

Bharati Vidyapeeth University Medical College Hospital & Research Centre, Pune Antimicrobial Policy and Antimicrobial Stewardship Program 2024-2025 Version - 10.0



Prepared by	Checked by	Verified by	Apprøved by
Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Arr	Dr sk tulwani wam Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2024-2025 Version - 10.0

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Air	Dr SK I	lwani lwani Director

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List of amendments in Version 10.0

- 1. Isolate listing and Antibiogram for the year 2023
- 2. Antibiogram : Sample type Pus has been separated into Wound and Tissue for Surgical and Orthopaedic specialities
- 3. Added: OPD cases of Acute gastroenteritis suggested antibiotic if required
- 4. UTI in Obst-Gynae : Added Table 4a Cefixime and Fosfomycin after 20wk pregnancy (added reference No 9)
- 5. Updated vancomycin dosing for surgical prophylaxis in Table 16
- 6. Added Table 20 for Cardiac device implantation prophylaxis (added reference No 10)
- 7. Renumbered older table No 20 to Table 21 (Spectrum of commonly used antimicrobials)
- 8. Added sample collection method for High Vaginal Swab Appx 4

Dr M Kumar
Prof Microbiology

Checked by

Verified by

Dr AK Verma Quality
Dr AK Verma Quality
Department by Ned Ar



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1. Introduction -

Over the last 60 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 re-administration 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.

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

Dr M Kumar
Prof Microbiology

Checked by

Verified by

Dr AK Verma Quality
Dr AK Verma Quality
Department by Ned Art

Medical Director



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2. Clinical Pathway

- 1. 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.
 - 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.
- 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 requires 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.
- 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. Infection control team will fill antimicrobial audit form and conduct regular department wise audits.
- 11. Findings of the audit will drive improvement in antimicrobial use.

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Air	Dr skrijiwani wam Medical Director



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3. Antimicrobial Stewardship



Antibiotic Stewardship



A set of coordinated strategies to improve the use of antimicrobials

Goal

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

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Checked by

Verified by

Dr M S Modak Prof Code

Dr AK Verma Quality

Prof Microbiology

Dr AK Verma Quality

Department by Ned Avi

Medical Director

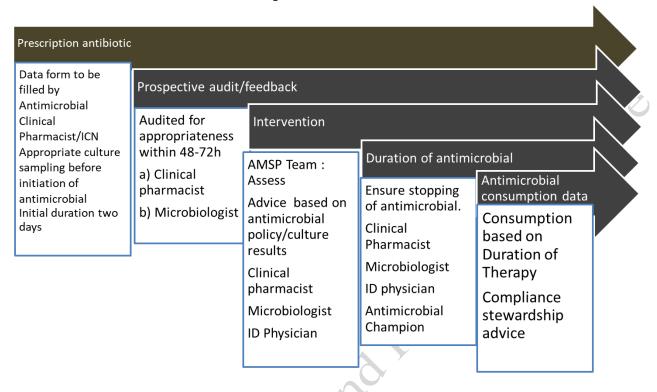


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Workflow of Antimicrobial stewardship





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

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by N. d Dir	Dr sk Tulwani Lvam Medical Director

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4. **List of Restricted Antimicrobials**

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

- 1. Piperacillin tazobactam
- 2. Carbapenems
- 3. Linezolid
- 4. Vancomycin
- 5. Teicoplanin
- 6. Daptomycin
- 7. Tigecycline
- 8. Polypeptides: Polymixin, Colistin
- 9. Echinocandins: Caspofungin, Micafungin
- 10. Ceftazidime avibactam
- 11. Aztreonam

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 penicillins; first-, second and third-generation cephalosporins; In addition, the plasmids bearing genesencoding ESBLs frequently also carry genes encoding 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.

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Arr	Dr sktolwani wam Medical Director



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MDR (Multi-drug resistance):

Isolates resistant to representatives of three or more classes of antimicrobial agents, 3harati Hospital and Research Centre

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

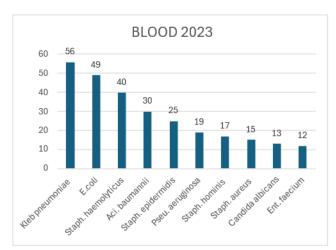
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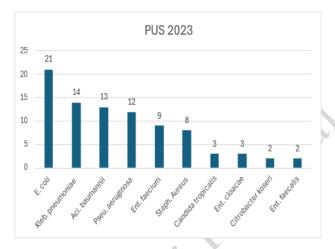


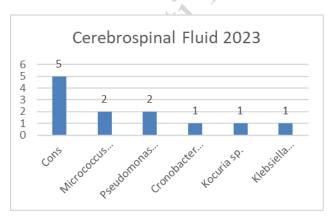
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5. Organisms commonly isolated and antibiogram: Area wise

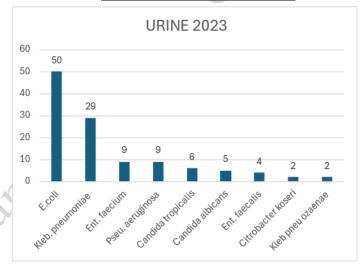
INTENSIVE CARE UNIT: TOP 10 ISOLATES BY SAMPLE

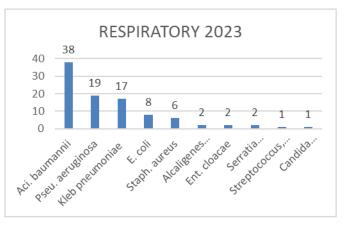






SPECIMEN	OVERALL NO. OF
TYPE	ISOLATES
BLOOD	442
PUS	104
RESPIRATORY	108
URINE	137
CSF	12





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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Dir	Dr SK Tulwani Wani Medical Director



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INTENSIVE CARE UNIT ANTIBIOGRAM GRAM NEGATIVE

Percentage susceptibility gram negative organism 2023

Location	Specimen type	Organism	Number of patients	LEVOFLOXACIN	CIPROFLOXACIN	COTRIAMOXAZOLE	AMOX/CLAV	Cefuroxime	CEFTAZIDIME	CEFTRIAXONE	CEFOTAXIME	CEFEPIME	CEF/SUL	PIPTAZ	AMIKACIN	GENTAMICIN	ERTAPENEM	MEROPENEM	IMIPENEM	TIGECYCLINE	FOSFOMYCIN	MINOCYCLINE	COLISTIN
		Klebsiella pneumoniae	56	0	27	44	29	36		27		34	36	36	47	47	44	46	47		80		100
	Blood	Escherichia coli	49		6	50	48	63		25		42	63	63	85	77	73	85	85		98		100
		Acinetobacter baumannii	30		10	33		10		10		7	10	10	10	13		13	10				100
		Burkholderia cepacia	12	80	0	90		20	60	0		0	20	20	40	20		80	20			100	
	Urine	Escherichia coli	50		4	58	38	58		24		30	58	58	80	74	74	78	76		98		100
		Klebsiella pneumoniae	29		14	55	28	28		21		28	28	28	38	45	41	45	41		79		100
	Respiratory	Acinetobacter baumannii	38		5	29		5		3		5	5	5	5	8		5	5				100
		Klebsiella pneumoniae	17		31	56	38	31		6		19	31	31	38	38	50	44	44		81		100
SSICU		Escherichia coli	21		0	43	19	0		5	0	24	57	57	91	91	62	71	71		91		100
	Pus	Klebsiella pneumoniae	14		14	50	29	14		14	14	21	36	36	43	43	36	36	36	36	43		100
		Acinetobacter baumannii	13		0	8				0	0	0	0	0	0	0		0	0		0		92
		Acinetobacter baumannii	10		10	30		10		10		10	10	10	10	10		10	10				100
	Other	Klebsiella pneumoniae	7		14	71	29	43		29		57	43	43	71	71	71	71	57	57	100		100
		Escherichia coli	5		0	50	50	60		0		20	60	60	50	0	50	50	80		100		100
	Fluid	Escherichia coli	12		10	40	36	67		20		58	67	67	90	80	80	90	92		91		100
	Cerebrospinal	Pseudomonas aeruginosa	2	50				100				100	100	100				50	0				100
	fluid	Cronobacter sakazakii	1		0	100	100			###					100	100	100	100			100		

Reserved / Restricted Drugs :Not to be used empirically 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

Most organisms other than Enterobacterales and Acinetobacter have not reached the statistically significant number of 30 isolates. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Dr M Kumar
Prof Microbiology

Checked by

Verified by

Dr AK Verma Quality
Dr AK Verma Quality
Department by Ned Ar

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INTENSIVE CARE UNIT ANTIBIOGRAM GRAM POSITIVE

Percentage susceptibility gram positive organism 2023

Location	Specimentype	Organism	Number of patients	NITROFURANTOIN	CEFOXITIN	OXACILLIN	PENICILLIN	CIPROFLOXACIN	LEVOFLOXACIN	MOXIFLOXACIN	CLINDAMYCIN	GENTAMICIN	GENTAMICIN HIGH LEVEL	ERYTHROMYCIN	TETRACYCLINE	Sulfa/Trimeth	TIGECYCLINE	DAPTOMYCIN	LINEZOLID	TEICOPLANIN	VANCOMYCIN
		Staphylococcus haemolyticus	40	97	8	8	5	8	8		40	21		3	68	53		92	90	95	100
		Staphylococcus epidermidis	25	96	4	4	0	24	24		52	56		16	64	52		92	96	92	100
	Blood	Staphylococcus hominis	17	87	21	20	13	40	40		80	93		7	67	67		93	93	80	100
	Dioou	Staphylococcus aureus	15	100	43	43	7	7	7		86	57		36	86	43		100	100	100	100
		Enterococcus faecium	12	8			0	0	0				100	0	0				92	75	75
		Staphylococcus capitis	7	100	86	86	29	100	100		100	100		86	86	100		86	100	100	100
SSICU	Urine	Enterococcus faecium	9	0			0	0	0				100	0	11				78	100	67
	D	Enterococcus faecium	9				0			0		0		100	0		100			100	100
	Pus	Staphylococcus aureus	8			38	0			100	100	0			100	63	100			100	100
	Respiratory	Staphylococcus aureus	6	100	67	67	33	0	0		100	100		33	83	33	100	100	100	100	100
		Staphylococcus haemolyticus	2	100	0	0	0	0	0		0	0		0	50	100	100	100	100	100	100
	Cerebrospinal fluid	Staphylococcus hominis	1	100	0	0	0	0	0		0	100		0	0	0	100	100	100	100	100
	rtala	Staphylococcus cohnii	1	0	0	0	0	0	0		0	100		0	100	100	100	100	0	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

Except Staphylococcus haemolyticus all other isolates have not reached the statistically significant number of 30 isolates. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern. Staphylococcus species other than S aureus are considered collection contamination/skin flora.

Prepared by

Checked by

Verified by

Approved by

Dr M S Modak Prof

Prof Microbiology

Dr AK Verma Quality

Department by Ned Air

Medical Director

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2024-2025

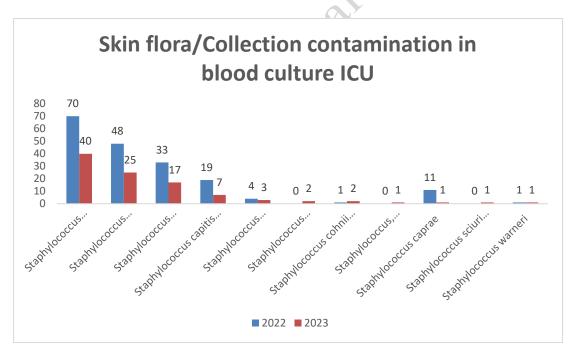
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Candida isolated in ICU

Location	Specimen type	Organism	Number of patients	FLUCONAZOLE	VORICONAZOLE	AMPHO B	CASPOFUNGIN	MICAFUNGIN
		Candida albicans	13	62	77	31	100	92
	Blood	Candida parapsilosis	8	43	57	57	100	100
	blood	Candida tropicalis	8	29	75	75	100	100
		Candida glabrata	1		0	100	100	100
	Pus	Candida tropicalis	3	100	100	100	100	100
SSICU	rus	Candida parapsilosis	1	0	0	100	100	100
	Respiratory	Candida tropicalis	1	100	100	100	100	100
		Candida tropicalis	6	60	80	80	100	100
	Urine	Candida parapsilosis	1	0	0	100	100	100
	Offile	Candida dubliniensis	1	0	0	100		
		Candida albicans	5	80	80	40	100	100

Please note: Individual isolates are less than 30 in number, not statistically significant. Antibiogram has been shown with available number of organisms.

