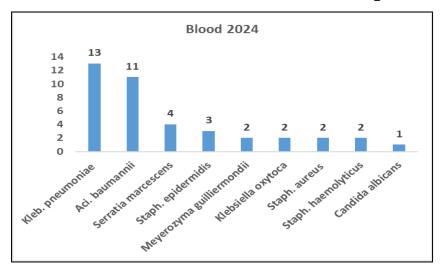
Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Naiwani Medical Pirector

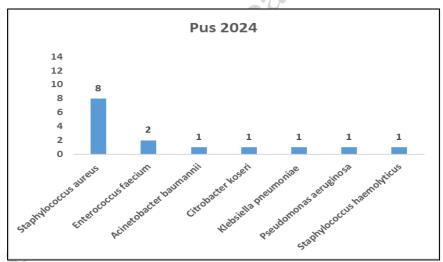


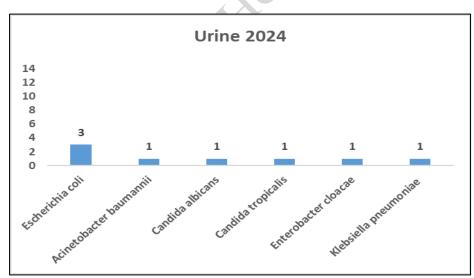
2025-2026 Version - 11.0

Neonatal Intensive Care Unit: Top 10 Isolates Sample wise:



Specimen	Overall No. of
Туре	Isolates
Blood	53
Pus	15
Urine	8





Dd but	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Simpa Acharya Ouality Department	Dr. SK Valwani Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

NICU Antibiogram:

Percentage Susceptibility Organism -NICU 2024

Organism	Number of patients	Cotrimoxazole	Amox/clav	Ciprofloxacin	Levofloxacin	Penicillin	Oxacillin	Cefuroxime	Cefuroxime Axet	Ceftriaxone	Cefepime	Cel/Sul	Piptaz	Amikacin	Gentamicin	Clindamycin	Erythromycin	Tetracycline		lmipenem	Meropenem	Teicoplanin	Linezolid	Daptomycin	Vancomycin	Tigecycline	Fosfomycin	Colistin
Klebsiella pneumoniae	15	67	50	40				40	36	53	60	64	64	86	100			100	79	71	71					100		100
Acinetobacter baumannii	13	54		23				0	0	36	25	33	33	23	31				0	18	25							100
Escherichia coli	5	80	80	0				0	0	0	25	50	50	100	80				10	0 100	100					100	100	100
Serratia marcescens	4	75	0	100				0	0		100			100	100				10	0 100	100							
Enterobacter cloacae	2	100	0	50				50	50	50	50	50	50	100	50				10	0 100	100					100		100
Klebsiella oxytoca	2	50	100	100				100	100	100	100	100	100	100	100				10	0 100	100					100		100
Staphylococcus aureus	9	13		0	0	0	14								75	100	33	100	Г			100	100	100	100	100		
Staphylococcus epidermidis	4	25		25	25	50	25								25	25	25	100				100	100	100	100	0		
Enterococcus faecium	3			0	0	0											0	67				67	100		100	0		
Staphylococcus haemolyticus	3	67		0	0	0	0								0	50	0	67				100	100	100	100	0		
Sphingomonas paucimobilis	2	50		50	100						100	100	100	100	50					50	100							

Reserved/ Restricted Drugs:Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Prepared by

Checked by

Verified by

Approved by

Dr. M Kumar

Prof & HoD

Prof Microbiology

Microbiology

Ochecked by

Verified by

Approved by

Dr. SK Ialwani

Medical Director



2025-2026

Version - 11.0

Percentage Susceptibility Gram Negative Organism 2024

L	ocation	Specimen type	Organism	Number of patients	Ciprofloxacin	Cotrimoxazole	Amox/Clav	Cefuroxime	Cefuroxime axetil	Ceftriaxone	Cefepime	Cef/Sul	Piptaz	Amikacin	Gentamicin	Ertapenem	Meropenem	Imipenem	Tigecycline	Colistin
		Dland	Klebsiella pneumoniae	13	46	69	50	39	33	54	62	69	67	92	100	75	75	75	100	100
	NICU	Blood	Acinetobacter baumannii	11	27	64				36	30	30	36	27	36		30	22		100

Reserved/ Restricted Drugs:Not to be used emperically unless justified
Will be useful clinically
Will be useful clinically in about 2/3 cases
Will not be useful clinically
Not tested/not appropriate antibiotic

Candida Isolates:

Candida Isolated in Neonatology

Location	Specimen type	Organism	Number of patients	Fluconazole	Voriconazole	Ampho B	Caspofungin	Micafungin
	Blood	Candida albicans	1	100	100	0	100	100
Nicu	blood	Meyerozyma guilliermondii	2	0	0	100	100	100
INICU	Urine	Candida albicans	1	100	100	0	100	100
	ornie	Candida tropicalis	1	100	100	100	100	100

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

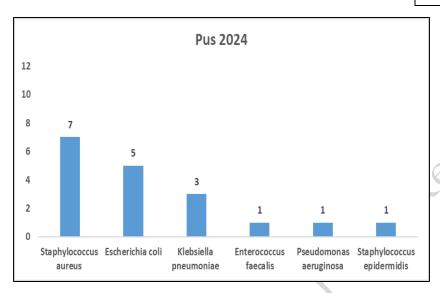
			/ .
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shipa Acharya Quality Department	Dr. SK Nalwani Medical Nirector

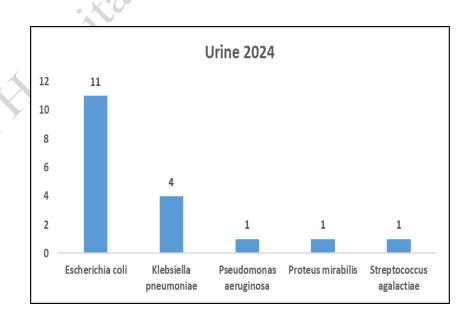


2025-2026 Version - 11.0

Obstetrics And Gynaecology: Top 10 Isolates Sample wise

Specimen	Overall No. of
Type	Isolates
Pus	18
Urine	18





			/ .
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Smipa Acharya Quality Department	Dr. SK Italwani Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Obstetrics And Gynaecology Antibiogram:

Percentage Susceptibility All Organism 2024

Organism	Number of patients	Nitrofurantoin	Ciprofloxacin	Levofloxacin	Moxifloxacin	Oxacillin	Penicillin	Cotrimoxazole	Amox/Clav	Cefuroxime	Cefuroxime axetil	Ceftazidime	Ceftriaxone	Cefotaxime	Cefepime	Cef/Sul	Piptaz	Amikacin	Gentamicin	Clindamycin	Erythromycin	Tetracycline	Teicoplanin	Linezolid	Daptomycin	Vancomycin	Ertapenem	Meropenem	Imipenem	Tigecycline	Fosfomycin	Colistin
Escherichia coli	20		10					55	55	5	6		21	0	32	0	74	85	85		100						94	94	94	100	100	100
Klebsiella pneumoniae	7		43					57	57	43	50		43	0	57	0	57	100	86								86	86	71	100	0	100
Pseudomonas aeruginosa	2		100	100								100			100	0	100	100	100									100	100			100
																					\vee											=
Staphylococcus aureus	9	100	0	0		67	0	56											89	89	44	100	100	100	86	100				100		
Enterococcus faecalis	2	100	50	50			100														0	0	100	100	100	100				100		
Streptococcus dysgalactiae	1			100	0		100						100	100						100	100	0		100		100				100		
Staphylococcus epidermidis	1	100	0	0		0	0	0											100	100	0	0	0	100	100	100				0		
Streptococcus agalactiae	1	100		0			100															0		100		100				100		
Staphylococcus warneri	1	100	100	100		100		100											100	100	100	100	100	100	100	100				0		

Reserved/ Restricted Drugs: Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

			/
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shipa Acharya Quality Department	Dr. SK Nalwani Medical Nirector

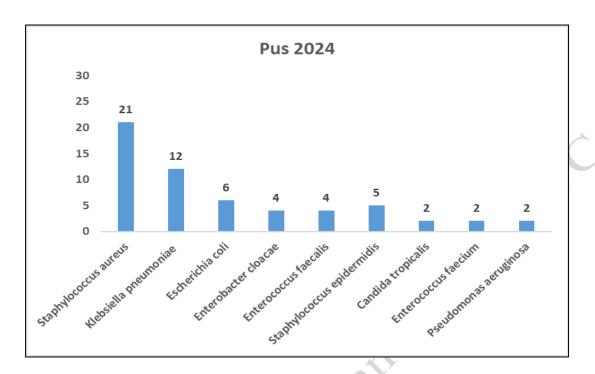


Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Orthopedics:

Top 10 Isolates Pus (n = 60)





2025-2026

Version - 11.0

Orthopedics Antibiogram:

Percentage Susceptibility Gram Negative Organism 2024

Location	Specimen type	Organism	Number of patients	Nitrofurantoin	Levofloxacin	Ciprofloxacin	Norfloxacin	Cotriamoxazole	Amox/Clav	Cefuroxime	Cefuroxime axetil	Ceftazidime	Ceftriaxone	Cefotaxime	Cefepime	Cef/Sul	Piptaz	Amikacin	Gentamicin	Ertapenem	Meropenem	Imipenem	Tigecycline	Fosfomycin	Minocycline	Colistin
		Escherichia coli	29			7.7		49	54	18	21		26		28	74	82	81	74	90	86	85	100			100
	Pus	Klebsiella pneumoniae	18			39		54	46	46	47		50		50	58	56	58	68	62	59	65	100			100
		Pseudomonas aeruginosa	13		63	67		50		100		90			94	79	89	90	80		89	84			100	100
Ortho		Escherichia coli	32	85	0	15	44	50	52	20	18	59	28	14	35	69	75	86	75	90	88	88		99	100	100
	United	Klebsiella pneumoniae	10	24		38	75	48	47	33	32	73	40	43	42	46	51	67	64	63	54	50				100
	Urine	Pseudomonas aeruginosa	15		26	33	100					46			47	35	80	58	46		37	40				100
		Enterobacter cloacae	13	0		85	67	85	8	60	50	67	62		100	100	85	92	92	85	100	100				100

Reserved/ Restricted Drugs: Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Percentage susceptibility Gram Positive organism 2024

Lo	cation	Specimen type	Organism	Number of patients	Cotrimoxazole	Penicillin	Oxacillin	Ciprofloxacin	Levofloxacin	Gentamicin	Clindamycin	Erythromycin	Tetracycline	Daptomycin	Tigecycline	Teicoplanin	Linezolid	Vancomycin
			Staphylococcus aureus	21	38	13	75	0	0	63	88	63	71	71	100	100	100	100
_		Dura	Enterococcus faecalis	4		100		33	50			0	0	100	100	100	100	100
O	rtho	Pus	Staphylococcus epidermidis	4	0	0	0	100	100	100	100	0	100	100	0	100	100	100
			Enterococcus faecium	2				0	0			0	100		0	100	100	100

Reserved/ Restricted Drugs: Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Prepared by

Checked by

Verified by

Approved by

Dr. M Kumar

Prof & HoD

Prof Microbiology

Microbiology

One M Checked by

Verified by

Approved by

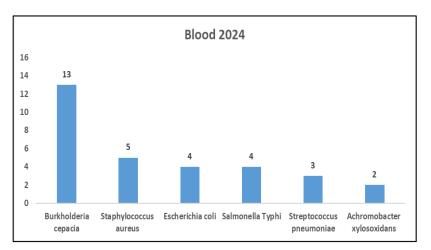
Dr. SK Latwani

Medical Director

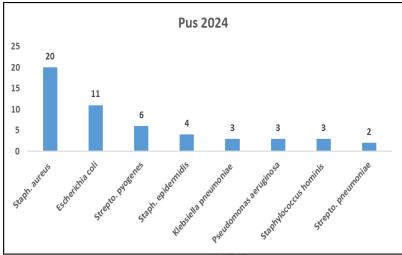


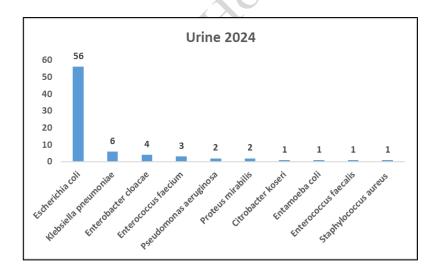
2025-2026 Version - 11.0

Paediatrics: Top 10 Isolates Sample wise



Specimen	Overall No
Туре	of Isolates
Blood	42
Pus	57
Urine	77





	Checked by	Verified by	Approved by
Prepared by		verified by	Appleted by
De	Dr. M S Modak	Jehary	4
Dr. M Kumar	Prof & HoD	Dr. Shilpa Acharya	Dr. SK Laiwani
Prof Microbiology	Microbiology	Quality Department	Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Paediatrics Antibiogram:

Percentage Susceptibility Organism -Peds 2024

Organism	Number of patients	Nitrofurantoin	Cotrimoxazole	Penicillin	Oxacillin	Ampicillin	Norfloxacin	Ofloxacin	Ciprofloxacin	Levofloxacin	Moxifloxacin	Amox/Clav	Cefuroxime	Cefuroxime axetil	Ceftazidime	Ceftriaxone	Cefotaxime	Cefepime	Cel/Sul	Piptaz	Amiacin	Gentamicin	Clindamycin	Erythromycin	Tetracycline	Ertapenem	Imipenem	Meropenem	Tigecycline	Minocycline	Fosfomycin	Colistin	Daptomycin	Teicoplanin	Vancomycin	Linezolid
Escherichia coli	72	80	51			56	64	67	11			57	12	11	55	23	0	22	83	83	89	86				94	93	96	100		100	100				
Burkholderia cepacia	13		83						0	75					100													91		88						
Klebsiella pneumoniae	11		55						55			73	50	50		64	0	64	73	73	82	73				82	73	82	100			100				
Enterobacter cloacae	6	0	100				100	100	50			0	20	20	100	50	0	60	83	83	100	50		100		100	100	100	100			100				
Pseudomonas aeruginosa	5								100	100					100			80	100	100	100	100					75	75				100				
Salmonella Typhi	4		75						0			0				100		0								100	100	100	100							
Staphylococcus aureus	27	100	50	4	48				15	17												82	82	59	100				100				100	100	96	100
Streptococcus pneumoniae	6		100							100	100					100	100						20		33				100						100	100
Streptococcus pyogenes	6			100		100				40	0					100	100						100	40	100				100						100	100
Staphylococcus epidermidis	5	100	80	33	40				80	75												100	100	25	100				0				100	100	100	100
Enterococcus faecium	3	0		0					0	0														0	0				0					100	100	100
Staphylococcus hominis	3		50	0	0				100	100												100	50	50	100				0					100	100	100
Enterococcus faecalis	2	100		100					50	50														0	0				100				100	100	100	100

Reserved/ Restricted Drugs: Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not approxitrofurantoin for urinary isolates only

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

			/
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Nalwani Medical Nirector



2025-2026

Version - 11.0

Paeds ICU Antibiogram:

Percentage Susceptibility Organism - PICU 2024

Organism	Number of patients	Cotrimoxazole	Amox/Clav	Penicillin	Ciprofloxacin	Levofloxacin	Cefuroxime	Cefuroxime Axetil	Ceftazidime	Ceftriaxone	Cefotaxime	Cefepime	Cef/Sul	Piptaz	Amikacin	Gentamicin	Clindamycin	Erythromycin	Tetracycline	Ertapenem	Imipenem	Meropenem	Teicoplanin	Tigecycline	Fosfomycin	Linezolid	Vancomycin	Colistin
Escherichia coli	12	33	36		27		27	22		27		40	0	73	91	83			100	63	78	78		100	100			100
Pseudomonas aeruginosa	12				50	55	33	0	50	0		50	0	63	60	0					42	42						100
Klebsiella pneumoniae	11	46	73		27		27	30		36	0	55	0	46	82	82				73	73	73		73				100
Acinetobacter baumannii	3	100			33					50		33	0	33	33	67					33	33						100
Burkholderia cepacia	4	100			0	33			67	0		0	0									75						
Enterobacter cloacae	2	100			100		100	100		50		100	0	100	100	100				100	100	100		100	0			
Staphylococcus aureus	16	31		0	19	21										94	94	56	100				100	100		100	100	
Streptococcus pneumoniae	6	60		100		83				100	100							17	40					100		100	100	

Reserved/ Restricted Drugs:Not to be used emperically unless justified

Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/not appropriate antibiotic

Please Note: Individual Isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Prepared by

Checked by

Verified by

Approved by

Dr. M Kumar

Prof & HoD

Prof Microbiology

Microbiology

Ochecked by

Verified by

Approved by

Dr. Sk Ialwani

Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Antimicrobial therapy in hospitalized patients

Antimicrobial therapy is used in hospitalized patients in three situations

- 1. **Empiric therapy** before the causative organism has been identified
- 2. **Definitive therapy** once the causative organism is identified
- 3. **Prophylactic therapy** to prevent infection, eg. surgical prophylaxis

Empiric therapy

Patient requiring empirical antimicrobial therapy should be classified into three types (Table 1) depending on the past history, prior exposure to health care, previous antimicrobials and associated comorbidities. Antimicrobial should then be chosen according to the site of infection and suspected microorganism based on local hospital microbiologic data (antibiogram). Appropriate cultures must be sent prior to antimicrobial therapy. Identification of the micro-organism will then dictate definitive therapy and also contribute to the hospital antibiogram for choosing empiric therapy.

Please note: Empiric therapy will be started only after appropriate samples for culture have been taken

TABLE 1- Patient Types for selecting empiric antimicrobial therapy

Patient Type 1	Patient Type 2	Patient Type 3
(Community acquired)	(Healthcare associated)	(Nosocomial Infections)
	Contact with health care system (e.g. recent hospital admission, nursing home, dialysis) without invasive procedure, within	
No contact with health	invasive procedure within last90 days. Current	Current hospitalization>7 days. Invasive procedures within last 90
care system	hospitalization less than 7 days	days
	40	Recent & multiple antimicrobial
No prior antimicrobial	Recent antimicrobial therapy	therapies within last 90 days Major
treatment	(within last 90 days)	invasive procedures done
No procedures done	Minimum procedures done	
		Cystic fibrosis, structural lung
Patient young with only a	Patient old with Multiple co-	disease, advanced AIDS, neutropenia
few co-morbid conditions.	morbidities.	other Severe immunodeficiency

Dunnamed by	Checked by	Verified by	Approved by	
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Smipa Acharya Quality Department	Dr. SK Laiwani Medical Director	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 37 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

6.Empiric Antimicrobial choice

Recommended antimicrobials for common conditions are listed below. This guide is a broad outline; not all-inclusive and; not meant to replace treating physician's judgment.

Table 1: Acute gastroenteritis

Name of condition	Patient Type 1	Patient Type 2 (Healthcare associated)	Patient Type 3 (Nosocomial Infections)
	(Community acquired)	(Healthcare associated)	(Nosocomial Infections)
Acute gastroenteritis	Most cases are self-		
	limited and require only		
	supportive treatment		
	' ') 7
	and hydration or being		
	viral in nature. Selected		
	sick patients can be		
	treated as per following		
	guidelines.		
	-		
	Tab Cefixime 200mg BD	A >	
	If at a all assessing at a se		
	If stool examination		
	shows		
	invasive diarrhoea (> 5		
	leucocytes /HPF or		
	blood in the stool).		
	Then consider		
	stool culture followed		
	by therapy as per AST		

Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Ouality Department	Dr. SK Natwani Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 2: Pneumonia

Name of condition	Patient Type 1	Patient Type 2	Patient Type 3
	(Community acquired)	(Healthcare associated)	(Nosocomial Infections)
Pneumonia	1] For non-ICU	Late Onset	Late onset HAP/VAP
	patients with	HAP/VAP (For more	suspected MDR Gram
	community acquired	than 48 hours of	negative –
	pneumonia	hospitalization but less	
	(CAP)Ceftriaxone (2g	than 7 days)	Imipenem (0.5-1 gm q6h
	IV q24h X 5-7 days)/		/Meropenem (1-2 g IV
	Amoxicillin/Clavulanic	If septic shock or	q8h)
	acid (1.2g q8h IV)	multisystem organ	Suspected XDR Gram
	\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	failure, Imipenem0.5-	negative
	+	1gm q6h or	.(3)
		Meropenem1-2 gm	Polymixin B 15 lac loading
	Macrolide	q8h	dose f/b 7.5 Lac mg BD
	(Azithromycin- 500mg	qon	Suspected MRSA-
	IV/PO once a day), x 5-7		Vancomycin (1g IV q12h
	days).		OR Teicoplanin (400mg
		A Y	IV for 3 doses, then
			q24h)
	2] ICU patients with		<u> </u>
	CAP		For suspected VRE-
			Linezolid (600mg IV/PO
	Ceftriaxone (2g IV q24hr		q12hr)**x 7-14 days
	X 5-7 days)/		For suspected Fungal
			(Filamentous
	Amoxicillin/Clavulanic		fungi/mould) infections-
	acid (1.2g q8hr IV)		infections-
	× V		Consider Antifungals in
6	+		Immunocompromised host.
	Macrolide: Azithromycin-		Add Liposomal
	500mg IV/PO q24h)/		Amphotericin B.
	Doxycycline 100mg PO		Substitute Voriconazole,
	q12h x 5-7 days).		if Aspergillus suspected
			on radiological evidence
	If aspiration is suspected		or galactomannan positive
	clindamycin 600mg q8h		
	Early onset HAP/VAP		If PCP suspected-
	·		add TMP-SMX or
	(less than 48 hours	<u> </u>	Clindamycin

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

Dr. Skilpa Acharya

Quality Department

Medical Prector



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

	<u> </u>	<u> </u>
	admission) Antimicrobial choice as	
	above unless	
	Pseudomonas or Gram	
	negative bacilli are	
	suspected. Then use	
	Cefoperazone-	
	Sulbactam* (1.5g-3gm	
	q6h) or piperacillin–	
	tazobactam (PIP-Taz)	
	4.5gm q6h	
	Look for typical viral	
	symptoms such as	
	sneezing and running	
	nose.	
	If fever, sore throat ,dry	2
	cough and viral	A . Y
	symptoms present, initiate	200
	Oseltamivir 75 mg BD x	
H1N1	5 d without waiting for	
Flu-like	illness confirmation by PCR	1

Note:

- 1. Fluoroquinolones should not be used for empiric treatment.
- 2. Fluoroquinolones should not be used routinely for treating Acute exacerbation of COPD
- 3. In the uncommon scenario of hypersensitivity to β-lactams, respiratory Fluoroquinolones (e.g. levofloxacin 750 mg daily) may be used if tuberculosis is not a diagnostic consideration at admission. Patients should also undergo sputum testing for acid-fast bacilli simultaneously if fluoroquinolones are being used in place of β-lactams.
- 4. **Patients with suspected MRSA infection, we recommend the use of empiric Vancomycin or Teicoplanin. The use of linezolid in India should be reserved because of its potential use in extensively drug-resistant tuberculosis.
- 5. Suspected viral pneumonia [influenza] Oseltamivir and/or Zanamavir should be given. It should be given within 48 hrs of onset of symptoms otherwise there is no clinical benefit.
- 6. In late HAP/VAP with suspected Acinetobacter infection combination of Colistin + carbepenem / sulbactam.
- 7. Duration of treatment for community acquired pneumonia should be minimum 5-7 days and patient should be afebrile 48-72 hours prior to stopping treatment.

Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Smipa Acharya Quality Department	Dr. SK Natwani Medical Nirector	

Version - 11.0



Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026

- 8. For ESBL / MRSA health care associated pneumonia minimum duration of treatment should be 10-14 days.
- 9. For proven pseudomonal / Acinetobacter health care associated pneumonia treatment should be for minimum 2 weeks and preferably combination of antimicrobial therapy should be used.
- 10. Colonization should be suspected if respiratory secretions culture show growth but following features are absent like Fever, leukocytosis, increased bronchorrhea, increasing oxygen requirement, new lung infiltrates.
- 11. In presence of Fever, leukocytosis, increased bronchorrhea, increasing oxygen requirement but absence of lung infiltrates with positive cultures [MDR GNB / MRSA] to be treated as health care associated tracheobronchitis with appropriate broad spectrum antimicrobials.
- 12. Aerosolised Tobramycin/Colistin can be added to IV antimicrobials as an adjunctive and Rese therapy for MDR gram negative infection with specialized nebulizers.

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

Dr. Ski laiwani

Medical Birector



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 3: Meningitis

Name of condition	Patient Type 1	Patient Type 2	Patient Type 3
	(Community acquired)	(Healthcare associated)	(Nosocomial Infections)
Meningitis	1] Age 2yrs-50yrs Vancomycin 1gm q12h + Ceftriaxone 2gm q12h 2] Age > 50yrs	Vancomycin 1gm q12h + cefepime 2gm q12h /Ceftazidime 2gm q8h	Empirical Therapy Vancomycin 1gm q12h + Colistin 9 MIU loading f/b 4.5 MU BD+/- Meropenem 2gm q8h. Consider
	Above Antimicrobials + Ampicillin 2gm q4h		Intrathecal Gentamicin/ Colistin 4.5 MIU BD
		a esect	Organism specific A] Suspected MRSA Meningitis — Vancomycin 1gm q12h +/- Rifampicin 600mg q12hor Linezolid 600mg q12h
			B] ESBL Gram negative/Pseudomonas or Acinetobacter (MDR / XDR) Meropenem 2gm q8h + Colistin 9 MIU loading f/b 4.5 MIU BD.

Note: Intrathecal/Intraventricular route dosage-

Vancomycin 10-20mgq24h;

Gentamicin 4-8 mgq24h;

Amikacin 30-50mg q24h;

Colistin 5-20mg q24h[1mg = 12,500 units]

IV Dexamethasone should be given in suspected pneumococcal meningitis before antimicrobial therapy and should be continued only if Gram stain / Culture confirms pneumococcal etiology

				/ '
Prepared by	Checked by	Verified by	Appr	oved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Smipa Acharya Ouality Department		K Latwani

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 42 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 4: Urinary tract infection

Name of	Patient Type 1	Patient Type 2	Patient Type 3
condition	(Community acquired)	(Healthcare associated)	(Nosocomial Infections)
UTI	Asymptomatic	Complicated UTI	Complicated UTI and
	bacteriuria	(Obstruction, reflux,	Pyelonephritis
	No empirical therapy.	azotemia, CAUTI)	(Suspected MDRO's/
	Send C/S.	IV Meropenem 1gm q8h/	Post renal transplant/
		IV Imipenem-cilastatin0.5	Recurrent UTI's)
	Non complicated UTI	gm q6h	IV Meropenem 1gm
	(Cystitis, Urethritis, No		q8h/IV Imipenem-
	evidence of obstructive		cilastatin 500mg q6h
	uropathy)	, C	+/-
	PO TMP SMX 160/800		IV Colistin 9 MIU loading f/b 4.5 MIU BD
	q12h / PO Nitrofurantoin		loading 1/0 4.5 Mile BD
	100 mg q12h		
	100 mg q12m		If MRSA or
	Acute Uncomplicated	Complicated	enterococcus,
	Pyelonephritis	Pyelonephritis	ConsiderVancomycin 1
	1 yelonephitels	1 yelonepin tus	gm q12h/ Teicoplanin
		(Obstruction, reflux,	400 mg 3doses f/b
		azotemia, CAUTI, Shock,	400 mg q24h
	Gentamicin 3 – 5 mg/kg	perinephric abscess)	
	q24h 5-7 days	Meropenem 1gm	
		q8h/Imipenem-cilastatin	look for obstruction.
A	Ceftriaxone 1gm q12h	0.5 gm q6h in presence of obstruction	Surgical management is mandatory to relieve obstruction
		q8h/Imipenem-cilastatin 0.5 gm q6h in presence of	Surgical managemen mandatory to relieve

			/	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Smipa Acharya Ouality Department	Dr. SK Latwani Medical Director	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 43 of 81