Skin flora/Collection contamination in blood culture ICU



Year	No of
	Contaminants
	isolated
2022	187
2023	100

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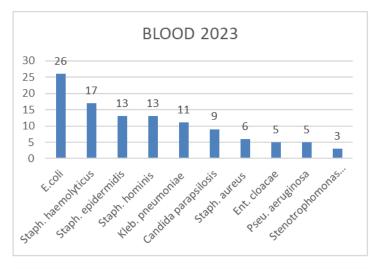


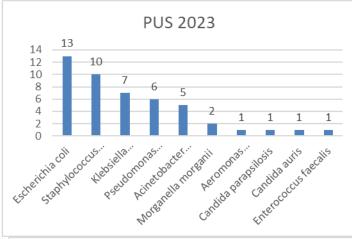
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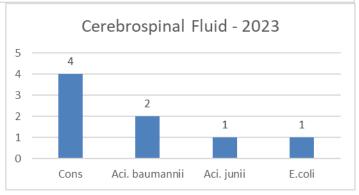
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MEDICINE

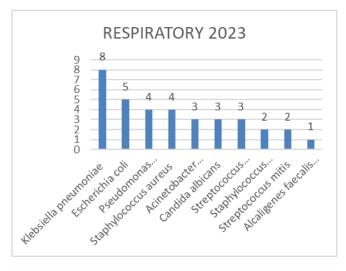
TOP 10 ISOLATES BY SAMPLE

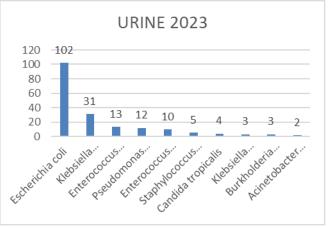






SPECIMEN	OVERALL NO. OF
TYPE	ISOLATES
BLOOD	166
PUS	56
RESPIRATORY	44
URINE	206
CSF	08





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Dr M Kumar Prof Microbiology	Dr M S Modak Prof Microbiology	Dr AK Verma Quality Department by Ned Ar	Dr sk iklwani wam Medicai Director

BHARAT

BHARATI VIDYAPEETH UNIVERSITY MEDICAL COLLEGE HOSPITAL AND RESEARCH CENTRE

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MEDICINE ANTIBIOGRAM GRAM NEGATIVE

Percentage Susceptibility Gram Negative Organism 2023

Location	Specimen type	Organism	Number of patients	LEVOFLOXAGN	NORFLOXACIN	GPROFLOXAGN	OFLOXACIN	NITROFURANTOIN	COTRIAMOXAZOLE	AMOX/CLAV	ŒFEPIME	ŒF/sur	ŒFUROXIME	ŒFTAZIDIME	ŒFTRIAXONE	PIPTAZ	AMIKAGN	GENTAMICIN	MEROPENEM	ERTAPENEM	IMIPENEM	MINOCYCLINE	TIGECYCLINE	FOSFOMYON	COLISTIN
	Blood	Escherichia coli	26			4			50	62	23	69	19		19	69	92	89	89	89	89			100	100
	Union	Escherichia coli	102		67	11	67	100	49	53	38	71	15	67	24	71	83	71	85	86	86			99	100
	Urine	Klebsiella pneumoniae	31	100	0	19	0	0	45	27	23	32	17	0	13	32	39	52	37	33	33	100		61	100
Medicine	Danimatani	Klebsiella pneumoniae	8			50			63	63	75	75	50		63	75	75	75	75	75	75		75	75	88
	Respiratory	Escherichia coli	5			0			60	20	40	20	20		20	20	60	60	60	60	60			100	100
	Dur	Escherichia coli	13			0			25	50	42	58	17		17	58	83	92	67	64	67			100	100
	Pus	Acinetobacter baumannii	5			0			20		0	0				0	0	0	0		0				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

Most organisms other than Enterobacterales have not reached the statistically significant number of 30 isolates. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

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MEDICINE ANTIBIOGRAM GRAM POSITIVE

Percentage Susceptibility Gram Positive Organism 2023

Location	Specimen type	Organism	Number of patients	OXACILLIN	CEFOXITIN	LEVOFLOXACIN	CIPROFLOXACIN	COTRIAMOXAZOLE	NITROFURANTOIN	PENICILLIN	GENTAMICIN	GENTAMICIN HIGH LEVEL	ERYTHROMYCIN	CLINDAMYCIN	TETRACYCLINE	TIGECYCLINE	LINEZOLID	DAPTOMYCIN	TEICOPLANIN	VANCOMYCIN
		Staphylococcus haemolyticus	17	6	0	0	0	40		0	33		13	47	80		87	94	93	93
	Blood	Staphylococcus epidermidis	13	23	23	23	23	69		8	69		23	46	69		100	100	92	100
		Staphylococcus hominis	13	25	27	33	33	42		8	75		0	58	67		92	83	83	92
		Staphylococcus aureus	6	33	33	17	17	67		0	50		33	83	83		100	100	100	83
Medicine		Enterococcus faecium	13			0	0		8	0		100	0		0		92		83	77
	Urine	Enterococcus faecalis	10			10	10		90	100		100	0		0		100	90	100	100
		Staphylococcus aureus	5	60	60	0	0	100	100	20	100		40	80	100		100	80	100	80
	Pus	Staphylococcus aureus	10	30	30	10	20	40		0	100		20	80	100	100	100	90	100	100
	Fluid	Staphylococcus haemolyticus	5	0	0	20	20	60		0	40		0	40	60	80	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.

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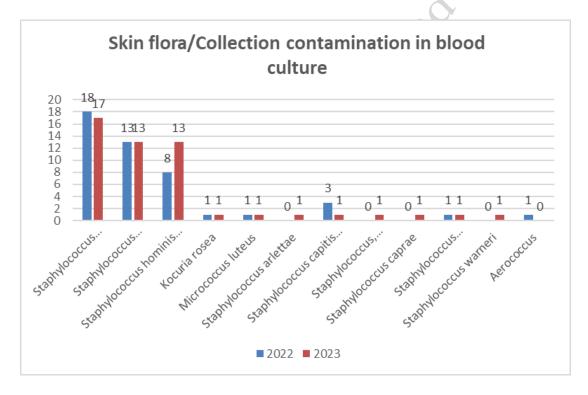
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CANDIDA ISOLATED IN MEDICINE

Location	Specimen type	Organism	Number of patients	FLUCONAZOLE	VORICONAZOLE	CASPOFUNGIN	MICAFUNGIN	АМРНО В
		Candida parapsilosis	9	67	78	100	100	89
	Blood	Candida albicans	2	100	100	50	50	100
		Candida tropicalis	1	0	0	100	100	0
	Urine	Candida tropicalis	4	100	100	100	100	100
Medicine	Urine	Candida albicans	2	100	100	100	100	0
	Dospiratory	Candida albicans	3	100	100	100	100	33
Respiratory		Candida tropicalis	1	0	100	100	100	100
	Fluid	Candida albicans	3	100	100	100	100	67
	Wound	Candida parapsilosis	1	0	100	100	100	100

Candida

Please note: Individual isolates are less than 30 in number, Not statistically significant. Antibiogram has been shown with available number of organisms.



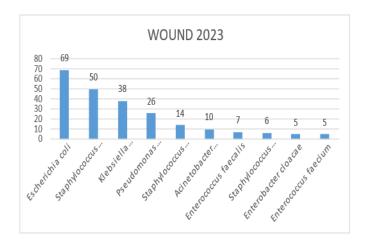
		No of
,	Year	Contaminants isolated
	2022	46
:	2023	51

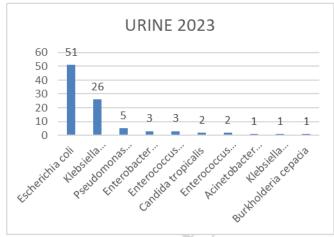
Prepared by	Checked by	Verified by 1 1	Apprøved by
Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Air	Dr sk thiwani wam Medical Director

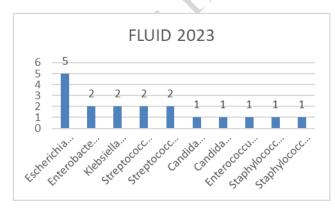


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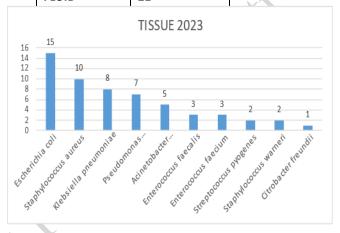
SURGERY: TOP 10 ISOLATES BY SAMPLE

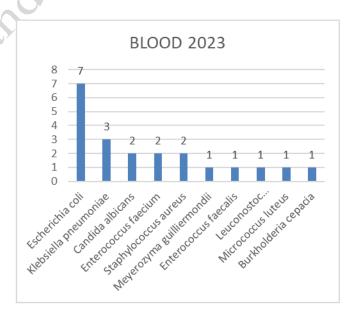






SPECIMEN	OVERALL NO. OF
TYPE	ISOLATES
BLOOD	26
WOUND	297
TISSUE	65
URINE	103
FLUID	21





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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Dir	Dr sk tellwani wam Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2024-2025

Version - 10.0

SURGERY ANTIBIOGRAM GRAM NEGATIVE

Percentage Susceptibility Gram Negative Organism 2023

Location	Specimen type	Organism	Number of patients	CIPROFLOXACIN	LEVOFLOXACIN	NORFLOXACIN	OFLOXACIN	FOSFOMYCIN	COTRIMOXAZOLE	NITROFURANTOIN	AMPICILLIN	AMOX/CLAV	CEFEPIME	CEFUROXIME	CEFTAZIDIME	CEFTRIAXONE	CE/SUL	PIPTAZ	AMIKACIN	GENTAMYCIN	MEROPENEM	IMIPENEM	ERTAPENEM	MINOCYCLINE	TIGECYCLINE	COLISTIN
	Blood	Escherichia coli	7	14					86			57	14	0		0	0	86	100	71	100	100				100
		Escherichia coli	69	4					42			49	29	12		17	0	64	88	67	84	84	81			100
	Wound	Klebsiella pneumoniae	38	40	0				61			51	53	38	0	46	0	53	66	66	63	61	65	100	58	100
		Pseudomonas aeruginosa	26	65	69								77		69		0	65	81	81	73	77				100
		Acinetobacter baumannii	10	0					44				0			0	0	0	0	0	0	0				100
C		Escherichia coli	15	0					33			47	20	0		0	0	60	73	80	80	80	73			100
Surgery	Tissue	Klebsiella pneumoniae	8	25					63			50	50	25		38	0	50	50	50	50	50	50		63	100
		Acinetobacter baumannii	5	20					40				20			0	0	20	20	20	20	20				100
	Urine	Escherichia coli	51	6		50	50	100	53	50	0	43	32	9	25	24	0	62	84	73	85	85	84			100
	Offile	Klebsiella pneumoniae	26	23		100	100	77	46	50		27	21	17	100	27	0	31	46	42	42	38	46			100
	Fluid	Escherichia coli	5	0					60			80	20	0		0	0	80	100	40	80	80	80			100
	Other	Escherichia coli	5	0					20			40	20	20		20	0	60	80	40	60	60	60			100
			265																							

_

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

Most organisms other than Enterobacterales have not reached the statistically significant number of 30 isolates. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Air	Dr SKTOWani WAYN Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2024-2025