UNIVERSITY

BHARATI VIDYAPEETH UNIVERSITY MEDICAL COLLEGE HOSPITAL AND RESEARCH CENTRE

Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026 Ven

Lower Urinary tract	OPD- Cap.Amoxicillin500 mg		
infection(UTI) in	q8h PO		
antenatal patients up to 20 weeks	In-patient		Meropenem1gm q8h
gestation	IV Ceftriaxone 1gm q12h		Or Colistin 4.5 MIUBD
Lower Urinary	OPD		
infection(UTI) in	Tab. Nitrofurantoin		
	SR100 mg BD oral/ Oral		
antenatal patients	Fosfomycin 1g sachet		
after 20 weeks	single dose/Oral Cefixime		
gestation	200mg BD x 3-5 days		
	Inpatient: Ceftriaxone	^	\sim
	1gm q12h	IV PIP-TZ 4.5 gm q6h	Meropenem 1gm q8h

Table 4 a : Pelvic Inflammatory Disease

		Levofloxacin Plus
		Metronidazole or
Pelvic inflammatory	Tab Cefixime	Ceftriaxone with
disease: Mild to	Plus Tab Metronidazole	doxycycline +/-
moderate	Plus Cap Doxycycline	metronidazole
Pelvic inflammatory	Clindamycin Plus	IV PIP-TZ 4.5 gm q6h or
disease: Severe	Gentamicin	Imipenem

			,	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Ouality Department	Dr. SK Nalwani Medical Nirector	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 44 of 81

Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Table 5- Skin & soft tissue infections

Name of condition	Patient Type 1 (Community acquired)	Patient Type 2 (Healthcare	Patient Type 3 (Nosocomial Infections)
Erysipelas / uncomplicated	IV Ceftriaxone 2 gm q24h	associated)	
cellulitis	If beta lactam allergy		
	IV Clindamycin 600 –		
	900 mg q8h		
Necrotizing	IV Ceftriaxone 2gm q12h		70'
infection of	+		
skin/fascia and	IV Clindamycin 600-		
muscle	900mg q8h / IV		
	Metronidazole 500mg		
	q6h		
	or		
	Inj Piperacillin-		
	Tazobactam +/- Inj		
	clindamycin If Suspected MRSA	2 eseati	
	IV Vancomycin1 gm	2	
	q12h/ IV Teicoplanin		
	400 mg q24h		
	Mixed aerobic and		
	anaerobic cover		
	including S.aureus		
	MRSA cover IV		-
	Vancomycin1gm q12h		
	If pseudomonas		
	suspected IV PIP-TZ		
Fournier gangrene	4.5gm q6h		
X			IV Meropenem 1gm q8h or
	_	© 1	IV Imipenem-Cilastatin
	q8h	If Suspected MRSA	1gm q6h. If MRSA
	if beta lactam allergy- IV		suspected IV Vancomycin
Diabetic foot	Clindamycin 600 q8h	Vancomycin1 gmq12h	1gm q12h

				/ '
Prepared by	Checked by	Verified by	Appr	oved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Smipa Acharya Ouality Department		K Latwani

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 45 of 81

Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 6- Bone and joint infections

Name of	Patient Type 1	Patient Type 2	Patient Type 3
condition	(Community	(Healthcare	(Nosocomial Infections)
	acquired)	associated)	
Acute	Ceftriaxone IV	-	-
Osteomyelitis	q12h OR Co-amoxiclav		
/ Septic	1.2 gm q8h with/without		
	Gentamicin 3 –5 mg/kg		
Arthritis	q24h		×
	If MRSA		
	suspected-		÷ (2)
	Vancomycin1gm		
	IVq12h		
		Usual Suspected	
		organism- Staph	C
		aureus/ MRSA	
		IV Vancomycin 1	
		gm q12h/	
		Teicoplanin(400mg	
		IV q12h for 3 doses, then q24h)	
		doses, then $q2411$	
		If Suspected MDR	
		Gram negative	
		organism	
Early	^	IV Meropenem	
implant	×2	1gm q8h	
associated		IV Imipenem +	
infection (< 3	.50°	Cilastatin 1gm q6h	
months)	- 65	IV/IV Colistin	-
	140		Usually low grade infection
	\\\\\		If Coagulase negative
	Y Y		staphylococcus suspected -
			IV Vancomycin 1 gm q12h /
Late implant	.0		Teicoplanin (400mg IV q12h for 3
associated	Y		doses, then q24h)
infection			If Anaerobe (Propionibacterium
(after 3			acne) suspected
months)	-	-	IV Clindamycin 600-900 mg q8h.

			/ '	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Naiwani Medical Director	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 46 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Table 7 Intra-abdominal infections -

Name of condition	Patient Type 1(Community	Patient Type 2	Patient Type 3
	acquired)	(Healthcare associated)	(Nosocomial Infections)
A) Extra – biliary	IV Ceftriaxone1-2 gm	IV Meropenem 1gm	IV Meropenem 1gm q8h IV
	q12h+IV	q8h/ IV Imipenem-	lmipenem -
	Metronidazole500mg q8h or	cilastatin 500mg q6h	cilastatin500mg q6h In
	IV PIP-TZ 4.5gm q6h		case of suspected
			Acinetobacter or XDR
			Gram negative organisms
			Colistin 4.5 MU BD If
			MRSA or Enterococcus
			suspected IV Vancomycin
			1 gm q12h /
			Teicoplanin(400mg IV
			q12h for 3 doses, then
			q24h) If VRE suspected
		AC.	Linezolid 600 mg IV q12h
			suspected, Add
			Fluconazole 400 mg IV
		5	q24h If non albicans
			Candida- IV Caspofungin
			70 mg stat and 50 mg
			q24h or Ampho B
B)Intra Abdominal	IV Ceftriaxone1-2 gm q12h +	IV Meropenem 1gm q8h	Eg- Acute cholangitis
Biliary	IV Metronidazole500mg q8h	/ IV Imipenem- cilastatin	following bilio enteric
	or IV PIP-TZ 4.5gm q6h	500mg q6h	anastomosis IV
			Meropenem 1gm q8h/ IV
			lmipenem -
			cilastatin500mg q6h .If
			MRSA or Enterococcus
			suspected IV Vancomycin
			1 gm q12h /
			Teicoplanin(400mg IV
	Y		q12h for 3 doses, then
	• *		q24h). If VRE suspected
	X >		Linezolid 600 mg IV q12h
2			If Fungal Infection
			suspected, Add
42.0			Fluconazole 400 mg IV
23hara			q24h
\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\			If non Albicans Candida IV
Y			Caspofungin 70 mg stat and
Noter: Metnonidszole	doeing based pr Phetroidek in	atiilistu idieslis ula 5 Cu nsa 24 hr	50 mg q34h Qr.Ampha.Bn

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

Prof Microbiology

One of the control of the

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 47 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 8: Infective Endocarditis

Name of condition			
Native Valve	IV Ceftriaxone	Alternative	
Endocarditis		Penicillin G2-3mu IV q4h or Vancomycin500 mg q12h for 4weeks	
		Ceftriaxone 2 gmq24h for 2 weeks plus	
		Gentamicin 3mg per kg divided into equal doses	
		q8h for 2 weeks	
Prosthetic valve	Cloxacillin 2gm IV q4h	IV Cefazolin 2g q8h	
endocarditis	for 4-6 weeks or IV		
	Vancomycin500 mg	$\nearrow \emptyset$	
	q12h for 4-6 weeks		

Note:-

If Penicillin resistant Streptococci - Ceftriaxone 2 gram per day IV q24h for 6 weeks plus Gentamicin 3mg per kg divided into equal doses q8h for 6 weeks

Enterococci – Ampicillin 2gm IV q4h + Gentamicin3mg per kg divided into equal doses q8hboth 4-6 weeks or Vancomycin 500 mg q12h + Gentamicin for 4weeks.

Staphylococci –Nafcillin or Oxacillin 2gm IV 4 hourly for 4-6 weeks or Vancomycin 15 mg /kg IV 12 hourly for 4-6 weeks

			/ `
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Natwani Medical Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 48 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 9: Malaria, Leptospirosis, Scrub Typhus, Enteric fever (IN LABORATORY CONFIRMED CASES)

Plasmodium Vivax		
Malaria	Chloroquine Sensitive	Chloroquine resistant –
		any of the ACT therapy excluding
	Chloroquine (10mg base/kg stat followed	SP
	by 5 mg/kg at 12,24,36 hours) plus	1. Artesunate +Amodiaquine
	Primaquine (7.5 mg (base) q12h PO	2. Artesunate +Mefloquine
	x14days) (Primaquine should not be given	3. Dihydroartemisin plus
	in severe G6PD deficiency)	piperaquine

Plasmodium Falciparum	OPD	Drug resistant Falciparum
Malaria	Artesunate(2.4 mg/kg at 12 & 24 hours)	Malaria
	plus	Artesunate 2.4 mg/kg for 7 days
	Sulfadoxine (25 mg/ kg) & Pyrimethamine	
	(1.25 mg/kg) as a single dose or	or Change (Total gray 12 2 2 101)
	Artesunate(same dose as above) plus	
	Amodiaquine(10mg) base per kg OD for	
	3 days (Fixed dose combinations are	
	available) or Artemether plus	
	Lumefantrine(1.5/9mg/kg BD for 3 days)	days plus one of the following
	Drug combination of A+L(mg)available	three
	40+240:60+360:80+480 or Artesunate	1. Tetracycline 4mg/kg Odx7
	+Mefloquine (25mg base/kg –total)	days
	(8mg/kg once a day for 3 days)	2. Doxycycline 3mg/kg OD x
	Hospitalized patient	7days
	Artesunate IV 2.4 mg/kg at 12 & 24 hours	3. Clindamycin 10mg/kg BD x
	and 2.4 mg/kg q24h X 5 days +	7days
	Doxycycline 100mg q12h x 7 days	·
		Alternative: Amoxicillin (500 mg)PO
Leptospirosis (Mild)	Doxycycline 100mg q12h x 7 days	TDS x7 days
^	X	Ampicillin (500mg)PO TDS x 7 days
Leptospirosis (Moderate	Ceftriaxone (1gm 12 hourly x7 days or	Alternative
or Severe)	Cefotaxime (1gm 6 hourly IV x 7 days	Penicillin (1.5 million units /IV
		/IM 6 hourly x7 days
	Doxycycline (100mg) BD x 7 to 15 days	
Scrub Typhus	or	Alternative
	Azithromycin (500mg) OD x 3days	Chloramphenicol (500mg)QID
		x7-15 days
Enteric Fever (OPD)	T. Cefixime 400 mg TDS for 14 days	Alternative
		T. Azithromycin (1gm)OD for 5
		days
Enteric Fever(IPD)	Ceftriaxone (4gm/day)IV for 7-14 days	

	1	1,,	
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Laiwani Medical Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 49 of 8



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 10 Pediatric Infections

Name of	Patient Type 1 (Community		
condition	acquired)	Patient Type 2	Patient Type 3
Condition	acquireu)	1 attent Type 2	1 attent Type 3
Pneumonia	Community acquired Pneumonia	Either Type II or Early HAP/VAP	Either Type III or late HAP/VAP, IV Meropenem (60-120 mg /kg/day
AGE: 3			
weeks	Ceftriaxone100mg/kg/d	Piperacillin-	divided 8 hrly) plusVancomycin
to 3 months	od or Cefotaxime	tazobactam 300	(40-60 mg/ kg/ day divided 6-8 hrly
	150mg/kg/d tds x 10-14	mg/kg/d qid	
	days and		IV Meropenem (60-120 mg /kg/day
	*Azithromycin		divided 8 hrly) plusVancomycin
	10mg/kg/day x		(40-60 mg/ kg/ day divided 6-8 hrly.
			Add Fluconazole 6-12 mg/kg/day or amphotericin B (if renal
		0	dysfunction)
		S	aystanedon)
AGE: 4	Lobar	Piperacillin-	
months to 5	pneumonia/effusion	tazobactam 300	
years	Ceftriaxone	mg/kg/d	Same as above
	100mg/kg/d od with	Qid plusVancomycin	
	Cloxacillin 100-	(40-60 mg/ kg/ day	
	200mg/kg/d	divided 6-8 hrly	
	~?	Y	
	Bronchopneumonia without		
	effusion Ampicillin		
	200mg/kg/d qid		
	days*consider adding	Ceftriaxone100mg/kg/	
		d od Or Piperacillin-	
	1	tazobactam 300	
		mg/kg/d qid Either type II/post	
		neurosurgical	Either type II/III or post shunt
Meningitis		meningitis	infection
	7	9 * **	IV Meropenem (120 mg /kg/day
	Cefotaxime 200 mg/kg/d		divided 8 hrly)/
7	qid/or Ceftriaxone100mg/kg/d		plusVancomycin60mg/kg/d qid with
	our our prins	IV Meropenem (120	or without rifampin 10 mg/kg (PO)
	, 4111 6111 661118, 118, 4	mg /kg/day divided 8	q12h x 7-10 days after shunt removal
	qid*Discontinue Vancomycin	nriy) plusVancomycin(60	Consider additional Intraventricular therapy Vancomycin 10mg or
	1 20	mg/ kg/ day divided 6	Gentamicin 1-2 mg or Polymixin B
Age > 3	e i	hrly +/- rifampin 10	2mg or Colistin/10mg $[1mg = 12,500]$
months d by	meningitidis, 55kH. Mfluenzae	• , •	units] Approved by
Urinary Tract	Dr. M. S. Modak	Achary	
Dr. M Kumar Prof Microbiolog	Prof & HoD	Dr. Shilpa Acharya Quality Departmen	Dr. SK Jalwani Medical Director
	Policy Var 11 Data of release: 01 0/		

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 50 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026 Version - 11.0

PLNE	11111111	nerobiar roncy and Antinnerobiar St	ewarusiip 110graii 202	5 2020 Version 11:0			
Infectio	n						
Cystitis		Co-trimoxazole 8-10 mg/kg/d of trimethoprim bd OR Amoxy-clav 30-40 mg/kg/d bd OR Cefixime 8-10 mg/kg/d od					
Pyelone	ephritis	Ceftriaxone100mg/kg/d od OR Cefotaxime150mg/kg/d tds OR Piperacillin- tazobactam 300 mg/kg/d tds/qid +/- Amikacin 15- Piperacillin- tazobactam 300 mg/kg/d	Piperacillin- tazobactam300 mg/kg/d tds/qid Or	Same as for type II			
		l	Meropenem120mg/kg/d	Same as for type II			
		2011g/Rg/4 04 7110 17 44ys	<u>u</u>				
	20mg/kg/d od A10-14 days d						

Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Natwani Medical Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 51 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2

2025-2026 Version - 11.0

Cloxacillin 200mg/kg/d	Infections	200mg/kg/d	Piperacillin-	IV Meropenem (120 mg /kg/day
Infections Orbital Ceftriaxone 100mg/kg/d od/bd x 10-14 days		nlang aidh an Cafadaani		
Orbital cellulitis		DILLE ALTHAR L'ATATAVIMA	tazobactam 300	• • •
CellulitisCeftriaxone 100mg/kg/d od/bd x 10-14 daysBone and JointCloxacillin 200mg/kg/dVancomycinIV Meropenem (120 mg /kg/da divided 8 hrly)/ plusInfectionsplus either Cefotaxime 150mg/kg/d tds or60mg/kg/d qid or Clindamycin 20-40 mg/kg/d tds/qidVancomycin 60mg/kg/d tds/qidAcuteCeftriaxone 100mg/kg/d od/bd + Vancomycin x eptic arthritisCeftriaxone 10-14 daysPlus eitherOsteochondritisPiperacillin- tazobactam300 mg/kg/d tds/qid or combination therapyPiperacillin- tazobactam300 mg/kg/d tds/qid or combination therapy	Orbitai	-		1
Bone and Joint Cloxacillin 200mg/kg/d Vancomycin IV Meropenem (120 mg /kg/d adivided 8 hrly)/ plus Vancomycin Somg/kg/d qid or Clindamycin 20-40 Mg/kg/d tds/qid Vancomycin Clindamycin 20-40 Mg/kg/d tds/qid Vancomycin Somg/kg/d qid or Clindamycin 20-40 Mg/kg/d tds/qid Vancomycin Somg/kg/d tds/qid Vancomycin Somg/kg/d tds/qid Somg/kg/d tds/qid or combination therapy Somg/kg/d qid or Somg/kg/d tds/qid Somg/kg/d tds/qid or combination therapy Somg/kg/d qid or Somg/kg/d qid or Somg/kg/d tds/qid or combination therapy Somg/kg/d qid or Somg/kg/d qid or Somg/kg/d qid or Somg/kg/d qid or Somg/kg/d tds/qid Somg/kg/d tds/qid or Somg/kg/d tds/qid or combination therapy Somg/kg/d qid or Somg/	aallydid			oomg/kg/a qia
10-14 days Cloxacillin 200mg/kg/d Vancomycin IV Meropenem (120 mg /kg/da divided 8 hrly)/ plus Vancomycin 150mg/kg/d tds or Clindamycin 20-40 mg/kg/d tds/qid 20-40 mg/kg/d tds/qid 20-40 mg/kg/d tds/qid Mays Plus either Cefotaxime 150mg/kg/d tds or Ceftriaxone 100mg/kg/d tds or Ceftriaxone 150mg/kg/d tds or Ceftriaxone 150mg/kg/d tds or Ceftriaxone 150mg/kg/d tds or Ceftriaxone 100mg/kg/d od/bd Osteochondritis Piperacillintazobactam300 mg/kg/d tds/qid or Combination therapy Piperacillintazobactam300 mg/kg/d tds/qid or Combination therapy Piperacillintazobactam300 mg/kg/d tds/qid or Ceftriaxone Ceftriaxon	cenunus			
Bone and Joint Cloxacillin 200mg/kg/d Vancomycin IV Meropenem (120 mg /kg/da divided 8 hrly)/ plus Vancomycin 60mg/kg/d qid or 150mg/kg/d tds or Ceftriaxone 100mg/kg/d od/bd + Vancomycin x 10-14 days Plus either Cefotaxime 150mg/kg/d tds or Ceftriaxone 100mg/kg/d tds or Ceftriaxone 150mg/kg/d tds or Ceftriaxone 150mg/kg/d tds or Ceftriaxone 150mg/kg/d tds or Ceftriaxone 100mg/kg/d od/bd Osteochondritis Piperacillin- tazobactam300 mg/kg/d tds/qid or combination therapy		0 0	bumg/kg/a qia	
Bone and Joint 200mg/kg/d Vancomycin IV Meropenem (120 mg /kg/da divided 8 hrly)/ plus	_			
Infections Description De				
Infections plus either Cefotaxime 150mg/kg/d tds or 150mg/kg/d tds or Clindamycin 20-40 mg/kg/d tds/qid 20-40 mg/kg/d tds/qid 20-40 mg/kg/d tds/qid 20-40 mg/kg/d tds/qid mg/kg/d tds/qid 20-40 mg/kg/d tds/qid mg/kg/d	Bone and Joint	200mg/kg/d	Vancomycin	
Acute Ceftriaxone 100mg/kg/d od/bd + Vancomycin x 10-14 days Plus either Ceftriaxone 150mg/kg/d tds/qid Plus either Cefotaxime 150mg/kg/d tds or Clindamycin 20-40 mg/kg/d tds/qid Plus either Cefotaxime 150mg/kg/d tds or Ceftriaxone 100mg/kg/d od/bd Osteochondritis Piperacillin- tazobactam300 mg/kg/d tds/qid or combination therapy Clindamycin 20-40 mg/kg/d tds/qid Domg/kg/d tds/qid Clindamycin 20-40 mg/kg/d tds/qid Domg/kg/d tds/qid Domg/kg/d od/bd				
Acute Ceftriaxone 100mg/kg/d od/bd + Vancomycin x 10-14 days Plus either Cefotaxime 150mg/kg/d tds or Ceftriaxone 100mg/kg/d tds or Ceftriaxone 100mg/kg/d tds or Ceftriaxone 100mg/kg/d od/bd Osteochondritis Piperacillin- tazobactam300 mg/kg/d tds/qid or combination therapy	Infections		60mg/kg/d qid or	
Acute Ceftriaxone 100mg/kg/d od/bd + Vancomycin x 10-14 days Plus either Cefotaxime 150mg/kg/d tds or Ceftriaxone 100mg/kg/d od/bd Osteochondritis Piperacillin- tazobactam300 mg/kg/d tds/qid or combination therapy mg/kg/d tds/qid		150mg/kg/d tds or		
Osteomyelitis/s eptic arthritis 100mg/kg/d od/bd + Vancomycin x 10-14 days Plus either Cefotaxime 150mg/kg/d tds or Ceftriaxone 100mg/kg/d od/bd Osteochondritis Piperacillintazobactam300 mg/kg/d tds/qid or combination therapy	A4-	C-6:	Clindamycin 20-40	20-40
Osteomyelitis/s eptic arthritis Osteomyelitis/s eptic arthritis Osteochondritis Osteochondritis Piperacillintazobactam300 mg/kg/d tds/qid or combination therapy	Acute		mg/kg/d tds/qid	mg/kg/d tds/qid
eptic arthritis 10-14 days Plus either Cefotaxime 150mg/kg/d tds or Ceftriaxone 100mg/kg/d od/bd Osteochondritis Piperacillin- tazobactam300 mg/kg/d tds/qid or combination therapy	0.4	0 0		
Cefotaxime 150mg/kg/d tds or Ceftriaxone 100mg/kg/d od/bd Osteochondritis Piperacillin- tazobactam300 mg/kg/d tds/qid or combination therapy				×0'
Osteochondritis Piperacillintazobactam300 mg/kg/d tds/qid or combination therapy	eptic arthritis	10-14 days	Plus either	
Osteochondritis Piperacillintazobactam300 mg/kg/d tds/qid or combination therapy			0	O
Ceftriaxone 100mg/kg/d od/bd Osteochondritis Piperacillin- tazobactam300 mg/kg/d tds/qid or combination therapy				ľ
Osteochondritis Piperacillin- tazobactam300 mg/kg/d tds/qid or combination therapy			150mg/kg/d tds or	
Osteochondritis Piperacillin- tazobactam300 mg/kg/d tds/qid or combination therapy			Ceftriaxone	
tazobactam300 mg/kg/d tds/qid or combination therapy			100mg/kg/d od/bd	
mg/kg/d tds/qid or combination therapy	Osteochondritis	Piperacillin-		
combination therapy		tazobactam300		
combination therapy		mg/kg/d tds/qid or		
With Cloracinin				
200mg/kg/d plus			7	
Ceftazidime				
100mg/kg/d tds		100mg/kg/d tds		
		5 40/1 0		
7-10 days after surgery		7-10 days after surgery		
		(1) (1):		
Cloxacillin W : 111 1 1 200				D: 311: 4 1 4 200
Skin and soft 200mg/kg/d Vancomycin Piperacillin- tazobactam 300	Skin and soft	200mg/kg/d	Vancomycin	1 ÷
mg/kg/d tds/qid or IV		C C 1: 40	60 /1 /1 11	
tissue or Cefazolin 60- 60mg/kg/d qid Meropenem	tissue	or Cefazolin 60-	60mg/kg/d qid	1 1
				(120 mg /kg/day divided 8 hrly
1 2 P 4 2 P 7	infections	100mg/kg/d or		1 *
infections 100mg/kg/d or plus		Clindamyoin 20 40		vancomycin 60mg/kg/d qid
1 2. P. 42 10 7	Y	CilliualityCIII 20-40		
infections 100mg/kg/d or Clindamycin 20-40 plus Vancomycin 60mg/kg/d qid	Y	·		
infections plus Vancomycin 60mg/kg/d aid	Y	mg/kg/d tds/qid x 7-10		

	1	1,,	
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Laiwani Medical Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 52 of 8



Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026 Version - 11.0

A . 11.4	A ' '11' / 1 1	A 1.	
Animal bite	Amoxicillin/clavulanate	Alternatives	
wounds (dog / cat)	50mg/kg/d tdsi.v or p.o	Piperacillin300mg/kg/d qid 7-10 days	NA
		Penicillin allergy	
		Clindamycin20- 40mg/kg tds/qid plus TMP /SMX 80mg/kg/ bd X 7-10 days (dog bites); or cefuroxime 20-30mg/kg/d x 7-10 days (cat bites)	CONTR
Vascular		Piperacillin-	Meropenem 120mg/kg/d tds plus Vancomycin 60mg/kg/d
catheter associated Infections		tazobactam300 mg/kg/d tds/qid + Vancomycin60mg/kg/d qid	qid
Severe Sepsis/septic shock	Cefotaxime 150 mg/kg/day divided 6-8 hrly OR Ceftriaxone 100 mg/kg/day divided 12 hrly +/- amikacin 15-20 mg/kg/d od	IV Piperacillin – Tazobactam 300-400 mg/kg/day divided 8 hrly + IV Vancomycin 45-60 mg/kg/day divided 6-8 hrly	IV Meropenem 80-120 mg/ kg/8 hrly + IV Vancomycin 45-60 mg/kg/day divided 6-8 hrly

				/ '
Prepared by	Checked by	Verified by	Appr	eved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department		K Natwani cal Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 53 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 11: Empiric Therapy of Neonatal Intensive Care Unit Sepsis and Meningitis

Diagnosis	Suspected organisms	Early onset	Late onset	Nosocomial	Community acquired	Duration
Sepsis	Klebsiella, Acinetoba cter, E.coli, Enterococc us, Others :Serratia, Burkholde ria, Pseudomo nas, Proteus	Gentamicin (for haemodynamical ly stable) Piperacillin- Tazobactum (for haemodynamical ly unstable)	1 st line :Piperacillin- Tazobactam 2 nd line: Meropenem 3 rd line: Colistin	1 st line Piperacillin- Tazobactam 2 nd line: Meropenem 3 rd line: Colistin	1 st line :Cefotaxime and Amikacin 2 nd line:Piperacillin- Tazobactam 3 rd line: Meropenem 4 th line: Colistin	10days
Pneumonia	E coli, Klebsiella, Acinetoba cter, Enterococc us, Staphyloco ccus (CONS) Others :Serratia, Burkholde ria, Pseudomo nas, Proteus	Gentamicin (haemodynamica lly stable) Piperacillin- Tazobactam (haemodynamica lly unstable)	1 st line :Piperacillin- Tazobactam 2 nd line: Meropenem 3 rd line: Colistin	1 st line Piperacillin- Tazobactam 2 nd line Meropenem 3 rd line Colistin	Ceftriaxone plus Azithromycin	7days
NEC	Proteus		1 st line Piperacillin- Tazobactam and Amikacin 2 nd line Meropenem 3 rd line Colistin	1 st line Piperacillin- Tazobactam 2 nd line Meropenem 3 rd line Colistin	1 st line Piperacillin- Tazobactam 2 nd line Meropenem 3 rd line Colistin	7-10days
Meningitis	For early onset: E coli, GBS, enteric bacilli, listeria,	1 st line: Cefotaxime plus Gentamicin 2 nd line: Meropenem Checked by	Meropenem	Meropenem	Ceftriaxone /cefotaxime	Gram Positive: 14-days Gram negative: 21 days#
Dr. M Kumar Prof Microbio	streptococ cus, H	Dr. M S Modak Prof & HoD Microbiology	Dr. S Qual	Milpar Acharya ity Department	Dr. SK Natwa Medical Bire	#Ventriculi

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 54 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026 Version - 11.0

	influenza,					tis/Brain
	Neisseria					abscess: 6-
	meningitid					8 weeks
	es.					
	For late					
	onset:					
	Klebsiella,					
	Acinetoba					
	cter,					
	E.coli,					~
	Enterococc					
	us,				ACC.	
	Staphyloco ccus				~ (2)	
	(CONS)					
	Others					
	:Serratia,					
	Burkholde					
	ria,					
	Pseudomo				, , , , , , , , , , , , , , , , , , ,	
	nas,					
TIOT	Proteus		1 ct 1 ·	1 et 1:	A '1 '	101
UTI	Enterococc		1 st line:	1 st line	Amikacin	10days
	us, E coli,		Piperacillin-	Piperacillin-		
	Enterobact		Tazobactam 2 nd line:	Tazobactam 2 nd line:		
	er		Meropenem	Meropenem		
			3 rd line:	3 rd line:		
			Colistin	Colistin		
Skin and	Staphyloco	• X	1stline:Cloxa	Vancomycin	Cloxacillin	7days
soft tissue	ccus		cillin	,		,
infection			2 nd line:Vanc			
			omycin			
Arthritis	Staphyloco	Y	1 st line	1 st line:	Ceftriaxone plus	Culture
	ccus	y	Piperacillin-	Piperacillin-	Vancomycin	Negative:
	,Klebsiella	>	Tazobactam	Tazobactam		2weeks
	2		2 nd line	2 nd line:		Culture
			Meropenem 3 rd line	Meropenem 3 rd line		positive: 3 weeks
	.00		Colistin	Colistin		weeks
	, ,		Consult	Consuli		
Osteomyeli	Staphyloco		1 st line	1 st line	Ceftriaxone plus	4 weeks
tis	ccus,		Piperacillin-	Piperacillin-	Vancomycin	-
	Gram		Tazobactam	Tazobactam		
	Negative		2 nd line	2 nd line		
	Bacilli		Meropenem	Meropenem		
			3 rd line	3 rd line		
			Colistin	Colistin		

	1	1,,	
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Laiwani Medical Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 55 of 8



Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026 Version - 11.0

Catheter	Staphyloco		1 st line:		10days
related	ccus(CON		Vancomycin		3 3 3 3
Infection	S),		and		
Infection					
	S.aureus,		Amikacin		
	Gram		2 nd		
	negative		line:Piperacil		
	bacteria		lin-Tazobact		
			3 rd line:		
			Meropenem		
			4 th line		7
			Colistin		~
			Collstill		
Fungal	Candida		Amphoterici		Depending
infection	albicans		n B or		on location
	and		Fluconazole		
	Candida		(depending		
	Non		on		
	albicans		Antifungal		
	aro rouris		susceptibility		
			report)	-9	
			report)		
				<u> </u>	
		110501			

			/	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Datwani Medical Director	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 56 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table: 12 Empiric therapy of Ophthalmic infections

Sr. No	Category	Suspected Organisms	First Line	Alternative
1	Bacterial	S aureus and albus	Topical Moxifloxacin 0.5% eyedrops 3-6 times per day	
	conjunctivitis	H Aegyptius H Influenzae, C diphtheriae	Tobramycin eye ointment at bed time Penicillin eye drops 10,000 units/ml	XSE
2	Bacterial	Pseudomonas,	Moxifloxacin eye drops 0.5% 1 hourly	Fortified
2	Dacterial	S.aureus	Moximoxaciii eye drops 0.5% 1 nourry	Vancomycin
	Keratitis	Pneumococcus	Fortified Tobramycin eye drops	eye drops
	Trotutio	N gonorrheae	Totalica Toolamyem eye arops	eye drops
		1 (Bononinous		Amikacin eye
			,0,	drops
3	Fungal	Aspergillus,	Natamycin eye drops 6 times a day	Amphotericin B
	Keratitis			eye drops
		Fusarium, Candida	Itraconazole eye drops /ointment at bed	Voriconozole eye
		albicans	time Tablet Fluconazole 150mg twice	drops
			a day& eye drops 4-6 times per day	Intracameral
			Nystatin eye ointment	Amphotericin B
4	Viral Keratitis	H Simplex	Acyclovir Tablet 800mg 5 times a day	Tablet
		** =	and ointment 5 times a day	Valacyclovir
		H Zoster	Gancyclovir ointment	1000mg 3 times a
_	T 1 1 1 1 1 1	C C	1	day
5	Endophthalmit is	S aureus	Intravitreal Vancomycin 1 mg /0.1 ml	Intravitreal
	18	Sepidermidis	andAmikacin 400microgrames /.ml	Vancomyci n 1mg /0.1ml and
		Streptococcus	and Annikacin 400microgrames /.im	Ceftriaxone
		Pseudomonas		2.25mg/0.1ml
		H Influenzae	Intravitreal Amphotericin B	2.23mg/0.1mm
		Candida /fusarium	maravarear ramphoteriem B	
6	Orbital	Staphylococci	Intravenous	Intravenous
	cellulitis	7 7 11111		Ceftriaxone
	4.0		Piperacillin and Tazobactam 4.5g	
			twice a day	
	10 -		Intravenous Metronidazole 100ml 3	
	D'		times a day	
	· ·	Mucormycosis/Asper	Intravenous Amphotericin B	
7		gillosis	T 11 . A	
7	Acute	Staphylococcus,	Tablet Amoxicillin and Clavulanic acid	
	Dacryocystitis	-	625 mg twice a day	
		Pneumococcus	Moxifloxacin eye drops 0.5% 3-6	
			times a day	
			a day	1

Prepared by	Checked by	Verified by	Approved by
D	Dr. M S Modak	Dr. Shilpa Acharya	
Dr. M Kumar	Prof & HoD		Dr. SK Lalwani
Prof Microbiology	Microbiology	Quality Department	Medical Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 57 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 13: ENT Infection

Name of	Patient Type 1	Patient Type 2	Patient Type 3
condition	(Community acquired)		
Acute infection	Inj Ampicillin 1 gm q6h	-	-
like acute	Amoxicillin		
membranous	+clavulanic acid 1.2		
tonsillitis, ASOM,	gm q8h		
Acute epiglottitis			
without			×
complication			
Acute infection	Addition	-	-
with	Of aminoglycoside for		
complications like	gram negative coverage		
acute mastoiditis,	and metronidazole for		~
Quinsy	anaerobic coverage		
Chronic infection	Amoxicillin	ID/ Medicine	ID/Medicine consult
without	+clavulanic acid 1.2	consult	
complication like	gm q8h IV		
CSOM, chronic	Ceftriaxone 1 gm q12h		
sinusitis	IV		
	Inj Ceftriaxone+ inj	2	
Chronic infection	amikacin	ID/ Medicine	ID/ Medicine consult
with	+ inj metronidazole	consult	
complications like	A	-0-	
meningitis, orbital		>	
cellulitis, brain	. X.O		
abscess			

			/ .	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Stilpa Acharya Ouality Department	Dr. SK Natwani Medical Director	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 58 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 14: Surgical site infection

Name	Type 1	Type 2	Type 3
Head & Neck	Ceftriaxone 1gm	Meropenem 2gm q8h	If fungal infection
	q12h IV	IV	suspected
	+	+	Ampho B
	Metronidazole	Vancomycin 1 gm	If VRE suspected
	Or	q12h IV	Linezolid
	PIP-TZ 4.5 gm q6h		If XDR or PDR
	IV		Gram negative
	If MRSA suspected		infection suspected
	Add Vancomycin		Colistin 4.5MUBD If
	1gm IV q12h		CNS infection
	If CNS infection		Add intrathecal
			antimicrobials as
	Ceftazidime 2 gm		above
	q8h IV instead of		
	Ceftriaxone/PIP-TZ		<i>J</i>
Other infections	Ceftriaxone 1gm	Meropenem 2gm q8h	If fungal infection
Sternal infections	q12h IV	IV	suspected
Chest	+	+	Ampho B
Abdominal	Metronidazole	Vancomycin 1 gm	If VRE suspected
Perineal	Or	q12h IV	Linezolid
	PIP-TZ 4.5 gm q6h		If XDR or PDR
	IV		Gram negative
	If MRSA suspected		infection suspected
	Add Vancomycin		Colistin 4.5MUBD If
	1gm IV q12h		clostridium difficile
	.40 ′		colitis or sepsis
	57		suspected
	4 O'		Oral Vancomycin
			250 mg q6h
•	Y		+ Matronidazala 500
X	>		Metronidazole 500
			mg q8h IV

Note:

Surgical debridement is almost always necessary. Any graft, device or foreign body must be removed.

			,	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Ouality Department	Dr. SK Nalwani Medical Nirector	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 59 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 15: Catheter related blood stream infections (CRBSI)

Name	Type 1	Type 2	Type 3
Peripheral catheter		Cloxacillin 1 gm q6h IV	Ceftriaxone 1gm - q12h IV
Central venous catheter (short term) Dialysis catheter	-	+ Meropenem 2gm q8h IV Vancomycin 1 gm	Meropenem 2gm q8h IV + Vancomycin 1 gm
(short term)		q12h IV	q12h IV
Dialysis catheter (long term) Hickman or other implanted catheter (long term)		and Reserve	If fungal infection (Non- AlbicansCandida suspected) Ampho B iv Or Caspopfungin 70 mg IV q24h flowed by 50 mg If VRE suspected Linezolid If XDR or PDR Gram negative infection suspected Colistin 4.5MUBD

Note:

Change catheter if signs of thrombophlebitis are present

Catheter cultures and blood cultures to be sent as per HICC protocol.

Catheter maybe kept in situ pending culture reports especially if CRBSI not strongly suspected and no other IV access is available

Remove catheter immediately if local signs of suppuration present or if central venous catheter and blood cultures are positive

			,	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Ouality Department	Dr. SK Nalwani Medical Nirector	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 60 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

DEFINITIVE THERAPY ONCE THE CAUSATIVE ORGANISM IS IDENTIFIED

It is vital to send cultures before empiric antimicrobials are prescribed. Once cultures results are available the next steps are

- 1. Decide whether the organism grown is a colonizer or an actual pathogen. Evaluate carefully if the site from which culture has been sent has active infection either from clinical signs or from elevated WBC counts or radiological evidence.
- 2. Don't treat colonizing organisms, Consult microbiology for the decision
- 3. Choose the simplest antimicrobial class to which the organism shows susceptibility
- 4. If the cultures show intermediate susceptible or multidrug resistant organism, consult infectious disease specialist for choice of appropriate antimicrobial.
- 5. Linezolid should be given only in culture confirmed MRSA infections after consultation with ID physician.
- 6. Levofloxacin is reserved for use in culture confirmed pulmonary infections only
- 7. Do not continue therapy beyond indicated duration. If the duration is to be exceeded then the clinician will justify the same and endorse it in the clinical notes

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Prof & HoD

Prof Microbiology

Dr. Skilpa Acharya

Quality Department

Prof Medical Pricetor



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table 16. Antimicrobial Prophylaxis for Surgery Clean and Clean Contaminated cases

Procedure	Antimicrobial	
Clean surgeries (example: elective		
hernia repair, breast surgeries)	Cefazolin / Cefuroxime	
	Cefazolin / Cefuroxime (add amikacin and metronidazole	
	in open fractures)	
Orthopedic surgery		
Cardiovascular / vascular surgery	Cefazolin / Cefuroxime	
Neurosurgery	Cefazolin / Cefuroxime	
Ophthalmic surgery	Topical quinolone. Systemic- Cefazolin / Cefuroxime	
	Cefazolin / Cefuroxime/Amoxyclav	
Head, neck and ENT surgery	(Ceftriaxone in cases involving dural exposure)	
Gastroduodenal	Cefuroxime / Cefazolin	
A constant of Colombia	Cofeesia (Cofeesia en IM-residente	
Appendicular / Colorectal surgery	Cefuroxime / Cefazolin and Metronidazole	
Biliary	Cefuroxime / Cefazolin/ cefoperazone- sulbactam	
Abdominal / Vaginal hysterectomy /	Sulvaciani	
Caesarian		
section/MTP/Sterilization/Laparoscopic		
procedures	Cefazolin / Cefuroxime +Metronidazole	
Urologic surgery	Cefuroxime (or as guided by urine culture)	
Opthalmology	Topical Quinolone eye drops	
Intraocular surgeries under LA	r Comments and analysis	
Surgeries under GA (Clean Surgeries)	IV Inj. Cefazoline / Cefuroxime Topical Quinolone eye drops	

Note: Preoperative dose of antimicrobial is to be given 15 to 60 minutes before incision

- Dose of Cefazolin 2 gm IV
- Dose of Cefuroxime 1.5 gm IV
- Dose is to be repeated if surgery> 4 hours
- If penicillin allergy: consider either clindamycin or Vancomycin (infusion to be started more than 60 min before incision for vancomycin in view of prolonged infusion time required) (WHO Guidelines on surgical prophylaxis 2018)
- Antimicrobial prophylaxis must not be continued for more than 24 hours after surgery
- In case patient is already on antimicrobials the same antimicrobial should be adjusted for the period of surgery.

Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Nalwani Medical Director	



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Table 17: Pediatric surgery

Procedure	Antimicrobial	Duration
Urosurgery	Amoxiclav	3 days
Genitourinary: Clitoroplasty	Amoxiclav	5days
Hypospadias, vaginoplasty (
plain)		
Vaginoplasty (use of bowel)	Ceftriaxone +	1 day prior
	Metronidazole	X
Hirschprung ARM pull-through	Ceftriaxone + metronidazole	3 days
Esophageal procedures	Amox-clav	$\nearrow \bigcirc$
Biliary tract	Cefoperazone -sulbactam	3 days
Appendix : Nonperforated	Cefuroxime	1 dose
Perforated	Ceftraxone + Metronidazole	
	Piptaz if required	χΟ.
Paediatric Tumours	Cefuroxime	3 doses

Clean surgeries:

Citali sui geries.		
Procedure	Antimicrobial	Duration
Circumcision, Orchiopexy	Cefuroxime	Single dose
Herniotomy, LN biopsy,	No antimicrobials	-
Thyroglossal cyst Lipoma,		
small lumps	4.0	
Bharait		

Prepared by

Checked by

Or. M S Modak

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

Checked by

Verified by

Approved by

Dr. SK laiwani

Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026 Version - 11.0

Table 18: Antimicrobial prophylaxis for GI endoscopy

Antimi	crobial prophylaxis and/o	or treatment to prevent le	ocal infections	
Patient condition	Procedure contemplated	Goal of prophylaxis	Antimicrobial	Periprocedural antimicrobial prophylaxis
Bile duct obstruction in absence of cholangitis	ERCP with complete drainage	Prevention of cholangitis	Nil	Not recommended
Bile duct obstruction in absence of cholangitis	ERCP with incomplete drainage	Prevention of cholangitis	Ceftriaxone	Recommended; contin antimicrobials after procedure X 3 days
Solid lesion in upper GI tract	EUS-FNA	Prevention of local infection	-	Not recommended
Solid lesion in lower GI tract	EUS-FNA	Prevention of local infection	-	Not recommended
Pancreatic cysts	EUS-FNA	Prevention of cyst infection	Ceftriaxone+ Metronidazole	Three doses
All patients	Percutaneous endoscopic feeding tube placement	Prevention of peristomal infection	Cefazolin/Cefuroxime	Recommended single dose
Cirrhosis with acute GI bleeding	Required for all patients regardless of endoscopic procedures	Prevention of infectious adverse events and reduction of mortality	Ceftriaxone	On admission
Synthetic vascular graft and other nonvalvular cardiovascular devices	Any endoscopic procedure	Prevention of graft and device infection	-	Not recommended
Prosthetic joints	Any endoscopic procedure	Prevention of septic arthritis	-	Not recommended
Peritoneal dialysis	Lower GI endoscopy	Prevention of peritonitis	Ceftriaxone + metronidazole	Suggested
EU	S-FNA, EUS-guided FNA.	1.0		

			,	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Nalwani Medical Nirector	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 64 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Table -19 FEBRILE NEUTROPENIA

Febrile Neutropenia-definition

- Neutropenia-ANC<500/mm3and expected to fall below 500/mm3 in 48hrs
- Fever-single oral temperature of 38.3oC(1010F) on one occasion or 38oC (100.40F) on atleast 2 occasions (1 hour apart)
- Neutropenic patients may not have usual signs of infection. Redness, tenderness and fever may be the only signs.