Version - 10.0

SURGERY ANTIBIOGRAM GRAM POSITIVE

Percentage Susceptibility Gram Positive Organism 2023

Location	Specimen type	Organism	Number of patients	COTRIAMOXAZOLE	NITROFURANTOIN	CEFOXITIN	OXACILLIN	PENICILLIN	GENTAMICIN	GENTAMICIN HIGH LEVEL	ERYTHROMYCIN	CLINDAMYCIN	CIPROFLOXACIN	LEVOFLOXACIN	TIGECYCLINE	TETRACYCLINE	DAPTOMYCIN	LINEZOLID	TEICOPLANIN	VANCOMYCIN
		Staphylococcus aureus	50	66	100	34	34	8	78		40	84	10	10	100	96	10	100	100	100
		Staphylococcus epidermidis	14	93	100	0	0	0	93		29	71	36	36	100	71	10	100	100	100
Cura	Wound	Enterococcus faecalis	7		100			71		100	0		43	43	100	29	57	100	100	100
Surg		Staphylococcus haemolyticus	6	33	100	0	0	0	33		0	17	0	0	100	100	83	83	83	83
		Enterococcus faecium	5		0			0		100	0		0	0	100	0		100	75	75
	Tissue	Staphylococcus aureus	10	80	100	50	50	10	90		30	90	10	20	100	90	90	100	100	90

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	АМРНО В	CASPOFUNGIN	MICAFUNGIN
	Blood	Candida albicans	2	50	100	0	100	100
	Бюба	Candida guilliermondii	1	0	0	100	100	100
Cura	Wound	Candida albicans	2	100	100	50	100	100
Surg	Urine	Candida tropicalis	2	100	100	100	100	100
	Fluid	Candida parapsilosis	1	0	100	100	100	100
	riulu	Candida tropicalis	1	100	100	100	100	100

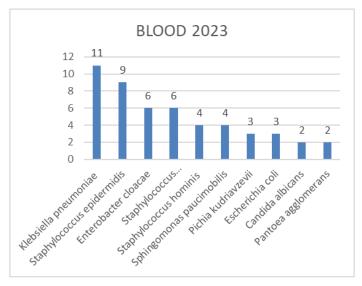
Please note: Individual isolates are less than 30 in number. Not statistically significant. Antibiogram has been shown with available number of organisms.

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Ar	Dr sk ulwani wam Medical Director

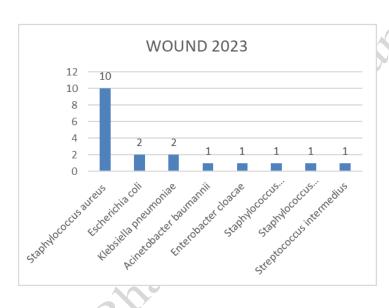


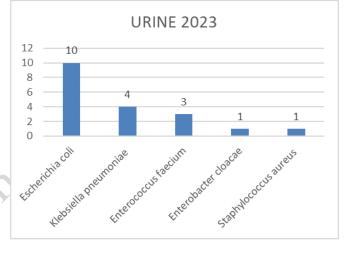
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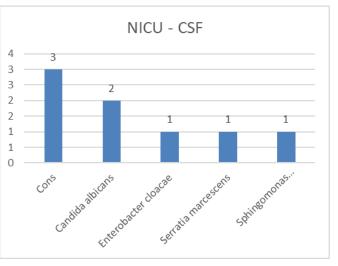
NEONATAL INTENSIVE CARE UNIT: TOP 10 ISOLATES SAMPLEWISE



SPECIMEN	OVERALL NO.
TYPE	ISOLATES
BLOOD	68
PUS	19
URINE	20
ENDOTRACHEAL	4
ASPIRATE	
	A () '







Prepared by	Checked by	Verified by	Apprøved by
Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by A.d A.	Dr sk tellwahi wam Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2024-2025

Version - 10.0

NICU ANTIBIOGRAM GRAM NEGATIVE

Percentage Susceptibility Gram Negative Organism 2023

Location	Specimen type	Organism	Number of patients	COTRIAMOXAZOLE	NITROFURANTOIN	OFLOXACIN	NORFLOXACIN	LEVOFLOXACIN	CIPROFLOXACIN	FOSFOMYCIN	AMOX/CLAV	CEFEPIME	CEFUROXIME	CEF/SUL	CEFTRIAXONE	PIPTAZ	AMIKACIN	GENTAMICIN	MEROPENEM	ERTAPENEM	IMIPENEM	MINOCYCLINE	COLISTIN
		Klebsiella pneumoniae	11	100					80	100	82	80	70	0	80	80	100	100	80	90	90		100
	Blood	Enterobacter cloacae	6	83					67	100	33	67	33	0	50	67	100	67	100	100	100		100
NICU		Sphingomonas paucimobilis	4	100				100	67			100		0	100	100	33	67	100		100	100	
	Urine	Escherichia coli	10	90	100	67	67		30	100	60	57	14	0	30	90	90	80	100	100	100		100
		Klebsiella pneumoniae	4	50					0	100	25	50	0	0	25	75	75	75	75	75	75		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.

Dr M Kumar
Prof Microbiology

Checked by

Verified by

Approved by

Dr AK Verma Quality
Department by Ned Air

Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2024-2025

Version - 10.0

NICU ANTIBIOGRAM GRAM POSITIVE

Percentage Susceptibility Gram Positive Organism 2023

	Location	Specimen type	Organism	Number of patients	OXACILLIN	CEFOXITIN	COTRIAMOXAZOLE	NITROFURANTOIN	PENICILLIN	CIPROFLOXACIN	LEVOFLOXACIN	ERYTHROMYCIN	CLINDAMYCIN	GENTAMICIN	TETRACYCLINE	TIGECYCLINE	DAPTOMYCIN	LINEZOLID	TEICOPLANIN	VANCOMYCIN	/
ĺ			Staphylococcus epidermidis	9	11	11	22	100	0	11	11	11	22	22	89		100	100	100	100	
	NICH	Blood	Staphylococcus haemolyticus	6	0	0	50	100	0	0	0	0	33	0	100		100	100	100	83	
	NICU		Staphylococcus hominis	4	25	25	50	100	0	50	50	0	25	100	75		100	100	100	100	
		Wound	Staphylococcus aureus	10	20	20	90	100	10	10	10	30	100	90	100	100	100	100	100	100	

Reserved / Restricted Drugs :Not to be used empirically 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

Candida isolated in NICU

Location	Specimen type	Organism	Number of patients	Fluconazole	Voriconazole	Ampho B	Caspofungin	Micafungin
		Candida albicans	2	100	50	0	100	100
		Candida parapsilosis	1	0			100	100
NICU	Blood	Candida tropicalis	1	100	100	100	100	100
	Cerebrospinal fluid	Candida albicans	2	50	100	50	100	100
	Fluid	Candida parapsilosis	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.

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Air	Dr SKT (wahi LVAY) Medical Director



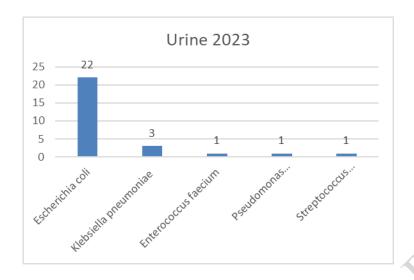
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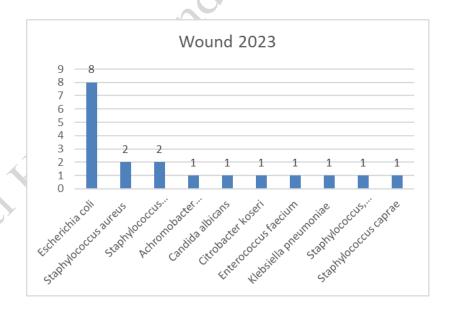
2024-2025

Version - 10.0

OBSTETRICS AND GYNAECOLOGY: TOP 10 ISOLATES SAMPLEWISE

URINE	28
WOUND	20
TYPE	ISOLATES
SPECIMEN	OVERALL NO OF





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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Air	Dr sktolwani wam Medical Director



2024-2025

Version - 10.0

OBSTETRICS AND GYNECOLOGY ANTIBIOGRAM GRAM NEGATIVE

Percentage Susceptibility Gram Negative Organism 2023

Location	Specimen type	Organism	Number of patients	COTRIAMOXAZOLE	NITROFURANTOIN	CIPROFLOXACIN	NORFLOXACIN	OFLOXACIN	LEVOFLOXACIN	FOSFOMYCIN	AMOX/CLAV	CEFUROXIME	CEF/SUL	CEFTAZIDIME	CEFTRIAXONE	CEFEPIME	PIPTAZ	AMIKACIN	GENTAMICIN	MEROPENEM	ERTAPENEM	IMIPENEM	TIGECYCLINE	COLISTIN
	Blood	Escherichia coli	2	100		0				100	100	0	0		0	0	100	100	100	100	100	100		100
		Escherichia coli	22	59	100	27	56	44		96	73	8	0	56	18	46	86	91	86	92	91	92		100
	Urine	Klebsiella pneumoniae	3	100	0	100	100	100		100	67	100	0	100	67	100	100	100	100	100	100	100		100
		Pseudomonas aeruginosa	1			0			0				0	0		0	0	0	0	0		0		0
Obgyn		Escherichia coli	8	50		0					25	0	0		0	13	63	75	63	88	88	88		100
Obgyii	Wound	Klebsiella pneumoniae	1	100		0					100	0	0		0	0	100	100	0	100	100	100	100	100
	wound	Achromobacter xylosoxidans	1	100		0			0				0	100		0	100	0	0	100		100		
		Citrobacter koseri	1	100		100					100	0	0		100	100	100	100	100	100	100	100		100
	Other	Escherichia coli	4	25		0					75	0	0		0	0	100	75	75	100	100	100		100
		Klebsiella pneumoniae	2	50		50					50	50	0		50	50	50	100	50	100	100	100	50	100

Reserved / Restricted Drugs :Not to be used empirically 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: Other than E. coli individual isolates are less than 30 in number. Not statistically significant numbers.

Dr M Kumar
Prof Microbiology

Checked by

Verified by

Dr AK Verma Quality
Department by Ned Dr

Medical Director



2024-2025

Version - 10.0

OBSTETRICS AND GYNECOLOGY ANTIBIOGRAM GRAM POSITIVE

No significant number of Gram Postive isolates in 2023

Percentage Susceptibility Gram Positive Organism 2023

Location	Specimen type	Organism	Number of patients	NITROFURANTOIN	COTRIAMOXAZOLE	OXACILLIN	CEFOXITIN	LEVOFLOXACIN	CIPROFLOXACIN	RIFAXIMIN	PENICILIIN	GENTAMICIN	GENTAMICIN HIGH LEVEL	ERYTHROMYCIN	CLINDAMYCIN	TETRACYCLINE	TEICOPLANIN	VANCOMYCIN	DAPTOMYCIN	LINEZOLID	TIGECYCLINE
	Blood	Staphylococcus hominis	1	100	0	100	100	0	0	100	100	100		0	0	100	100	100	100	100	
		Streptococcus agalactiae	1	100				100			100					0		100		100	
		Enterococcus faecium	1	0				0	0		0		100	0		0	0	0		0	100
		Staphylococcus, coagulase negative	1	100	100	100	100	100	100	100	0	100		100	100	100	100	100	100	100	100
Obgyn	Urine	Streptococcus agalactiae	1	100				100			100					0		100		100	
Obgyii		Enterococcus faecium	1	0				0	0		0		100	0		0	100	100		100	
		Staphylococcus haemolyticus	2	100	50	50	50	50	0	100	50	100		50	100	100	100	100	100	100	100
	Other	Enterococcus faecium	1	0				0	0		0		100	0		0	0	0		100	100
		Enterococcus faecalis	2	100				50	50		100		100	0		0	100	100	0	50	100
		Staphylococcus aureus	1	100	100	100	100	100	100	100	0	100		0	100	100	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 appropriate antibiotic

Please note: Individual isolate numbers are in single digits. Hence these isolates may not be truly representative of the antimicrobial susceptibility pattern.