Protocol:

- Critical examination of areas usually harboring infections, including but not limited to, oral cavity, axillary region, scalp, groin, perineal region.
- Send blood Cultures 2 sets (each bottle 10ml x 4 bottles)
- Other relevant investigations: urea, creatinine, ALT, urine culture ,Chest Xray, separate culture from central line, etc.

Patient-Hemodynamically stable

- Blood culture 2 sets
- Start IV Cefoperazone sulbactam 1.2gm IV 8 hourly
- No need to add glycopeptides in the initial regimen (except in specific situations, given below)

Patient-Hemodynamically unstable

• Start BL-BLI agent(Cefoperazone-Sulbactam 1.2gm IV 8 hourly/ piperacillin- tazobactam 4.5gm IV 8 hourly) OR

Carbapenem (meropenem 1gm IV 8 hourly/imipenem 500mg IV 6 hourly/doripenem 500mg IV 6 hourly)

No need to add glycopeptides in the initial regimen (except in specific situations, givenbelow)

Reassess after 48 hours:

If blood cultures are negative, hemodynamically stable but still febrile

- Reculture blood
- Add amikacin 500mg IV BD for 3days
- Add colistin (instead of amikacin) if indicated (see below)

If blood cultures are negative, hemodynamically unstable but still febrile

Inj Colistin (+/-Carbapenem) + glycopeptides + Echinocandin/ L-AmphoB

Blood culture growing Gram negative bacilli

Patient afebrile- continue the empirical antimicrobial till antimicrobial sensitivity is available.

 Rationalise as 	s per susceptibility profiles			
Prepared by	Checked by	Verified by	Approved by	
When to add glycope	eptides r. M S Modak	Dehary	4	
Dr. M Kumar	Prof & HoD	Dr. Shilpa Acharya	Dr. SK Dalwani	
Prof Microbiology	Microbiology	Quality Department	Medical Director	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 65 of 81

Version - 11.0



Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026

- 1. Haemodynamic instability, or other evidence of severe sepsis, septic shock or pneumonia
- Colonisation with MRSA or penicillin-resistant S. pneumonia
- 3. Suspicion of serious catheter-related infection e.g. chills or rigors within fusion through catheter and cellulitis around the catheter exit site
- 4. Skin or soft-tissue infection at any site
- 5. Positive blood culture for gram-positive bacteria, before final identification and susceptibility testing is available
- 6. Severe mucositis

When to add empirical colistin in febrile neutropenic patients?

- 1. Hemodynamic instability.
- 2. Colonization with carbapenem resistant gram-negative bacteria.
- 3. Previous infection with carbapenem resistant gram-negative bacteria.
- 4. GNB in blood, sensitivity pending, persistent fever with hemodynamic instability.

Empirical Antifungal Therapy

- No response to broad spectrum antimicrobials (3-5days)- add L-AmphoB/echinocandin
- When a patient is located at a remote area and may not have access to emergency healthcare services, febrile neutropenia can be life threatening. Under such circumstances, availability of broadspectrum oral antimicrobials with the patient can help them gain time to reach emergency healthcare service.

Useful tips

- Febrile after 72hrs- CT chest and consider empirical antifungal.
- If fever persists on empirical antimicrobials, send two sets blood cultures/day for 2 days
- Send further cultures if clinical deterioration
- Unexplained persistent fever in otherwise stable patient doesn't require change in empirical antimicrobial regimen.

Continue the regimen till ANC is >500cells/mm3

- If glycopeptides started as a part of empirical regimen, STOP after 48hrs, if no evidence of Gram positive infection
- Antimicrobial treatment should be given for at least seven days with an apparently effective antimicrobial, with at least four days without fever.
- Once Neutrophil count has recovered, with no culture positivity and hemodynamically stable; antimicrobials can be stopped and patient observed, even if remains febrile. Evaluate for fungal infection, if at risk.

			/ '	
Prepared by	Checked by	Verified by	Approved by	
Artiviral prophylaxis	Dr. M S Modak	Jehary	42	
Dr. M Kumar Prof Microbiology	Prof & HoD Microbiology	Ouality Department	Medical Director	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 66 of 81

Version - 11.0



Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026

- For HSV IgG positive patients undergoing allo-HSCT or leukemia induction needs acyclovir prophylaxis
- All patients being treated for cancer need to receive annual influenza vaccination with an inactivated

vaccine.

• Neutropenic patients presenting with influenza like illness should receive empirical treatment with neuraminidase inhibitor.

Antifungal prophylaxis

- a) Induction chemotherapy of Acute Leukemia: Posoconazole
- b) Post allo BMT

Pre engraftment:

Voriconazole/ echinocandin

Post engraftment:

Posoconazole

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

Dr. Skilpa Acharya

Quality Department

Medical Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 67 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Table - 20 Cardiac Device implant Prophylaxis

A risk assessment score is proposed with one point for each

a) Diabetes mellitus

b) Heart failure

c) Oral anticoagulation therapy

d) Chronic corticosteroid use

e) Renal insufficiency/failure

f) Prior CIED infection

g) Presence of more than two leads implanted

h) Presence of epicardial leads

k) Use of temporary pacemaker

I) Replacement, or upgrade procedure.

According to the score, patients will be stratified in two groups and the **same documented in the case notes**:

- (a) low infective risk (score <3)
- (b) high infective risk (score ≥3)

Patients in the 'low-risk' group will treated with only two doses of antimicrobials, both intravenous, of whom the first 15-60min before skin incision and the second after 8 hours. Patients in the 'high-risk' group will get intravenous prophylaxis for two full days (of whom the first administration 15-60min before skin incision

Antimicrobial prophylaxis will be amoxicillin/clavulanic acid unless the patient had a history of allergic reactions to penicillin. The dosage will be dependent on renal function:

- (a) intravenous amoxicillin/clavulanic acid 2/0.2 g in patients with creatinine clearance (CrCl) >30 mL/min and 1/0.2 g in patients with CrCl <30mL/min
- (b) oral amoxicillin/clavulanic acid 875/125 mg every 8 h in patients with CrCl >30 mL/min, and 875/125 mg every 12 h in patients with CrCl <30mL/min.
- (c) In case of penicillin allergy Intravenous clindamycin day for 2 doses for the low risk group and two days for the high risk group.

Prepared by

Checked by

Verified by

Approved by

Dr. M S Modak

Prof & HoD

Prof Microbiology

Microbiology

One Medical Director

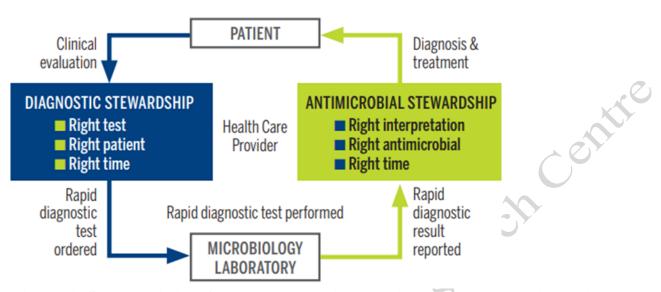
Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 68 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Diagnostic Stewardship



Diagnostic Stewardship is defined as the co-ordinated guidance and interventions to improve appropriate use of microbiological diagnostics to guide therapeutic decisions. It should promote appropriate, timely diagnostic testing, including specimen collection and pathogen identification and accurate , timely reporting of results to guide patient treatment

	1	1,,	
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Laiwani Medical Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 69 of 81

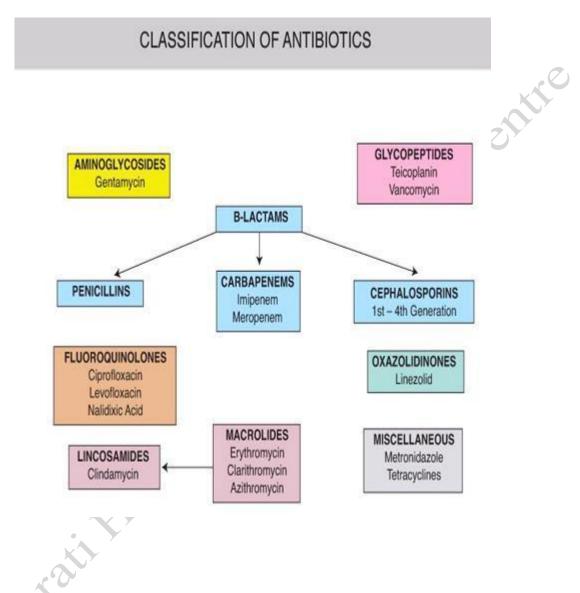


Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Appendix 1

Commonly used antimicrobials



			/ `
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Smipa Acharya Quality Department	Dr. SK Naiwani Medical Nirector

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 70 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Table 21 : Spectrum of commonly used antimicrobials:

Antimicrobial				
Class	Name	Organisms	Indication & Dose	Side effects
Penicillins				Allergy
β-lactamase susceptible	Penicillin G Penicillin V Ampicillin. Amoxicillin (PO)	Gram +ve Gram +ve Gram +ve & Gram -ve Gram +ve	Not easily available 1-2 gms q6h 500 mg q8h	N.C
β – lactamase resistant	Cloxacillin	Gram +ve	0.5-1gm q6h	
β-lactam/ β- lactam inhibitor combination	Piperacillin- tazobactam. Ampicillin- sulbactam. Amoxicillin- clavulanate (IV)	ESBL Gram –ve organisms ESBL Gram –ve organisms Gram +ve & Haemophilus. influenzae	4.5 gm q6h as infusion 1 gm q6h 1.2 gm q8h	
Cephalosporins				
1 st Generation	Cefazolin (IV) Cephalexin (PO)	Gram +ve	1gm q8h 500 mg q8h	
2 nd Generation	Cefadroxil (PO) Cefuroxime (PO & IV)	Gram +ve Gram +ve	500 mg q12h 750 mg q8h	
3 rd Generation	Cefotaxime Ceftriaxone Ceftizoxime Ceftazidime Cefixime (PO) Cefpodoxime (PO) Cefdinir (PO)	Gram +ve & Gram -ve Gram +ve & Gram -ve Gram +ve & Gram -ve Gram +ve & Gram -ve Anti-pseudomonas	1 gram q6h 1-2gm q12h 1 gm q12h 1-2 gm q8h 200 mg q12h	

				/ '
Prepared by	Checked by	Verified by	Appr	oved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Smipa Acharya Ouality Department		K Latwani

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 71 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program 2025-2026 Version - 11.0