Dr M Kumar
Prof Microbiology

Checked by

Verified by

Approved by

Dr AK Verma Quality
Department by Ned Ar

Date of release: 01.04.2025 Valid till Mar 2025 unless amended earlier

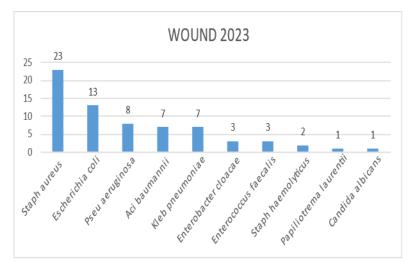
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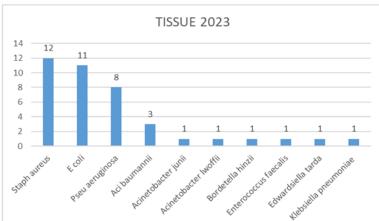
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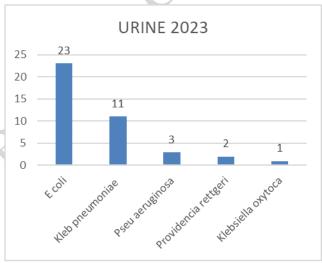
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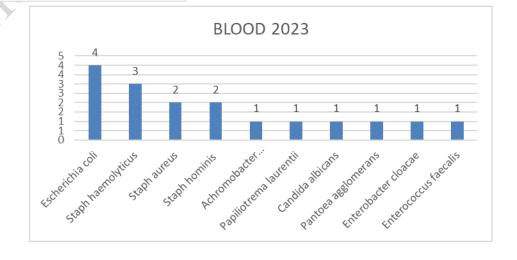
ORTHOPAEDICS: TOP 10 ISOLATES SAMPLEWISE



SPECIMEN	OVERALL NO OF
TYPE	ISOLATES
WOUND	77
TISSUE	46
URINE	42
BLOOD	22







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Dr M Kumar Prof Microbiology	Dr M S Modak Prof Microbiology	Dr AK Verma Quality Department by Ned Ar	Dr sk tellwani wam Medical Director



2024-2025

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ORTHOPAEDICS ANTIBIOGRAM GRAM NEGATIVE

Percentage Susceptibility Gram Negative Organism 2023

Location	Specimen type	Organism	Number of patients	COTRIAMOXAZOLE	CIPROFLOXACIN	LEVOFLOXACIN	FOSFOMYCIN	AMOX/CLAV	CEFUROXIME	CEF/SUL	CEFTAZIDIME	CEFTRIAZONE	CEFEPIME	PIPTAZ	AMIKACIN	GENTAMICIN	MEROPENEM	ERTAPENEM	IMIPENEM	TIGECYCLINE	COLISTIN
	Urine	Escherichia coli	23	22	4		96	13	0	0		4	14	38	48	44	39	39	46		100
	Offic	Klebsiella pneumoniae	11	46	27		82	36	27	0		18	36	46	46	55	46	46	46		100
		Escherichia coli	13	54	8		100	39	8	0		15	39	54	100	92	77	77	77		100
ortho	Wound	Pseudomonas aeruginosa	8		75	75				0	63		63	63	88	75	75		88		100
Ortho	vvounu	Acinetobacter baumannii	7	43	0					0		0	0	0	0	0	0		0		100
		Klebsiella pneumoniae	7	57	14		71	29	14	0		14	14	14	57	57	43	43	43	57	100
	Tissue	Escherichia coli	11	27	0		100	27	0	0		0	0	36	100	91	55	55	55		100
	rissue	Pseudomonas aeruginosa	8	,	75	63				0	75		75	63	88	88	63		75		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	Dr AK Verma Quality Department by Ned Air	Dr SKTOWani WAYN Medical Director



Antimicrobial Policy and Antimicrobial Stewardship Program

2024-2025

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ORTHOPAEDICS ANTIBIOGRAM GRAM POSITIVE

Percentage Susceptibility Gram Positive Organism 2023

Location	Specimen type	Organism	Number of patients	COTRIAMOXAZOLE	NITROFURANTOIN	OXACILLIN	CEFOXITIN	PENICILLIN	CIPROFLOXACIN	LEVOFLOXACIN	GENTAMICIN	GENTAMICIN HIGH LEVEL	CLINDAMYCIN	TETRACYCLINE	DAPTOMYCIN	LINEZOLID	TIGECYCLINE	TEICOPLANIN	VANCOMYCIN	
	Wound	Staphylococcus aureus	23	74	100	39	39	9	13	13	78	48	96	96	96	100	100	100	100	
ortho	Tissue	Staphylococcus aureus	12	58	100	50	50	17	17	25	75	42	100	92	100	100	100	100	100	
	Other	Staphylococcus aureus	5	60	100	60	60	0	20	20	40	60	100	100	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 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.

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Air	Dr sktolwani wam Medical Director

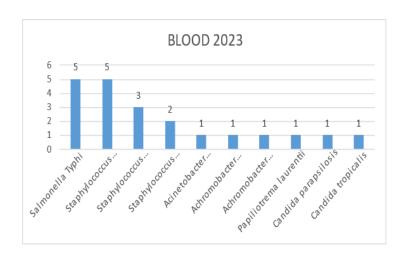




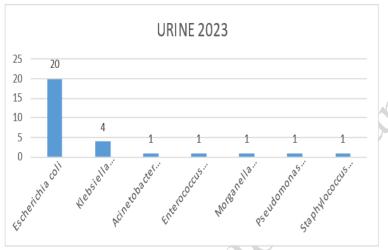
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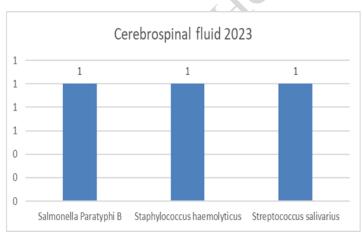
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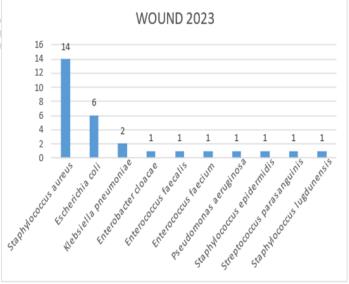
PEDIATRICS& PICU: TOP 10 ISOLATES SAMPLEWISE



SPECIMEN	OVERALL NO OF
TYPE	ISOLATES
BLOOD	36
WOUND	31
URINE	30







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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Air	Dr SKT (wani LVM) Medical Director



2024-2025

Version - 10.0

Paediatrics Antibiogram

Percentage Susceptibility Gram Negative Organism 2023

Location	Specimen type	Organism	Number of patients	COTRIAMOXAZOLE	CIPROFLOXACIN	LEVOFLOXACIN	FOSFOMYCIN	AMOX/CLAV	AZTREONAM	CEFEPIME	CEF/SUL	CEFTRIAXONE	CEFUROXIME	PIPTAZ	AMIKACIN	GENTAMICIN	IMIPENEM	MEROPENEM	ERTAPENEM	MINOCYCLINE	COLISTIN
	Blood	Salmonella Typhi	5	100	0	0		100		100		100		100			100	100	100		
	Wound	Escherichia coli	6	0	0	0		50	0	33	17	25	25	17	100	80	50	60	50	100	100
David		Klebsiella pneumoniae	2	50	0			50		50	100	0	0	100	100	100	50	100	100		100
Paed	Urine	Escherichia coli	20	37	5		100	20		47	47	32	16	47	90	63	74	68	68		100
		Klebsiella pneumoniae	4	25	0		50	0		0	0	0	0	0	25	0	0	25	25		100
	Respiratory	Pseudomonas aeruginosa	2		100	100				100	100			100	0	0	0	0			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.

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Air	Dr sktolwani wam Medical Director

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Please note: Individual isolates are less than 30 in number. Antibiogram has been shown with available number of organisms.

Percentage Susceptibility Gram Positive Organism 2023

																		- 4	
Location	Specimen type	Organism	Number of patients	CEFOXITIN	OXACILLIN	COTRIAMOXAZOLE	NITROFURANTOIN	CIPROFLOXACIN	LEVOFLOXACIN	PENICILLIN	GENTAMICIN	ERYTHROMYCIN	CLINDAMYCIN	TETRACYCLINE	TIGECYCLINE	LINEZOLID	DAPTOMYCIN	TEICOPLANIN	VANCOMYCIN
		Staphylococcus aureus	2	0	0	0	100	0	0	0	100	0	100	50		100	100	100	100
Paed	Blood	Staphylococcus hominis	3	33	33	100	100	100	100	33	67	33	67	100		67	67	67	67
Paeu		Staphylococcus epidermidis	5	0	0	80	100	60	60	0	60	20	40	100		100	100	60	100
	Wound	Staphylococcus aureus	14	50	50	50	100	0	0	0	93	43	86	100	100	100	93	100	93

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

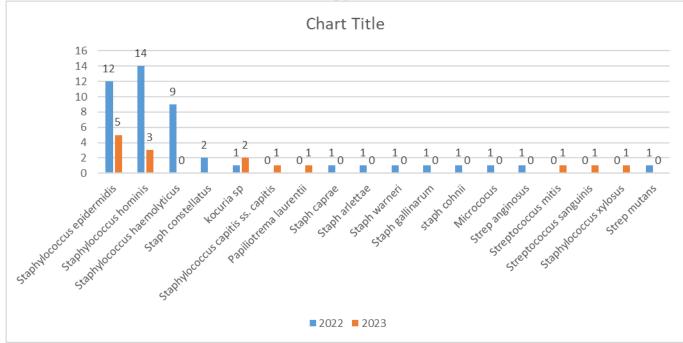
Will be useful clinically

Will be useful clinically in about 2/3 cases

Will not be useful clinically

Not tested/ not appropriate antibiotic

Skin flora/Collection contamination in blood culture – Paediatrics & PICU



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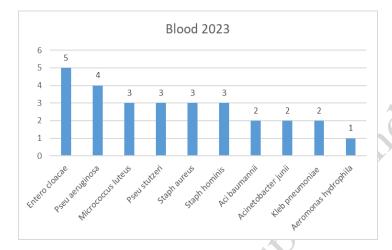
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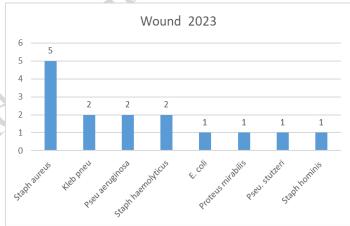
ONCOLOGY

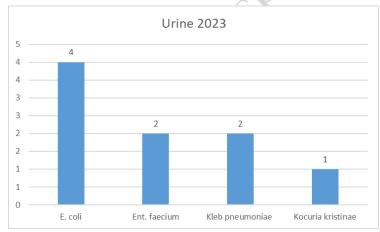
Please note: Individual isolates are less than 30 in number. Antibiogram has been shown with available number of organisms.

Please prescribe empiric therapy as per policy guidelines outlined in the section on therapy of common conditions and Febrile neutropenia

	I
SPECIMEN	NO OF
TYPE	ISOLATES
BLOOD	42
PUS/TISSUE	19
URINE	10







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ONCOLOGY

Percentage Susceptibility Gram Negative Organism 2023

Location	Specimen type	Organism	Number of patients	COTRIAMOXAZOLE	CIPROFLOXACIN	LEVOFLOXACIN	AMOX/CLAV	CEFEPIME	CEF/SUL	CEFTRIAXONE	CEFUROXIME	CEFTAZIDIME	PIPTAZ	AMIKACIN	GENTAMICIN	MEROPENEM	ERTAPENEM	IMIPENEM	TIGECYCLINE	MINOCYCLINE	FOSFOMYCIN	COLISTIN
		Enterobacter cloacae	5	40	40		0	40	0	20	20		40	40	40	40	40	40			20	40
		Pseudomonas aeruginosa	4		25	25		50	0			50	50	100	100	50		50				25
		Pseudomonas stutzeri	3	0	0	0		100	0			100	100	100	100	100		100				
		Acinetobacter baumannii	2	50	50			50	0	50			50	50	50	50		50				100
		Acinetobacter junii	2	50	100	100		100	0	0		0	0	0	100	0		0		100		0
	Blood	Klebsiella pneumoniae	2	0	0		0	0	0	0	0		0	50	50	50	50	50			100	0
		Escherichia coli	1	100	0		100	0	0	0	0		100	100	100	100	100	100			100	100
		Pseudomonas alcaligenes	1	0	0	0		100	0			100		100	0	100		100				
		Proteus mirabilis	1	0	0		0	0	0	0	0		100	0	0	0	0	0			100	
		Aeromonas hydrophila	1	100	100	100		100	0			100	0	100	100	0		0				
ONCO		Salmonella enterica	1	100	100		100	100	0	100	100		100	100	100	100	100	100			100	100
		Klebsiella pneumoniae	1	0	0		0	0	0	0	0		0	100	100	0	0	0	100		100	100
	Tissue	Citrobacter freundii	1	0	0		100	0	0	0	0		0	100	0	0	100	0			100	0
		Escherichia coli	1	0	0		0	0	0	0	0		0	0	0	0	0	0			100	100
		Klebsiella pneumoniae	2	50	0		0	50	0	0	0		50	50	50	50	50	0	50		100	100
		Pseudomonas aeruginosa	2		100	50		0	0			0	0	100	100	100		100				0
	Wound	Escherichia coli	1	0	0		0	0	0	0	0		0	100	100	100	0	100			100	100
		Proteus mirabilis	1	100	0		100	100	0	100	100		100	100	100	100	100	0			100	
		Pseudomonas stutzeri	1	0	0	100		100	0			100	100	100	0			0				
	l luiu a	Escherichia coli	4	0	0		0	25	0	0	0		50	100	75	100	100	100			100	100
	Urine	Klebsiella pneumoniae	2	0	0		50	50	0	0	0		50	100	100	100	100	100			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.