Clase Name Cefoperazone Sulbactam	Antimicrobial				Side
Cefoperazone beta lactamase inhibitor Sulbactam	Class	Name	Organisms	Indication& Dose	effects
beta lactamase inhibitor Sulbactam Sulbactam Sulbactam Sulbactam Sum q12h Sum q12h Sum q12h Sum q12h Sum q24h S	Cephalosporin Plus	Cefoperazone			
inhibitor Aminoglycosides Streptomycin Kanamycin Gram -ve 3mg/kg q24h s Smg/kg q24h s Smg/kg q24h s Smg/kg q24h Smg/kg q24h q4h Smg/kg q24h Smg/kg q24h Smg/kg q24h Smg/kg q24h Smg		/sulbactam	-	1.5 gm –	
Aminoglycosides Kanamycin Gentamicin Amikacin Tobramycin Netilmicin Ciprofloxacin Clarbapenems Imipenem- cilastatin Meropenem Doripenem Ertapenem Ertapenem Polymyxins Polym	inhibitor				
Kanamycin Gentamicin Amikacin Gram -ve Smg/kg q24h Muscle weaknes Smg/kg q24h Weaknes Smg					
Gentamicin Amikacin Tobramycin Netilmicin Gram -ve 3mg/kg q24h Muscle weaknes Smg/kg q24h Westlimicin Gram -ve 5mg/kg q24h Westlimicin Gram -ve 5mg/kg q24h Westlimicin Smg/kg q24h Westlimicin Smg/	Aminoglycosides	Streptomycin	Gram –ve	0.75 -1gm q24h	Deafnes
Amikacin Tobramycin Netilmicin Quinolones Nalidixic acid Norfloxacin Ciprofloxacin Ofloxacin Spectrum Carbapenems Imipenem- cilastatin Meropenem Doripenem Doripenem Doripenem Ertapenem Ertapenem Ertapenem Ertapenem Carbapenems Ertapenem Doripenem Carbapenems Doripenem Doripenem Carbapenems Ertapenem Doripenem Carbapenems Ertapenem Doripenem Carbapenems Ertapenem Doripenem Carbapenems Ertapenem Doripenem Carbapenems Doripenem Carbapenems Ertapenem Doripenem Carbapenems Ertapenem Doripenem Carbapenems Ertapenem Doripenem Carbapenems Ertapenem Ertapenem Does not cover Pseudomonas, Acinetobacter & Enterococcus Polymyxins Polymyxins Polymyxins Polymyxins Polymyxins Polymyxins Polymyxins Colistin ESBL, Metalloproteinase Colistin Corm +ve and anerobes Colomg q8h Muscle weakness Colistin		Kanamycin	Gram –ve	3mg/kg q24h	S
Tobramycin Netilmicin Gram -ve Smg/kg q24h weaknes S		Gentamicin	Gram –ve	13mg/kg q24h	Vertigo
QuinolonesNetilmicinGram -vesNalidixic acid Norfloxacin Ciprofloxacin Offloxacin Doffloxacin Doripenem1 gm q6h 400 mg q12h 500 mg q12h 750 mg q24h 400 mg q24hSeizuresExtended spectrumLevofloxacin Moxifloxacin750 mg q24h 400 mg q24hSeizuresCarbapenems Imipenem- cilastatin Meropenem DoripenemGram +ve except MRSA, ESBL Gram -ve except Stenotrophomonas, Burkholderia, Corynebacterium, Enterococcus faecium not covered0.5gm -1gm q6h 1 - 2 gm q8hErtapenemDoes not cover Pseudomonas, Acinetobacter & Enterococcus1 - 2 gm q8hEnterococcus1 gm q24hPolymyxin BESBL, MetalloproteinasePolymyxin B 15Lac unit Loading f/b 7.5 lac unit BDPolymyxin BESBL, MetalloproteinaseColistin 4.5MUBD (loading dose of 9MU is mandatory 10xicityColistin Lincosamide ClindamycinRenal toxicityC. difficile		Amikacin	Gram –ve	3mg/kg q24h	Muscle
QuinolonesNalidixic acid Norfloxacin Ciprofloxacin Ofloxacin Levofloxacin Bextended spectrum1 gm q6h 400 mg q12h 500 mg q12h 200 mg q12h 750 mg q24h 400 mg q24hSeizuresExtended spectrumLevofloxacin MoxifloxacinGram +ve except MRSA, ESBL Gram -ve except Stenotrophomonas, Burkholderia, Corynebacterium, Enterococcus faecium not covered0.5gm -1gm q6h 1 - 2 gm q8hSeizuresErtapenemDoes not cover Pseudomonas, Acinetobacter & Enterococcus1 - 2 gm q8hMuscle weaknessPolymyxins Polymyxin BPolymyxin B 1 - 2 gm q24hMuscle weaknessPolymyxins Polymyxin BESBL, Metalloproteinase producing Gram -veColistin 4.5MUBD (loading dose of 9MU is mandatory 9MU is mandatoryMuscle weaknessLincosamide ClindamycinCoram +ve and anerobes600mg q8hdifficile		Tobramycin	Gram –ve	5mg/kg q24h	weaknes
Norfloxacin Ciprofloxacin Ciprofloxacin Ciprofloxacin Ciprofloxacin Coloxacin Coloxacin Coloxacin Coloxacin Moxifloxacin Carbapenems Gram +ve except MRSA, ESBL Gram –ve except Stenotrophomonas, Burkholderia, Corynebacterium, Enterococcus faecium not covered 1 – 2 gm q8h Folymyxin		Netilmicin	Gram -ve		S
Extended Levofloxacin Ofloxacin Moxifloxacin Carbapenems Imipenem- cilastatin Meropenem Doripenem Ertapenem Ertapenem Ertapenem Ertapenem Carbapenems Ertapenem Ertapenem Ertapenem Corynebacterium, Enterococcus faecium not covered Enterococcus Enterococcus Enterococcus Enterococcus Enterococcus Enterococcus Enterococcus Enterococcus Enterococcus Corynebacterium, Enterococcus Enterococcus Enterococcus Enterococcus Corynebacterium, Enterococcus Enterococcus Enterococcus Enterococcus Colistin ESBL, Metalloproteinase Colistin Colistin Corynebacter & Colistin 4.5MUBD (loading dose of opm of toxicity) Espace Acinetobacter Ertapenem Colistin Colistin Colistin Corynebacterium, Enterococcus Enterococcus Colistin Colistin Colistin Colistin Colistin Colistin Colistin Comp q8h Colificile	Quinolones	Nalidixic acid		1 gm q6h	
Extended spectrum		Norfloxacin		400 mg q12h	
Extended spectrum Moxifloxacin Moxifloxacin 750 mg q24h 400 mg q24h Carbapenems Imipenem- cilastatiin ESBL Gram -ve except MRSA, ESBL Gram -ve except Stenotrophomonas, Burkholderia, Corynebacterium, Enterococcus faecium not covered Ertapenem Does not cover Pseudomonas, Acinetobacter & Enterococcus 1gm q24h Polymyxins Polymyxin B Polymyxin B ESBL, Metalloproteinase ESBL, Metalloproteinase Colistin Producing Gram -ve foom g q24h Corynebacterium, Enterococcus faecium not cover Pseudomonas, Acinetobacter & Igm q24h Polymyxin B ISLac unit Loading f/b 7.5 lac unit BD Muscle weakness Colistin Forducing Gram -ve foom g q8h Colistin Corynebacterium, Enterococcus faecium not cover Pseudomonas, Acinetobacter & Igm q24h Polymyxin B ISLac unit BD Muscle weakness Colistin Forducing Gram -ve foom g q8h Colistin Colistin Colistin A.5MUBD (loading dose of g of g y y y y y y y y y y y y y y y y y y		Ciprofloxacin		500 mg q12h	Seizures
spectrumMoxifloxacin400 mg q24hCarbapenems Imipenem- cilastatinGram +ve except MRSA, ESBL Gram -ve except Stenotrophomonas, Burkholderia, Corynebacterium, Enterococcus faecium not covered1 - 2 gm q8hErtapenemDoes not cover Pseudomonas, Acinetobacter & Enterococcus1 gm q24hPolymyxins Polymyxin BPolymyxin B 15Lac unit Loading f/b 7.5 lac unit BDMuscle weaknessPolymyxin BESBL, MetalloproteinaseColistin 4.5MUBD (loading dose of 9MU is mandatory 9MU is mandatory 10 mandatory 9MU is mandatory 10 mandatory<		Ofloxacin		200 mg q12h	
Carbapenems Gram +ve except MRSA, ESBL Gram -ve except Seizures Meropenem Doripenem Stenotrophomonas, Burkholderia, Corynebacterium, Enterococcus faecium not covered 1 - 2 gm q8h Ertapenem Does not cover Pseudomonas, Acinetobacter & Enterococcus 1gm q24h Polymyxins Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 9MU is mandatory) Colistin 9MU is mandatory Lincosamide C. Clindamycin Gram +ve and anerobes 600mg q8h difficile	Extended	Levofloxacin		750 mg q24h	
Imipenem- cilastatin Meropenem Doripenem Doripenem Ertapenem Does not cover Pseudomonas, Acinetobacter & Enterococcus Enterococcus Enterococcus Enterococcus Enterococcus Enterococcus Enterococcus Enterococcus Colistin ESBL, Metalloproteinase Colistin ESBL, Metalloproteinase Colistin ESBL, Metalloproteinase Colistin ESBL, Metalloproteinase Colistin Colistin ESBL, Metalloproteinase Colistin ESBL, Metalloproteinase Colistin Colisti	spectrum	Moxifloxacin		400 mg q24h	
cilastatin ESBL Gram –ve except 1 – 2 gm q8h Meropenem Stenotrophomonas, 1 – 2 gm q8h Burkholderia, Corynebacterium, Enterococcus faecium not covered Ertapenem Does not cover Pseudomonas, Acinetobacter & Enterococcus 1 gm q24h Polymyxins B Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin Colistin 4.5MUBD (loading dose of 9MU is mandatory producing Gram –ve Renal toxicity Lincosamide Clindamycin Gram +ve and anerobes 600mg q8h difficile	Carbapenems				Seizures
Meropenem Doripenem Burkholderia, Corynebacterium, Enterococcus faecium not covered Does not cover Pseudomonas, Acinetobacter & Enterococcus Enterococcus Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin Producing Gram –ve Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Lincosamide Clindamycin Gram +ve and anerobes Comp q8h	_		- // V V	0.5gm -1gm q6h	
Burkholderia, Corynebacterium, Enterococcus faecium not covered Does not cover Pseudomonas, Acinetobacter & Enterococcus Igm q24h Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Lincosamide Clindamycin Corynebacterium, Enterococcus faecium not covered Does not cover Pseudomonas, Acinetobacter & Enterococcus Igm q24h Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity C. difficile	cilastatin				
Ertapenem Does not cover Pseudomonas, Acinetobacter & Enterococcus Enterococcus Igm q24h Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin Producing Gram –ve Polymyxin Acinetobacter & Colistin Acineto	Meropenem			1-2 gm q8h	
Ertapenem Does not cover Pseudomonas, Acinetobacter & Enterococcus Enterococcus Polymyxins Polymyxins Polymyxin B ESBL, Metalloproteinase Colistin Producing Gram –ve Polymyxin B Colistin Producing Gram –ve Polymyxins Producing Gram –ve Polymyxins Colistin A:5MUBD (loading dose of 9MU is mandatory Renal toxicity) Esmandatory Colistin Colindamycin Colidamycin Colimanycin Colimanycin Colimanycin Covered Does not cover Pseudomonas, Acinetobacter & Igm q24h Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Colistin A:5MUBD (loading dose of 9MU is mandatory Renal toxicity) Colimanycin Colimanyc	Doripenem				
Ertapenem Does not cover Pseudomonas, Acinetobacter & Igm q24h Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin Poducing Gram –ve Polymyxin B ESBL, Metalloproteinase Colistin Forducing Gram –ve Polymyxin B Colistin Forducing Gram –ve					
Ertapenem Does not cover Pseudomonas, Acinetobacter & Enterococcus 1gm q24h Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 9MU is mandatory toxicity Lincosamide Clindamycin Gram +ve and anerobes 600mg q8h Colistin C. difficile			Enterococcus faecium not		
Pseudomonas, Acinetobacter & Enterococcus 1gm q24h Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 4.5MUBD (loading dose of 9MU is mandatory toxicity Lincosamide Clindamycin CC. Gram +ve and anerobes Colographic form of the first of			covered		
Pseudomonas, Acinetobacter & Enterococcus 1gm q24h Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 4.5MUBD (loading dose of 9MU is mandatory toxicity Lincosamide Clindamycin CC. Gram +ve and anerobes Colographic form of the first of					
Acinetobacter & Enterococcus 1gm q24h Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Lincosamide Clindamycin Gram +ve and anerobes 600mg q8h difficile	Ertapenem	• X			
Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of loading dose of load			· ·		
Polymyxin B 15Lac unit Loading f/b 7.5 lac unit BD Muscle weakness Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Lincosamide Clindamycin Cam +ve and anerobes Colistin 4.5MUBD (loading dose of 9MU is mandatory for toxicity Cam +ve and anerobes Colistin 4.5MUBD (loading dose of 9MU is mandatory for toxicity Cam +ve and anerobes Colistin 4.5MUBD (loading dose of 9MU is mandatory for toxicity Cam +ve and anerobes					
Polymyxins Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Lincosamide Clindamycin Company to the control of the co		() y		,	
Polymyxins Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Lincosamide Clindamycin Gram +ve and anerobes Colistin 4.5MUBD (loading dose of 9MU is mandatory toxicity C.					
Polymyxins Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Lincosamide Clindamycin Gram +ve and anerobes Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity C.		Y		_	M1-
Polymyxin B ESBL, Metalloproteinase Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Lincosamide Clindamycin Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Colistin 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity	Polymyving			Iac unit BD	
Colistin producing Gram –ve 4.5MUBD (loading dose of 9MU is mandatory Renal toxicity Lincosamide Clindamycin Gram +ve and anerobes 600mg q8h difficile		\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	ESBI Metalloprotoinese	Colictin	weakiiess
Colistin producing Gram –ve (loading dose of 9MU is mandatory Renal toxicity Lincosamide Clindamycin Gram +ve and anerobes 600mg q8h difficile	1 Olymyxiii D		ESBE, Micianoprotemase		
Colistin producing Gram –ve 9MU is mandatory Renal toxicity Lincosamide Clindamycin Gram +ve and anerobes 600mg q8h difficile					
LincosamideC.ClindamycinGram +ve and anerobes600mg q8hdifficile	Colistin		producing Gram _ve		Renal
LincosamideC.ClindamycinGram +ve and anerobes600mg q8hdifficile	Consun			71410 is mandatory	
Clindamycin Gram +ve and anerobes 600mg q8h difficile	Lincocamido				•
			Gram +ve and anerobes	600mg a8h	
I I Colific	Cilidaniyeni		Grain +ve and ancroves	ooonig qon	colitis

	1	1,,	
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Laiwani Medical Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 72 of 8



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Antimicrobial			Indication&	Side
Class	Name	Organisms	Dose	effects
Glycopeptides				Renal
Vancomycin		MRSA	1gm q12h	toxicity
			400 mg 3	•
			doses f/b 400	
Teicoplanin			mg	
			q24h	
Oxazolidinedione				Thromb
Linezolid		VRE	600 mg	ocytope
			q12h	nia
Lipopeptides			4-6mg/kg	
Daptomycin		MRSA	q24h	
Antifungals				
Fluconazole		Candida albicans	400 mg	
*7 ' 1			q12h	
Voriconazole		Aspergillus	6mg/kg	
		3	q12h first	
			day then	
Comofinacia		Non albicans candida	4mg/kg	
Caspofungin		Non afficans candida	70mg IV then 50 mg	
			q24h	
Anidulafungin		Non albicans candida	q2411	
Amadiardigiii		Tion aroleans candida		
AmphoB aqueous		Broad spectrum covers all	Refer	
AmphoB colloidal	•)	above + Mucor etc	product	
AmphoB liposomal			insert	
r r				
	(O 2)			,

				/ '
Prepared by	Checked by	Verified by	Appr	eved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department		K Natwani cal Director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 73 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026

Version - 11.0

Appendix 2 Duration of therapy for various clinical conditions

Short course therapy is equally	v effective		
Condition	Short Course (Days)	Long Course (Days)	Outcome
Chronic Bronchitis and	<=5	>=7	Equivalent
obstructive pulmonary			
disease, acute exacerbations			Most patients do not require antimicrobials at all
			require anumicrobiais at an
Pneumonia community	<=8	10-15	Equivalent
acquired		11.01	
Possible pneumonia NICU	3	14-21	Equivalent
Atypical CAP	1	3	Equivalent
VAP	5-8	10-15	Equivalent
Empyema	14-21	21-42	Equivalent
Neutropenic fever	Until afebrile and stable	Until non neutropenic	Equivalent
Osteomyelitis, Chronic	42	84	Equivalent
Osteomyelitis removed	28	42	Equivalent
implant Debrided diabetic osteo	10-21	42-90	Equivalent
Debrided diabetic osteo	10-21	42-90	Equivalent
Septic arthritis	14	28	Equivalent
Intra abdominal infection	4	8-10	Equivalent
Complex appendicitis	1-2	5-6	Equivalent
Pyelonephritis /cUTI	5-7	10-14	Equivalent
Skin Infections (Cellulitis,	5-6	10-14	Equivalent
Major abscesses, wound infections)	40		
Sinusitis, acute bacterial	5	10	Equivalent
Bacteremia (Non Staph aureus)	7	14	Equivalent
Bacterial meningitis(peds)	4-7	7-14	Equivalent
Cystic fibrosis exacerbation	10-14	14-21	Equivalent
NT. 4			

Note

- Recommended duration of therapy for candidemia without persistent fungemia or metastatic complications is for 2 weeks after documentation of negative cultures of candida from the bloodstream. (IDSA). Repeat blood culture after three days of starting therapy
- In cases of CRE, MRSA, CRAB bacteremia, repeat blood culture after 3 days of initiating appropriate antibiotics as per AST

			/ '	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Ouality Department	Dr. SK halwani Medical Director	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 74 of 81



Prepared by

Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

BHRC / HIC / F23 / (V-3)

Appendix 3 Antimicrobial Agent Form

PICU/NICU/1/2/3/6/8 inician/Unit Head : Patient ty	7pe: 1 2 3 V	Wt. of Patientkg	minated O	atient code here
Patient ty Type CT CRP/PCT Route, Freq and	rpe: 1 2 3 V	Wt. of Patientkg	minated O	Dirty DATE
Patient ty Type CT CRP/PCT Route, Freq and	rpe: 1 2 3 V	Wt. of Patientkg	minated O	Dirty DATE
Patient ty Type CT CRP/PCT Route, Freq and	rpe: 1 2 3 V	Wt. of Patientkg	minated O	Dirty DATE
Type CT CRP/PCT Route, Freq and	rpe: 1 2 3 V :: Clean Cle RADIOL	Wt. of Patientkg	minated O	Dirty DATE
Type CT CRP/PCT Route, Freq and	RADIOI	ean cont. Contar	minated O	Dirty DATE
Type CT CRP/PCT Route, Freq and	RADIOI	ean cont. Contar	IONS WITH	H DATE
CT CRP/PCT	RADIOL Start Stop	ean cont. Contar	IONS WITH	H DATE
CT CRP/PCT	RADIOI	LOGICAL EXAMINATI	IONS WITH	H DATE
CT CRP/PCT	RADIOI	LOGICAL EXAMINATI	IONS WITH	H DATE
Route, Freq and	Start Stop			
Route, Freq and	Start Stop			
	CONT. 100 CONT.	ID ref/AMSP Advio	ce	IV to
	CONT. 100 CONT.	ID ref/AMSP Advio	ce	IV to
	CONT. 100 CONT.	ID ref/AMSP Advio	ce	IV to
	CONT. 100 CONT.	ID ref/AMSP Advice	ce	IV to
				Oral
아이는 아이들이 아이를 보면서 모양에 되어 있다.		TT 4. Central Line	5. Drains	6. HD Cathete
roorganism isolated received (DD/MM/)	Sensiti		Change of AMA Y/N	Response after change Y/N
	roorganism isolated	roorganism isolated Sensit	roorganism isolated Sensitivity Pattern	roorganism isolated Sensitivity Pattern Change received (DD/MM/) (Name of imp antibiotics) of AMA

Dr. M S Modak
Prof & HoD
Microbiology

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 75 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

1) Appropriate	Score	Description	
	A	Empiric	
	В	Definitive	
	С	Surgical prophylaxis	
	D	De-escalation/Escalation	
2) Inappropriate	A	No Indication	
	В	Empiric	
	С	Definitive (No de-escalation/ escalation based on culture/advice)	
	D	Surgical Prophylaxis	
	Е	Duration longer than required	
	F	Duration longer for surgical prophylaxis	
	G	Dose/Freq (underdosing/overdosing)	
	Н	Irrational Combination	
3) Others	A	Indication not documented	
	В	ID reference	
	С	Referred from outside	
	D	Readmitted	

^{*} E- Empiric, D-Definitive, SP- Surgical prophylaxis, MP- Medical prophylaxis

List of high-end antibiotics:

- 1. Carbapenems 2. Piptaz 3. Levofloxacin 4. Colistin / Polymyxin-B 5. Fosfomycin 6. Daptomycin
- 7. Teicoplanin 8. Vancomycin 9. Tigecycline/Minocycline 10. Linezolid 11. Echinocandins
- 12. Voriconazole/Posaconazole 13. Amphotericin B 14. Cefta/Avi Aztreonam

				/ '
Prepared by	Checked by	Verified by	Appr	oved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Ouality Department		K Latwani

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 76 of 81

^{**} B - Blood, P - PUS, U - Urine, Resp - Sputum, ETT, BAL, CSF. - Cerebro Spinal Fluid, F - Fluids.



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Appendix 4

Operative Procedure for Sample Collection

1. URINE CULTURE

(a) Items required

The following items should be available as preparation for collecting a catheter specimen of urine for analysis:

- · Sterile gloves;
- Alcohol-saturated swab;
- Gate clip or non-traumatic clamps;
- Sterile Universal specimen container;



(b) Collection of urine sample from indwelling

Use sterile precautions

Clean hands using hand rub

Clamp the distal portion of the catheter.

Disconnect urine bag.

Disinfect distal portion of the catheter using 70% alcohol swab. Allow it to dry thoroughly.

Do not allow the distal end of the catheter to touch body or clothes of the patient.

After 10 minutes release the clamp and collect urine in a sterile urine collection container

(c) Urine must be transported to the lab without delay. If delay is unavoidable, sample should be stored in the refrigerator at 4°C.

catheter:



Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 77 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

2. BLOOD CULTURE

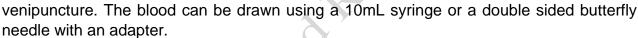
(a) Hands should be clean and dry and gloved before taking sample.







- (b) Prepare Blood culture vials by proper labeling
- (c) The culture bottle should be brought to room temperature before the sample inoculation and should not be refrigerated after inoculation.
- (d) Remove flip-off caps and wipe the rubber bung with a 70% alcohol swab
- (e) Disinfect skin at venepuncture site by wiping with 70% alcohol in a circular motion from centre to periphery and then with 1% iodine. Wait for at least one minute, allow to dry before











- (f) After venepuncture carefully withdraw the needle and compress the venepuncture site.
- (g) The number of sets to be collected is as follows.
 - (i) Febrile episode: 2 bottles from separate sites within 10 min
 - (ii) Acute endocarditis: 6 bottles from separate sites at 30 min intervals
 - (iii) Central Line related: 1 bottle drawn from the central line and 1 bottle from a peripheral venipuncture site
- (h) The various culture bottles for Automated Blood Culture System and the volume of blood to be added to them are as under.
- (i) BacT/Alert/BACTEC Aerobic (30 ml): 10 ml blood (optimal)
 - (ii) BacT/Alert/BACTEC Paediatric (20 ml): 4 ml blood (optimal)

	1	1,, .,, .,	- Land bu	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Latwani Medical Director	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 78 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Vei

3. ENDOTRACHEAL/TRACHEOSTOMY ASPIRATE CULTURES

- (a) The collection of endotracheal aspirate will be performed by the Anaesthetist or the Intensivist.
- (b) Do not use a swab for collecting sample.
- (c) Use a BAL collection trap to collect the endotracheal aspirate.
- (d) The aspirate is to be drawn as such in case it is less tenacious or alternatively, 5 ml of sterile saline is instilled and withdrawn immediately through the disposable sterile endotracheal aspiration catheter.



(e) The material is to be sent to laboratory within the next one hour.

4. PUS CULTURE

(a) Clean surface of wound or abscess with 70% alcohol and allow to dry; aspirate pus or fluid from the depth of the wound in a place in a sterile container and laboratory.

disposable syringe, send to the

Cotton swab to be used only if pus cannot be collected in sterile bottle or syringe.



5. Body fluids

Sample-Amniotic fluid, Synovial fluid, pericardial fluid, Pleural fluid, peritoneal fluid Sterile fluids are usually collected by a trained, qualified physician.

Aseptically collect at least 1 ml of fluid into a new disposable sterile container.

6. Bronchoalvelar Lavage

BAL should be collected under aseptic conditions preferably with a protected specimen brush if available. The material should be collected in a BAL trap container.

7. High vaginal swabs (HVS)

Use speculum to separate the vaginal walls.

Wipe away any excess cervical mucus with a sterile cotton swab.

Use the sterile swab stick to collect sample as high as possible in the vaginal vault.

				/ '
Prepared by	Checked by	Verified by	Appr	oved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Ouality Department		K Latwani

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 79 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

Appendix 5

List of BL-BLI combination not recommended by WHO

- amoxicillin/sulbactam
- cefadroxil/clavulanic acid
- cefepime/sulbactam
- cefepime/tazobactam
- cefixime/clavulanic acid
- cefixime/clavulanic acid/lactobacillus acidophilus
- cefotaxime/sulbactam
- cefpodoxime proxetil/clavulanic acid
- cefoperazone/tazobactam
- ceftazidime/sulbactam
- ceftazidime/tazobactam
- ceftriaxone/sulbactam
- ceftriaxone/tazobactam
- cefuroxime axetil/clavulanic acid
- cefuroxime/clavulanic acid
- cefuroxime/sulbactam
- meropenem/sodium/sulbactam
- meropenem/sulbactam
- piperacillin/sulbactam

			/ '
Prepared by	Checked by	Verified by	Approved by
Dr. M Kumar Prof Microbiology	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK halwani Medical director

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 80 of 81



Antimicrobial Policy and Antimicrobial Stewardship Program

2025-2026 Version - 11.0

References:

- 1. National treatment guidelines for antimicrobial use in infectious disease, version 1 (2016)
- 2.ICMR Treatment guidelines for antimicrobial use in common syndromes, 2022 2nd edition
- 3.ICMR Antimicrobial Stewardship Program Guideline 2018
- 4. Antimicrobial stewardship programmes in health-care facilities in low-and middle-income countries: a WHO practical toolkit 2019
- 5.Antimicrobial stewardship: Systems and processes for effective antimicrobial medicine use. NICE guideline 2015 (updated Jan 2018)
- 6.IDSA: New Antibiotic Stewardship Guidelines Focus on Practical Advice for Implementation 2016
- 7. Harrison's Principles of Medicine 20th ed
- 8. ASGE guidelines for antimicrobial prophylaxis in GI Endoscopic procedures
- 9. Urinary Tract Infections in Pregnant Individuals. Obstet Gynecol. 2023 Aug 1;142(2):435-445. doi: 10.1097/AOG.000000000005269. PMID: 37473414.
- 10. Blomstro m-Lundqvist C, Traykov V, Erba PA, Burri H, Nielsen JC, Bongiorni MG et al. European Heart Rhythm Association (EHRA) international consensus document on how to prevent, diagnose, and treat cardiac implantable electronic de- vice infections-endorsed by the Heart Rhythm Society (HRS), the Asia Pacific Heart Rhythm Society (APHRS), the Latin American Heart Rhythm Society (LAHRS), International Society for Cardiovascular Infectious Diseases (ISCVID), and the European Society of Clinical Microbiology and Infectious Diseases (ESCMID) in collaboration with the European Association for Cardio-Thoracic Surgery (EACTS). Europace 2020;22:515–49.
- 11. 2019 WHO AWaRe Classification Database of Antibiotics for evaluation and monitoring of use
- 12. el Moussaoui R, de Borgie CA, van den Broek P, et al. Effectiveness of discontinuing antibiotic treatment after three days versus eight days in mild to moderate-severe community acquired pneumonia: randomized, double blind study. Bmj 2006; 332(7554): 1355.
- 13. Schonwald S, Kuzman I, Oresković K, et al. Azithromycin: single 1.5 g dose in the treatment of patients with atypical pneumonia syndrome--a randomized study. Infection 1999; 22(3):198-202.\
- 14. Chastre J, Wolff M, Fagon JY, et al. Comparison of 8 vs 15 days of antibiotic therapy for ventilator-associated pneumonia in adults: a randomized trial. JAMA: the journal of the American Medical Association 2003; 290(19): 2588-98.
- 15. Drekonja DM, Trautner B, Amundson C, et al. Effect of 7 vs 14 Days of Antibiotic Therapy on Resolution of Symptoms Among Afebrile Men With Urinary Tract Infection: A Randomized Clinical Trial. JAMA 2021; 326(4):324-331.
- 16. Bernard L, Dinh A, Ghout I, et al. Antibiotic treatment for 6 weeks versus 12 weeks in patients with pyogenic vertebral osteomyelitis: an open-label, non-inferiority, randomized, controlled trial. Lancet 2015. 385:875-82.
- 17. Yahav D, Franceschini E, Koppel F, et al. Seven versus fourteen Days of Antibiotic Therapy for uncomplicated Gram-negative Bacteremia: a Non-inferiority Randomized Controlled Trial. Clin Infect Dis 2019 69:1091-8.

			,	
Prepared by	Checked by	Verified by	Approved by	
Dr. M Kumar	Dr. M S Modak Prof & HoD Microbiology	Dr. Shilpa Acharya Quality Department	Dr. SK Nalwani Medical Nirector	

Antimicrobial Policy Ver.11 Date of release: 01.04.2025 Valid till Mar 2026 unless amended earlier Page 81 of 81