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ONCOLOGY GRAM NEGATIVE

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 2023

Location	Specimen type	Organism	Number of patients	COTRIAMOXAZOLE	NITROFURANTOIN	OXACILLIN	CEFOXITIN	PENICILLIN	CIPROFLOXACIN	LEVOFLOXACIN	GENTAMICIN	GENTAMICIN HIGH LEVEL	ERYTHROMYCIN	CLINDAMYCIN	TETRACYCLINE	DAPTOMYCIN	LINEZOLID	TIGECYCLINE	VANCOMYCIN	TEICOPLANIN
		Staphylococcus aureus	3	100		33	33	0	0	0	100		33	100	100	100	100		100	100
	Blood	Staphylococcus epidermidis	1	0		0	0	0	0	0	0		0	0	100	100	100		100	100
		Staphylococcus hominis	3	50		33	0	0	0	0	100		0	50	0	67	50		100	100
onco		Staphylococcus aureus	5	100		20	20	0	20	20	100		40	100	100	100	100	100	100	100
	Wound	Staphylococcus haemolyticus	2	50		0	0	0	0	0	0		0	50	100	100	50	100	100	100
		Staphylococcus hominis	1			100										100		100		
	Urine	Enterococcus faecium	2		0			0	0	0		100	0		0		100		50	50

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

Dr M Kumar
Prof Microbiology

Checked by

Verified by

Dr AK Verma Quality
Dr AK Verma Quality
Department by Ned Ar



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Skin flora/Collection contamination in blood culture - Oncology

ONCOLOGY GRAM POSITIVE

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 2023

										\mathcal{I}										
Location	Specimen type	Organism	Number of patients	COTRIAMOXAZOLE	NITROFURANTOIN	OXACILLIN	CEOXITIN	PENICILLIN	CIPROFLOXACIN	LEVOFLOXACIN	GENTAMICIN	GENTAMICIN HIGH LEVEL	ERYTHROMYCIN	CLINDAMYCIN	TETRACYCLINE	DAPTOMYCIN	LINEZOLID	TIGECYCLINE	VANCOMPCIN	TEICOPLANIN
		Staphylococcus aureus	3	100		33	33	0	0	0	100		33	100	100	100	100		100	100
	Blood	Staphylococcus epidermidis	1	0		0	0	0	0	0	0		0	0	100	100	100		100	100
	Staphylococcus hominis	3	50		33	0	0	0	0	100		0	50	0	67	50		100	100	
onco		Staphylococcus aureus	5	100		20	20	0	20	20	100		40	100	100	100	100	100	100	100
	Wound	Staphylococcus haemolyticus	2	50		0	0	0	0	0	0		0	50	100	100	50	100	100	100
		Staphylococcus hominis	1			100										100		100		
	Urine	Enterococcus faecium	2		0			0	0	0		100	0		0		100		50	50
			Staph	`	Rese	rve d	/ Res	چ ^{يزون} tricte	ed Dr	ugs:	- _S zapřn ^{AV} Notto	be ι	چن s ed	em p e	ericall	yunle	ss ju	stifie	· d	
					Will	be us	eful	clini			2 -	2023								
							eful	clini	cally	in ak	oout 2	2/3 ca	ses							
							Will not be useful clinically													
		Not t	este	d/no	tapp	ropri	ate a	an ti bi	otic											

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Dr M Kumar	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Air	Dr sk delwani wam
Prof Microbiology	Microbiology		Medical Director

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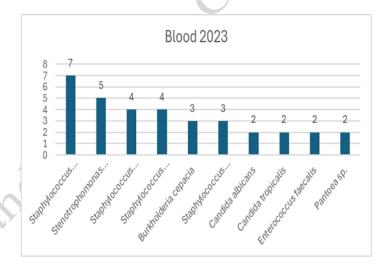
ALL OPD

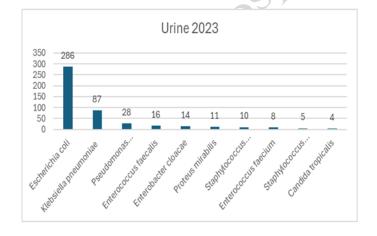
SPECIMEN	NUMBER OF
TYPE	ISOLATES
BLOOD	59
WOUND	362
SPUTUM	12
URINE	515

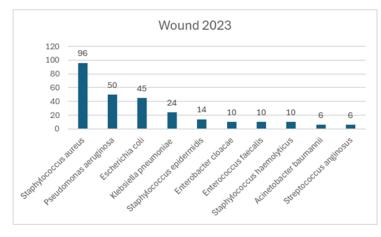
Blood isolates are from cultures taken in the Emergency Department

OPD: TOP 10 ISOLATES SAMPLEWISE

Skin flora/Collection contamination in EMD							
Staphylococcus epidermidis	7						
Staphylococcus haemolyticus	4						
Staphylococcus hominis ss. hominis	3						
Kocuria kristinae	1						
Kocuria rosea	1						
Micrococcus luteus	1						
Staphylococcus arlettae	1						
Staphylococcus saprophyticus ss. saprophyticus	1						
Staphylococcus caprae	1						
Streptococcus intermedius	1						
Staphylococcus xylosus	1						
Total	22						







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ALL OPD GRAM NEGATIVE

Percentage susceptibility gram negative organism 2023

Location	Specimen type	Organism	Number of patients	COTRIAMOXAZOLE	NITROFURANTOIN	OFLOXACIN	LEVOFLOXACIN	CIPROFLOXACIN	NORFLOXACIN	FOSFOMYCIN	AMOX/CLAV	CEFEPIME	Cefuroxime oral	CEFUROXIME	CEFTAZIDIME	CEFTRIAXONE	CEF/SUL	PIPTAZ	AMIKACIN	GENTAMICIN	MEROPENEM	ERTAPENEM	IMIPENEM	TIGECYCLINE	FOSFOMYCIN	COLISTIN
		Escherichia coli	286	51	86	36		13	45	99	55	37	15	27	64	28	78	78	81	73	92	92	92		99	100
		Klebsiella pneumoniae	87	48	33	74	0	29	74	76	38	30	18	41	63	39	45	45	65	64	55	60	56		76	100
	Urine	Pseudomonas aeruginosa	28				31	32				52			48		46	46	52	60	46		48			100
		Enterobacter cloacae	14	79	33	83		57	100	50	7	88	13	50	67	71	71	71	86	93	88	93	88		50	100
		Proteus mirabilis	11	46		0		27	33	100	82	88	88	100	100	91	100	100	82	64	100	100	38		100	
OPD		Pseudomonas aeruginosa	50				66	69				88			86		79	79	82	71	74		82			100
		Escherichia coli	45	71				4			44	49	13			29	78	78	91	76	93	91	96		100	100
	Wound	Klebsiella pneumoniae	24	58	0	100		42	100		61	55	41	100	100	54	65	65	78	83	86	71	77	65	87	100
		Enterobacter cloacae	10	90				80			10	90	10			60	90	90	90	100	90	90	90	90	80	100
		Proteus mirabilis	5	20			0	0			0	40	50		0	50	80	80	40	40	80	75	20		100	
	Other	Escherichia coli	10	40				30			70	40	30			30	80	80	100	90	100	90	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: Some 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.

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Ar	Dr SKT (wani LVA)





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ALL OPD GRAM POSITIVE

Percentage susceptibility gram positive organism 2023

Location	Specimen type	Organism	Number of patients	CEFOXITIN	OXACILLIN	COTRIAMOXAZOLE	NITROFURANTOIN	RIFAMPICIN	PENICILLIN	CIPROFLOXACIN	LEVOFLOXACIN	GENTAMICIN	GENTAMICIN HIGH LEVEL	ERYTHROMYCIN	CLINDAMYCIN	TETRACYCLINE	VANCOMYCIN	TEICOPLANIN	DAPTOMYCIN	LINEZOLID	TIGECYCLINE
		Staphylococcus aureus	96	35	35	61		97	5	11	12	78		48	87	93	100	100	95	99	100
	Wound	Staphylococcus epidermidis	14	29	29	71		79	0	36	43	86		14	57	93	100	100	93	93	100
		Enterococcus faecalis	10						100	44	44		100	11		0	100	100	67	100	100
		Enterococcus faecalis	16				94		81	6	6		100	19		0	94	94	69	100	
	Urine	Staphylococcus aureus	10	50	50	60	100	100	10	0	0	80		40	100	100	100	100	100	100	
	Offile	Enterococcus faecium	8				0		0	0	0		100	14		29	57	57		71	
		Staphylococcus saprophyticus	5	60	60	80	100	100	0	100	100	100		0	20	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 appropriate antibiotic

Please note: Some 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.

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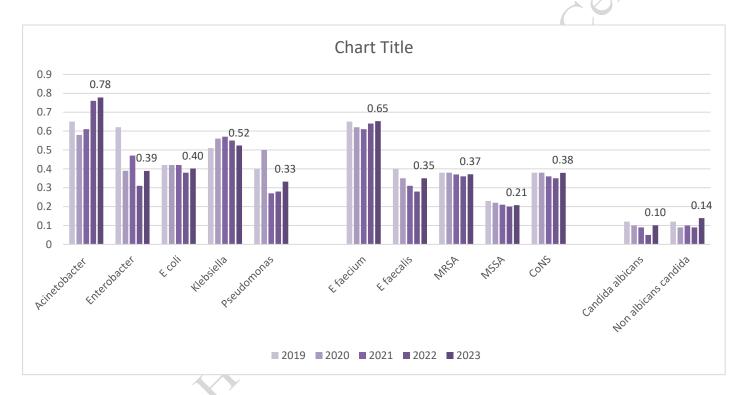
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Index of multidrug resistance Shows the trend of multidrug resistance of common pathogens for the years 2019-2023

Multi drug resistance index is calculated by formula Number of drugs resistant
Number of drugs tested

Value closer to 0.2 is considered the best







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6. 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	Current hospitalization>7 days.
No contact with health	last90 days. Current	Invasive procedures within last 90
care system	hospitalization less than 7 days	days
		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
	morbidities.	other Severe immunodeficiency

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by N. d. D. ?	Dr sk Talwahi Lum Medical Director



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

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Air	Dr sk tallwani wam Medical Director

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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	Inings on (0.5) 1 cm of
	(CAP)Ceftriaxone (2g	than 7 days)	Imipenem (0.5-1 gm q6h
	IV q24h X 5-7 days)/	TC 1 1	/Meropenem (1-2 g IV
	Amoxycillin/Clavulanic	If septic shock or	q8h)
	acid (1.2g q8h IV)	multisystem organ	Suspected XDR Gram
		failure, Imipenem0.5-	negative
	+	1gm q6h or	negative
	Macrolide	Meropenem1-2 gm	Colistin 4.5 MU/BD
	(Azithromycin- 500mg	q8h	
			Suspected MRSA-
	IV/PO once a day), x 5-7		Vancomycin (1g IV q12h
	days).		OR Teicoplanin (400mg
		2	IV q12h for 3 doses, then
			q24h)
	2] ICU patients with		E 4 LVDE
	CAP		For suspected VRE-
			Linezolid (600mg IV/PO
	Ceftriaxone (2g IV q24hr		q12hr)**x 7-14 days For suspected Fungal
	X 5-7 days)/		(Filamentous
			fungi/mould)
	Amoxycillin/Clavulanic		infections-
	acid (1,2g q8hr IV)		
	+		Consider Antifungals in
			Immunocompromised hos
Shara	Macrolide: Azithromycin-		Add Liposomal
	500mg IV/PO q24h)/		Amphotericin B.
	Doxycycline 100mg PO		Substitute Voriconazole,
	q12h x 5-7 days).		if Aspergillus suspected
7	TC		on radiological evidence
	If aspiration is suspected		or galactomannan positive
	clindamycin 600mg q8h		
ĺ			
	Early onset HAP/VAP		If PCP suspected- add TMP-SMX or

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	(less than 48 hours 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		3n.
	Look for typical viral symptoms such as sneezing and running nose.	2 esem	
H1N1	If fever, sore throat ,dry cough and viral symptoms present, initiate Oseltamivir 75 mg BD x 5 d without waiting for		
Flu-like illness	confirmation by PCR		

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.
- 6. In late HAP/VAP with suspected Acinetobacter infection combination of Colistin + carbepenem / sulbactam.

			/ .
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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Air	Dr SK Wwani WAY

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- 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.
- 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 therapy for MDR gram negative infection with specialized nebulisers.

Prepared by

Checked by

Verified by

Dr M S Modak Prof

Prof Microbiology

Dr AK Verma Quality

Dr AK Verma Quality

Department by Ned Ar



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Table 3: Meningitis

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

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof Microbiology	Dr AK Verma Quality Department by Ned Air	Dr SK Tulwani LVAM Medical Director



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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	,0	q8h/IV Imipenem-
	evidence of obstructive		cilastatin 500mg q6h
	uropathy)		+/-
	PO TMP SMX 160/800		IV Colistin 4.5 MUBD
	q12h / PO Nitrofurantoin	2	
	100 mg q12h	A Y	
			If MRSA or
	Acute Uncomplicated	Complicated	enterococcus,
	Pyelonephritis	Pyelonephritis	ConsiderVancomycin 1
	Fluroquinolones Ofloxacin	(Obstruction, reflux,	gm q12h/ Teicoplanin
	400 mg q12h OR	azotemia, CAUTI, Shock,	400 mg q24h
	Gentamicin 3 – 5 mg/kg	perinephric abscess)	
	q24h 5-7 days	Meropenem 1gm	
	If hospitalized	q8h/Imipenem-cilastatin	look for obstruction.
252	Ceftriaxone 1gm q12h	0.5 gm q6h in presence of obstruction	Surgical management is mandatory to relieve obstruction
1			

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Dr M Kumar	Dr M S Modak Prof	Dr AK Venna Quality	Dr skylivanilvam
Prof Microbiology	Microbiology	Department by Ned Dir	Medical Director

O INVERSITY

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em1gm q8h
in 4.5MUBD
X
7,7
em 1gm q8h

Table 4 a: Pelvic Inflammatory Disease

	Tummutory Biscuse		
Pelvic inflammatory	Tab Cefixime	Levofloxacin Plus	
disease: Mild to	Plus Tab Metronidazole	Metronidazole or	
moderate	Plus Cap Doxycycline	Ceftriaxone	
		differ	
Pelvic inflammatory	Clindamycin Plus	IV PIP-TZ 4.5 gm q6h or	
disease: Severe	Gentamicin	Imipenem	

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Dr M Kumar	Dr M S Modak Prof	Dr AK Venna Quality	Dr skilwani livam
Prof Microbiology	Microbiology	Department by Ned Dir	Médical Director

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Table 5- Skin & soft tissue infections

Name of condition	Patient Type 1	Patient Type 2	Patient Type 3
	(Community acquired)	(Healthcare	(Nosocomial Infections)
		associated)	
Erysipelas /	IV Ceftriaxone 2 gm q24h		
uncomplicated			
cellulitis	If beta lactam allergy		×
	IV Clindamycin 600 –		
	900 mg q8h		
Necrotizing	IV Ceftriaxone 2gm q12h	,	
infection of	+		
skin/fascia and	IV Clindamycin 600-		
muscle	900mg q8h / IV	70°	
	Metronidazole 500mg	a eseal.	
	q6h		
	If Suspected MRSA		
	IV Vancomycin1 gm		
	q12h/ IV Teicoplanin	2	
	400 mg q24h	O '	
	Mixed aerobic and		
	anaerobic cover		
	including S.aureus		
	MRSA cover IV		
	Vancomycin1gm q12h If pseudomonas		
	suspected IV PIP-TZ		
Fournier gangrene	4.5gm q6h		
- Januar Sangrene			IV Meropenem 1gm q8h or
•	IV Co-amoxiclav 1.2 gm		IV Imipenem-Cilastatin
X	q8h	If Suspected MRSA	1gm q6h. If MRSA
	if beta lactam allergy- IV		suspected IV Vancomycin
Diabetic foot		Vancomycin1 gmq12h	1gm q12h

Prepared by	Checked by	Verified by	Approved by
Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Air	Dr sk tallwani wam Medical Director



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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		$\sim \mathcal{O}_{\lambda}$
	If MRSA		
	suspected-		
	Vancomycin1gm		
	IVq12h		, O ^y
		Usual Suspected	
		organism- Staph	7)0
		aureus/ MRSA	
		IV Vancomycin 1	~
		gm q12h/	
		Teicoplanin(400mg	
		IV q12h for 3 doses, then q24h)	
		doses, then q2411)	
		If Suspected MDR	
		Gram negative	
	. x ° C	organism	
Early		IV Meropenem	
implant		1gm q8h	
associated	OS Y	IV Imipenem +	
infection (< 3		Cilastatin 1gm q6h	
months)	-	IV/IV Colistin	-
			Usually low grade infection
			If Coagulase negative
A			staphylococcus suspected -
	Y		IV Vancomycin 1 gm q12h /
Late implant			Teicoplanin (400mg IV q12h for 3
associated			doses, then q24h)
infection			If Anaerobe (Propionibacterium
(after 3			acne) suspected
months)	-	-	IV Clindamycin 600-900 mg q8h.

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Dr M Kumar	Dr M S Modak Prof	Dr AK Verma Quality	Dr skyllyanilvam
Prof Microbiology	Microbiology	Department by Med Dir	Medical Director



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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
			Linezolid 600 mg IV q12h
			suspected, Add
			Fluconazole 400 mg IV
			q24h If non albicans
		2	Candida- IV Caspofungin
			70 mg stat and 50 mg
		, y	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
			Imipenem -
	5		cilastatin500mg q6h .If
	40 .		MRSA or Enterococcus
			suspected IV Vancomycin
	•		1 gm q12h /
).			Teicoplanin(400mg IV
			q12h for 3 doses, then
			q24h). If VRE suspected
			Linezolid 600 mg IV q12h
			If Fungal Infection
\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\			suspected, Add Fluconazole 400 mg IV
Y			q24h
			If non Albicans Candida IV
			Caspofungin 70 mg stat and
NT-4 N/5-4 * 1 - 4 1 - 1	recing based on Paletensidek in a	includical for AA o	
Covers Alpinaudobiden	reemanbaseet on Ipharemadek Traf	gnsyddeslower gnsyffbri d	Istracumator o radabaket am

Dr M S Modak Prof Dr AK Verma Quality
Prof Microbiology

Dr M S Modak Prof Dr AK Verma Quality
Department by Ned Art Medical Director



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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	
	q12h for 4-6 weeks	, 0'

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

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof Microbiology	Dr AK Verma Quality Department by Ned Ar	Dr SKTUWani Wam Medical Director



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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
DI 1' D.1.'	OPP	D ' - D 1 '

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	or Quinine (10mg/kg TDS for 7
	(1.25 mg/kg) as a single dose or	.0
	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	
	and 2.4 mg/kg q24h X 5 days +	7days
	Doxycycline 100mg q12h x 7 days	
	11 100 101 7.1	Alternative: Amoxicillin (500 mg)PO
Leptospirosis (Mild)	Doxycycline 100mg q12h x 7 days	TDS x7 days
		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
	D1: (100) DD 7 (15 1	/IM 6 hourly x7 days
Scrub Typhus	Doxycycline (100mg) BD x 7 to 15 days	Alternative
Scrub Typhus	Or Agithmomyoin (500mg) OD v 2days	
	Azithromycin (500mg) OD x 3days	Chloramphenicol (500mg)QID
Enteric Fever (OPD)	T. Cefixime 400 mg TDS for 14 days	x7-15 days Alternative
Enteric rever (OPD)	1. Cerixinie 400 mg 1DS for 14 days	
		T. Azithromycin (1gm)OD for 5 days
Enteric Fever(IPD)	Ceftriaxone (4gm/day)IV for 7-14 days	luays
	, , , , , , , , , , , , , , , , , , ,	

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof Microbiology	Dr AK Verma Quality Department by Ned Air	Dr SK Tulwani LVAM Medical Director

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Table 10 Pediatric Infections

Name of	Patient Type 1 (Community		D. I III 0
condition	acquired)	Patient Type 2	Patient 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 to 3 months	Ceftriaxone100mg/kg/d od or Cefotaxime 150mg/kg/d tds x 10-14 days and *Azithromycin 10mg/kg/day x	Piperacillin- tazobactam 300 mg/kg/d qid	divided 8 hrly) plusVancomycin (40-60 mg/ kg/ day divided 6-8 hrly IV Meropenem (60-120 mg /kg/day divided 8 hrly) plusVancomycin (40-60 mg/ kg/ day divided 6-8 hrly. Add Fluconazole 6-12 mg/kg/day or amphotericin B (if renal dysfunction)
AGE: 4	Lobar	Piperacillin-	
months to 5 years	macrolide(azithromycin,) to cover Pertussis in partially unimmunized with DPT	tazobactam 300 mg/kg/d Qid plusVancomycin (40-60 mg/ kg/ day divided 6-8 hrly Ceftriaxone100mg/kg/d od Or Piperacillintazobactam 300 mg/kg/d qid	Same as above
		Either type II/post neurosurgical	Either type II/III or post shunt
Meningitis		meningitis	infection
Age > 3	Cefotaxime 200 mg/kg/d qid/or Ceftriaxone100mg/kg/d	IV Meropenem (120	IV Meropenem (120 mg /kg/day divided 8 hrly)/ plusVancomycin60mg/kg/d qid with
Age > 3 months	±	plusVancomycin(60	or without rifampin 10 mg/kg (PO)
шошшь	vancomychi oonig/kg/u	prus v ancomycin(00	or without mampin to mg/kg (FO)

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof Microbiology	Dr AK Verma Quality Department by Ned Air	Dr SK Tulwani LVAM Medical Director

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Urinary Tract Infection 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 Uncomplicated Amoxy-clav 30-40 mg/kg/d bd OR Ceftriaxone100mg/kg/d od OR Ceftriaxone100mg/kg/d dtds Pyelonephritis Complicated: Ceftriaxone100mg/kg/d dod OR Cefotaxime150mg/kg/d dod OR Cefotaxime150mg/kg/d tds OR Piperacillin-tazobactam 300 mg/kg/d tds/qid +/- Amikacin 15- Piperacillin-tazobactam300 mg/kg/d tds/qid+/- Amikacin 15- Qumg/kg/d dds/qid Or Meropenem120mg/kg/ Same as for type II		= = = = = = = = = = = = = = = = = = = =	mg/ kg/ day divided 6 hrly +/- rifampin 10 mg/kg(PO) q12h	q12h x 7-10 days after shunt removal Consider additional Intraventricular therapy Vancomycin 10mg or Gentamicin 1-2 mg or Polymixin B 2mg or Colistin 10mg 1mg = 12,500
Infection 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 Uncomplicated Amoxy-clav 30-40 mg/kg/d bd OR Ceftriaxone100mg/kg/d od OR Ceftriaxone100mg/kg/d tds Pyelonephritis Complicated: Ceftriaxone100mg/kg/d od OR Cefotaxime150mg/kg/d tds OR Piperacillin- tazobactam 300 mg/kg/d tds/qid +/- Amikacin 15- Piperacillin- tazobactam300 mg/kg/d tds/qid+/- Amikacin 15- Meropenem120mg/kg/ Same as for type II Piperacillin- tazobactam300 mg/kg/d tds/qid Or Meropenem120mg/kg/ Same as for type II				0- 0-
of trimethoprim bd OR Amoxy-clav 30-40 mg/kg/d bd OR Cefixime 8-10 mg/kg/d od Uncomplicated Amoxy-clav 30-40 mg/kg/d bd OR Ceftriaxone100mg/kg/d od OR Cefotaxime150mg/kg/d tds Pyelonephritis Complicated: 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 tds/qid+/- Amikacin 15- Meropenem120mg/kg/ Same as for type II				×
Uncomplicated Amoxy-clav 30-40 mg/kg/d bd OR Ceftriaxone100mg/kg/d od OR Cefotaxime150mg/kg/d tds Complicated: 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 tds/qid+/- Amikacin 15- Meropenem120mg/kg/ Same as for type II	Cystitis	of trimethoprim bd OR Amoxy-clav 30-40 mg/kg/d bd OR Cefixime 8-10 mg/kg/d		CENT
OR Piperacillin- tazobactam 300 mg/kg/d tds/qid +/- Amikacin 15- Piperacillin- tazobactam 300 mg/kg/d tds/qid+/- Amikacin 15- Meropenem120mg/kg/ Same as for type II	Pyelonephritis	30-40 mg/kg/d bd OR Ceftriaxone100mg/kg/d od OR Cefotaxime150mg/kg/d tds Complicated: Ceftriaxone100mg/kg/d od OR	and Res	Same as for type II
		300 mg/kg/d tds/qid +/- Amikacin 15- Piperacillin- tazobactam 300 mg/kg/d tds/qid+/- Amikacin 15-	tazobactam300 mg/kg/d tds/qid Or Meropenem120mg/kg/	Same as for type II

Prepared by	Checked by	Verified by	Apprøved by
Dr M Kumar	Dr M S Modak Prof	Dr AK Venna Quality	Dr skillyanilvam
Prof Microbiology	Microbiology	Department by Med Dir	Medical Director

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	Cloxacillin		
HEENT	200mg/kg/d	Piperacillin-	IV Meropenem (120 mg /kg/day divided 8 hrly)/ plus
Infections	plus either Cefotaxime	tazobactam 300	Vancomycin
Orbital	150mg/kg/d tds or	mg/kg/d tds/qid plus	60mg/kg/d qid
cellulitis	Ceftriaxone	Vancomycin	oomg, kg, a qia
	100mg/kg/d od/bd x	60mg/kg/d qid	
	10-14 days	oomg/kg/a qia	
	Cloxacillin		
Bone and Joint	200mg/kg/d	Vancomycin	IV Meropenem (120 mg /kg/day divided 8 hrly)/ plus
Infections	plus either Cefotaxime	60mg/kg/d qid or	Vancomycin
	150mg/kg/d tds or	oomg kg a qia oi	60mg/kg/d qid or Clindamycin
	150mg/kg/d tds of	Clindamycin 20-40	20-40
Acute	Ceftriaxone	mg/kg/d tds/qid	mg/kg/d tds/qid
Osteomyelitis/s	100mg/kg/d od/bd x	mg/kg/u tus/qiu	mg/kg/u tus/qiu
eptic arthritis	10-14 days	Plus either	
		Trus cities	
		Cefotaxime	
		150mg/kg/d tds or	
		Ceftriaxone	
		100mg/kg/d od/bd	
Osteochondritis	Piperacillin-		
	tazobactam300		
	mg/kg/d tds/qid or		
	combination therapy	y y	
	with cloxacillin		
	200mg/kg/d plus		
	Ceftazidime		
	100mg/kg/d tds		
	7-10 days after surgery		
	, 10 days after surgery		'
	Cloxacillin		
Skin and soft	200mg/kg/d	Vancomycin	Piperacillin- tazobactam 300
200	_ = = = = = = = = = = = = = = = = = = =	, , ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	mg/kg/d tds/qid or IV
tissue	or Cefazolin 60-	60mg/kg/d qid	Meropenem
		<i>8 8 1-2</i>	(120 mg /kg/day divided 8 hrly
infections	100mg/kg/d or		plus
			Vancomycin 60mg/kg/d qid
	Clindamycin 20-40		
	mg/kg/d tds/qid x 7-10		
	days		

Prepared by	Checked by	Verified by	Apprøved by
Dr M Kumar	Dr M S Modak Prof	Dr AK Verma Quality	Dr sk mlyahi
Prof Microbiology	Microbiology	Department by Ned Dir	Medical Director

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Animal bite	Amoxicillin/clavulanate	Alternatives	
wounds	50mg/kg/d tdsi.v or p.o		
(dog / cat)		Piperacillin300mg/kg/d	NA
		qid 7-10 days	
		Penicillin allergy	
			.0
		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	A 0
		days (cat bites)	
			Meropenem 120mg/kg/d
Vascular		Piperacillin-	tds plus
v uscului			Vancomycin 60mg/kg/d
catheter		tazobactam300	qid
associated		mg/kg/d tds/qid +	qu
Infections		Vancomycin60mg/kg/d	
linections			
		qid	
			IV Meropenem 80-120
Severe	Cefotaxime 150 mg/kg/	IV Piperacillin –	mg/ kg/8 hrly
Sepsis/septic	day divided 6-8 hrly	Tazobactam	+
	. X.	7	IV Vancomycin 45-60
shock	OR	300-400 mg/kg/day	mg/kg/day
	Ceftriaxone 100 mg/kg/	divided 8 hrly +	divided 6-8 hrly
	day divided 12 hrly	IV Vancomycin 45-60	
	+/-	mg/kg/day divided 6-8	
	amikacin 15-20	hrly	
	mg/kg/d od	,	

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Dr M Kumar	Dr M S Modak Prof	Dr AK Verma Quality	Dr skylivani livam
Prof Microbiology	Microbiology	Department by Ned Dir	Medical Director

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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			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

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Meningitis	For early onset: E coli, GBS, enteric bacilli, listeria, streptococ cus, H influenza, Neisseria meningitid es. For late onset: Klebsiella, Acinetoba cter, E.coli, Enterococc us, Staphyloco ccus (CONS) Others :Serratia, Burkholde ria, Pseudomo nas, Proteus	1st line: Cefotaxime plus Gentamicin 2nd line: Meropenem	Meropenem	Meropenem		Gram Positive: 14-days Gram negative: 21 days# #Ventriculi tis/Brain abscess: 6- 8 weeks
UTI	Enterococc us, E coli, Enterobact er	41051	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	Amikacin	10days
Skin and soft tissue infection	Staphyloco ccus		1 st line:Cloxa cillin 2 nd line:Vanc omycin	Vancomycin	Cloxacillin	7days
Arthritis	Staphyloco ccus ,Klebsiella		1 st line Piperacillin- Tazobactam 2 nd line Meropenem	1 st line: Piperacillin- Tazobactam 2 nd line: Meropenem	Ceftriaxone plus Vancomycin	Culture Negative: 2weeks

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Add Dir	Dr SK Ulwahi Wan Medical Director

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		3 rd line Colistin	3 rd line Colistin		Culture positive: 3 weeks
Osteomyeli tis	Staphyloco ccus, Gram Negative Bacilli	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 Vancomycin	4 weeks
Catheter related Infection	Staphyloco ccus(CON S), S.aureus, Gram negative bacteria	1 st line: Vancomycin and Amikacin 2 nd line:Piperacil lin-Tazobact 3 rd line: Meropenem 4 th line Colistin	Constill		10days
Fungal infection	Candida albicans and Candida Non albicans	Amphoterici n B or Fluconazole (depending on Antifungal susceptibility report)			Depending on location
8	arati				

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Dr M Kumar	Dr M S Modak Prof	Dr AK Verma Quality	Dr skyllyanilvam
Prof Microbiology	Microbiology	Department by Med Dir	Medical Director

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Table: 12 Empiric therapy of Ophthalmic infections

Sr. No	Category	Suspected Organisms		Alternative
1	Bacterial	S aureus and albus	Topical Moxifloxacin 0.5% eyedrops 3-6 times per day	
	conjunctivitis	H Aegyptius		
		H Influenzae,	Tobramycin eye ointment at bed time	
		C diphtheriae	Penicillin eye drops 10,000 units/ml	
2	Bacterial	Pseudomonas,	Moxifloxacin eye drops 0.5% 1 hourly	Fortified
	IZ	S.aureus	Endical Talanamia and dama	Vancomycin
	Keratitis	Pneumococcus N gonorrheae	Fortified Tobramycin eye drops	eye drops
				Amikacin eye
				drops
3	Fungal	Aspergillus,	Natamycin eye drops 6 times a day	Amphotericin B
	Keratitis			eye drops
		Fusarium, Candida	Itraconazole eye drops /ointment at bed	
		albicans	time Tablet Fluconazole 150mg twice	drops
			a day& eye drops 4-6 times per day	Intracameral
4	Vinel Venetitie	II Cimmless	Nystatin eye ointment	Amphotericin B Tablet
4	Viral Keratitis	H Simplex	Acyclovir Tablet 800mg 5 times a day	Valacyclovir
		H Zoster	and ointment 5 times a day Gancyclovir ointment	1000mg 3 times a
		H Zoster	Gancyclovir omunent	day
5	Endophthalmit	Saureus	Intravitreal Vancomycin 1 mg /0.1 ml	Intravitreal
3	is	5 dureus	intravirear vancomyem i mg/0.1 mi	Vancomyci
		Sepidermidis	andAmikacin 400microgrames /.ml	n 1mg/0.1ml and
		Streptococcus		Ceftriaxone
		Pseudomonas		2.25mg/0.1ml
		H Influenzae	Intravitreal Amphotericin B	C
		Candida /fusarium		
6		Staphylococci	Intravenous	Intravenous
	cellulitis		_,	Ceftriaxone
			Piperacillin and Tazobactam 4.5g	
			twice a day	
	1		Intravenous Metronidazole 100ml 3	
		N / (· / A	times a day	
		Mucormycosis/Asper gillosis	Intravenous Amphotericin B	
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	

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Air	Dr sk thiwani wan



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Table 13: ENT Infection

Name of	Patient Type 1	Patient Type 2	Patient Type 3
condition	(Community acquired)	71	31
Acute infection	Inj Ampicillin 1 gm q6h	-	-
like acute	Amoxicillin		
membranous	+clavulanic acid 1.2		, C
tonsillitis, ASOM,	gm q8h		X
Acute epiglottitis			
without			
complication			
Acute infection	Addition	-	-
with	Of aminoglycoside for		
complications like	gram negative coverage		· C y
acute mastoiditis,	and metronidazole for		
Quinsy	anaerobic coverage	0	
			/
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		
Chronic infection	amikacin	ID/ Medicine	ID/ Medicine consult
with	+ inj metronidazole	consult	
complications like			
meningitis, orbital			
cellulitis, brain			
abscess	4077		

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by N. d. D. ?	Dr sktolwani wam Medical Director



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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	0.	
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
	(0)		colitis or sepsis
	\		suspected
•	Y		Oral Vancomycin
X	>		250 mg q6h
			+
			Metronidazole 500
100			mg q8h IV

Note:

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

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof Microbiology	Dr AK Verma Quality Department by Ned Ar	Dr SKTUWahi Wam Medical Director



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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 (short term)	-	+ Meropenem 2gm q8h IV Vancomycin 1 gm q12h IV	Meropenem 2gm q8h IV + Vancomycin 1 gm 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

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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

Dr M S Modak Prof Dr AK Verma Quality
Prof Microbiology

Dr M S Modak Prof Dr AK Verma Quality
Department by Ned Art Medical Director



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

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

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Arr	Dr sk tellwani wam Medical Director



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Table 17 : Pediatric surgery

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

Clean surgeries:

Procedure	Antimicrobial	Duration
Circumcision, Orchiopexy	Cefuroxime	Single dose
Herniotomy, LN biopsy,	No antimicrobials	-
Thyroglossal cyst Lipoma,		
small lumps	. x 0	
Bharath		

Dr M Kumar
Prof Microbiology

Checked by

Verified by

Dr AK Verma Quality
Dr AK Verma Quality
Department by Ned Ar

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Table 18: Antimicrobial prophylaxis for GI endoscopy

Antimic	crobial prophylaxis and/	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; continue 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	- "C),	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	Prevention of septic	-	Not recommended
Peritoneal dialysis	Lower GI endoscopy	Prevention of peritonitis	Ceftriaxone + metronidazole	Suggested
EUS	-FNA, EUS-guided FNA.			•
Bharai	Hosk			

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Air	Dr sk tallwani wam Medical Director

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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-Haemodynamically 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-Haemodynamically 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, haemodynamically 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, haemodynamically unstable but still febrile

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

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Air	Dr sk udlyani wan Medical Director

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Blood culture growing Gram negative bacilli

- Patient afebrile- continue the empirical antibiotic till antibiotic sensitivity is available.
- Rationalise as per susceptibility profiles

When to add glycopeptides?

- 1. Haemodynamic instability, or other evidence of severe sepsis, septic shock or pneumonia
- 2. 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. Heamodynamic instability.
- 2. Colonisation with carbapenem resistant gram-negative bacteria.
- 3. Previous infection with carbapenem resistant gram-negative bacteria.
- 4. GNB in blood, sensitivity pending, persistent fever with haemodynamic instability.

Empirical Antifungal Therapy

- No response to broad spectrum antibiotics (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 antibiotics 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 antibiotics, 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 antibiotic regimen.

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Venna Quality Department by Ned Dr?	Dr sk tellwani wam Medical Director



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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
- Antibiotic treatment should be given for at least seven days with an apparently effective antibiotic, with at least four days without fever.
- Once Neutrophil count has recovered, with no culture positivity and hemodynamically stable; antibiotics can be stopped and patient observed, even if remains febrile. Evaluate for fungal infection, if at risk.

Antiviral prophylaxis

- 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

Dr M Kumar
Prof Microbiology

Checked by

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Approved by

Dr AK Verma Quality
Department by Ned Dr

Medical Director



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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 antibiotics, 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

Antibiotic 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 Kumar

Prof Microbiology

Dr M S Modak Prof Lodd

Dr AK Verma Quality

Department by Nod Air

Medical Director

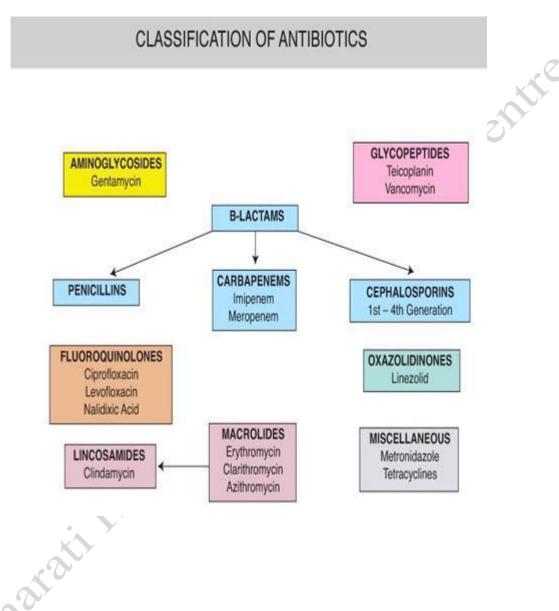


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Appendix 1

Commonly used antimicrobials



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Dr M Kumar	Dr M S Modak Prof	Dr AK Verma Quality	Dr sk Wani Lutam
Prof Microbiology	Microbiology	Department by Ned Dir	Medical Director



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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. Amoxycillin (PO)	Gram +ve Gram +ve Gram +ve & Gram –ve Gram +ve	Not easily available 1-2 gms q6h 500 mg q8h		
β – lactamase resistant	Cloxacillin	Gram +ve	0.5-1gm q6h		
β-lactam/ β- lactam inhibitor combination	Piperacillin- tazobactam. Ampicillin- sulbactam. Amoxycillin- 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		-00			
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		

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof Microbiology	Dr AK Verma Quality Department by Ned Ar	Dr SKTUWahi Wam Medical Director

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Antimicrobial				Side
Class	Name	Organisms	Indication& Dose	effects
Cephalosporin Plus	Cefoperazone	Anti-pseudomonas	q12h	
beta lactamase	/sulbactam	-	1.5 gm –	
inhibitor			3gm q12h	
minortor				
Aminoglycosides	Streptomycin	Gram –ve	0.75 -1gm q24h	Deafnes
	Kanamycin	Gram –ve	3mg/kg q24h	S
	Gentamicin	Gram –ve	13mg/kg q24h	Vertigo
	Amikacin	Gram –ve	3mg/kg q24h	Muscle
	Tobramycin	Gram –ve	5mg/kg q24h	weaknes
	Netilmicin	Gram -ve		S
Quinolones	Nalidixic acid		1 gm q6h	
-	Norfloxacin		400 mg q12h	
	Ciprofloxacin		500 mg q12h	Seizures
	Ofloxacin		200 mg q12h	
Extended	Levofloxacin		750 mg q24h	
spectrum	Moxifloxacin		400 mg q24h	
Carbapenems				Seizures
Imipenem-		Gram +ve except MRSA,	0.5gm -1gm q6h	
cilastatin		ESBL Gram –ve except		
Meropenem		Stenotrophomonas,	1-2 gm q8h	
Doripenem		Burkholderia,		
•		Corynebacterium,		
		Enterococcus faecium not		
	• X	covered		
Ertapenem		Does not cover		
_		Pseudomonas,		
		Acinetobacter &		
	Y	Enterococcus	1gm q24h	
				Muscle
Polymyxins	C ^y			weakness
Polymyxin B		ESBL, Metalloproteinase	Colistin	
			4.5MUBD	
			(loading dose of	
Colistin		producing Gram –ve	9MU is mandatory	Renal
				toxicity
Lincosamide				C.
Clindamycin		Gram +ve and anerobes	600mg q8h	difficile
				colitis

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof Microbiology	Dr AK Verma Quality Department by Ned Ar	Dr SKTUWahi Wam Medical Director

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Antimicrobial Class	Name	Organisms	Indication& Dose	Side effects
Class	Name	Organisms	Dose	cifects
Glycopeptides Vancomycin Teicoplanin		MRSA	1gm q12h 400 mg	Renal toxicity
1			q24h	A ()
Oxazolidinedione Linezolid		VRE	600 mg q12h	Thromb ocytope nia
Lipopeptides Daptomycin		MRSA	4-6mg/kg q24h	
Antifungals) '	
Fluconazole		Candida albicans	400 mg	
Voriconazole		Aspergillus	q12h 6mg/kg q12h first day then	
Caspofungin		Non albicans candida	4mg/kg 70mg IV then 50 mg q24h	
Anidulafungin		Non albicans candida	 	
AmphoB aqueous	•)	Broad spectrum covers all	Refer	
AmphoB colloidal AmphoB liposomal	(1050)	above + Mucor etc	product insert	

Prepared by	Checked by	Verified by	Apprøved by
Dr M Kumar	Dr M S Modak Prof	Dr AK Venna Quality	Dr skillyanilvam
Prof Microbiology	Microbiology	Department by Med Dir	Medical Director



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Appendix 2

Duration of therapy for various clinical conditions

Short course therapy is	equally effective		
Condition	Short Course (Days)	Long Course (Days)	Outcome
Chronic Bronchitis and obstructive pulmonary disease, acute exacerbations*	<=5	>=7	Equivalent Most patients do not require antimicrobials at all
Intra-abdominal infection	4	10	Equivalent
Neutropenic fever	Until afebrile and stable	Until non neutropenic	Equivalent
Osteomyelitis, Chronic	42	84	Equivalent
Pneumonia community acquired	<=8	10-15	Equivalent
Pyelonephritis	5-7	10-14	Equivalent
Skin Infections (Cellulitis, Major abscesses, wound infections)	5-6	10-14	Equivalent
Sinusitis, acute bacterial	5	10	Equivalent

Reference: Harrison's principle of Internal Medicine 20th ed

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof Microbiology	Dr AK Verma Quality Department by Ned Air	Dr SK Tulwani LVAM Medical Director

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Appendix 3 **Antimicrobial Agent Form**

0 6	E	HARAT	I VIDYAPE	ETH UNIV	ERSITY M	EDICAL CO	OLLEGE HO	SPITAL A	ND RESI	EARCH CEN	TRE
(0)				AN	TIMICROBI	AL AGEN	T FORM				
No. of Concession, Name of Street, or other Persons, Name of Street, or ot	-				BHR	C/HIC/F23(V-2))
	high end an										
1. Carl 9.Tige	bapenems 2 cycline/ Mir	2. Piptaz locycline	3. Levofloxac 10. Teicoplar	in ,4. Colistir nins 11. Line	n, 5. Polymyxir ezolids 12 .Ech	B , 6. Fosfo ninocandins	mycin/dapton 13. Voriconaz	nycin 7.Teico ole/ Posacor	oplanins nazole 14	8.Vancomyci I.Amphoterici	n, n B
	1 : ICU I/II/I No :	_	Ortho) N	Med 3/6/10	/11 Oth	er Wards:_			Гр.	aste Patient	\neg
Date	of Admissio				Date of	_				arcode here	1
	Confirmed		site of infect	tion:	Fat	ient type :	1 2	3	Wtof	patientk	kg.
	Date of Su	rgery:								2) Y	
[Time of in	cision	Time of	f 1 st dose	Time of	2 nd dose	Duration	of surgery	HRS		
	AMA Deta	ails:	1.	High End A	ntibiotics		2. Other Ant	ibiotics [
AMA(Generic name)	E/D/SP*		ite , Freq d Dose	Start Date	Day 1	Day3	Day7	IV to Oral	Stop Date	CHA AMA	NGE OF
		(mg	(/g)			CREATININE				(ren	narks)
						20					
		+			- x 2	-					
				1.0	>						
					Prophylaxis						
			al Line 2. Pe	-	e 3. HD Cath of removal_		ary Catheter	5. E11/11	6.Dra	ains	
					ES / NO; If Y		Report:				
(B/P/U/Resp/ and Date(DD/	CSF)* MM/YY)	Date	rganism isol: d(DD/MM/Y		Sensitivity P (Name of im			AMA Y/N		Response aft change Y/N	er
(1BEFORE,2	AFTER)	9									$\overline{}$
											\neg
	*=B- Bloo	d ; P- Pu	s ; U- Urine	; Resp – Spi	utum,ETT,BA	I ; CSF- Cer	ebro spinal f	luid			
Collected	Data by : _				Prescriber sig						
										/ .	
Prepared by	/		Checked b	ру		Verified b	DY LAL		Apprø	red by	
Dr M Kumai Prof Microb			Dr M S M Microbiol		Hodol.	Dr AK Ver Departm	ma Quality ent by Ned	Ma_	Dr SKT	lwani W Director	ran

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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 catheter:

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

swab Allow it to dry thoroughly.

70% alcohol

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.

Prepared by

Checked by

Verified by

Dr M S Modak Prof

Prof Microbiology

Dr AK Verma Quality

Dr AK Verma Quality

Department by Med Ard

Medical Director



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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 venipuncture.





- (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)

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Dr M Kumar Prof Microbiology	Dr M S Modak Prof	Dr AK Verma Quality Department by Ned Arr	Dr sk tulwani wam Medical Director



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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 disposable syringe, place in a sterile container and send to the laboratory.

(b) 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 sample as high as possible in the vaginal vault.

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Dr M Kumar Prof Microbiology		Dr AK Verma Quality Department by Ned Air	Dr skt (lwahi Lwam) Medical Director



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Dr M Kumar
Prof Microbiology

Checked by

Verified by

Dr AK Verma Quality
Dr AK Verma Quality
Department by Ned Ar