Interpreting Microbiology Results

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2025 Nebraska Antimicrobial Stewardship Summit LTC Fundamentals in Antibiotic Stewardship Track Friday, May 30, 2025



DEPT, OF HEALTH AND HUMAN SERVICES



NEBRASKA INFECTION CONTROL ASSESSMENT AND PROMOTION PROGRAM ANTIMICROBIAL STEWARDSHIP ASSESSMENT AND PROMOTION PROGRAM

Relevant Financial Disclosures

No Disclosures

None of the faculty or planners for this activity have relevant financial relationship(s) to disclose with ineligible companies whose primary business is producing, marketing, selling, re-selling, or distributing healthcare products used by or on patients



Objectives

- 1. Equip infection preventionists (IPs) with the knowledge to accurately interpret microbiology results, including distinguishing between colonization and active infection.
- 2. Provide IPs with practical examples of microbiology results and guide them on how to interpret these findings in the context of infection prevention.
- 3. Highlight the critical factors that influence the interpretation of microbiology results.



Key Aspects of Interpreting Microbiology Results



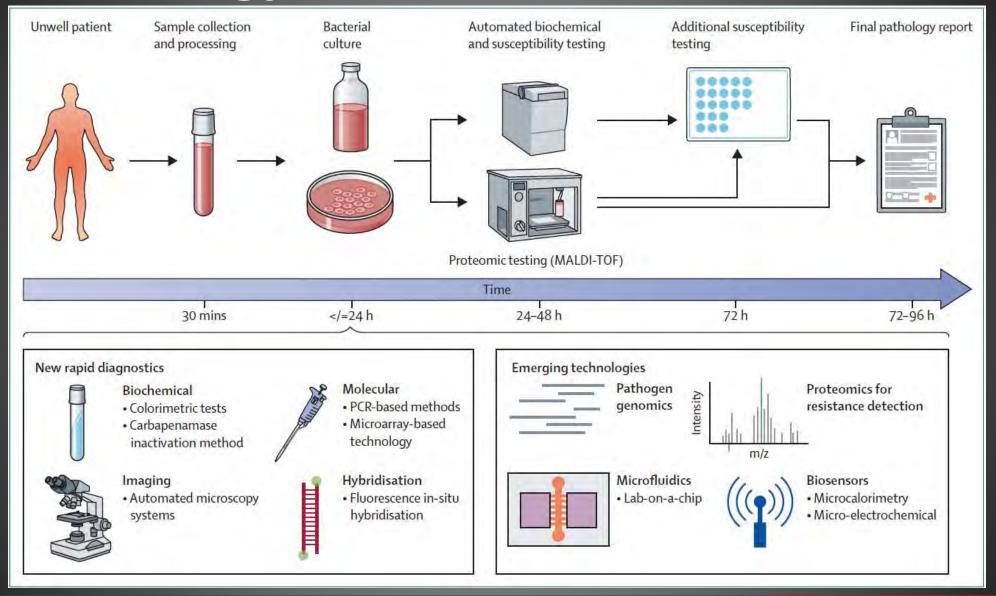
The Microbiology Lab Process³

Isolate Organism Identify Organism

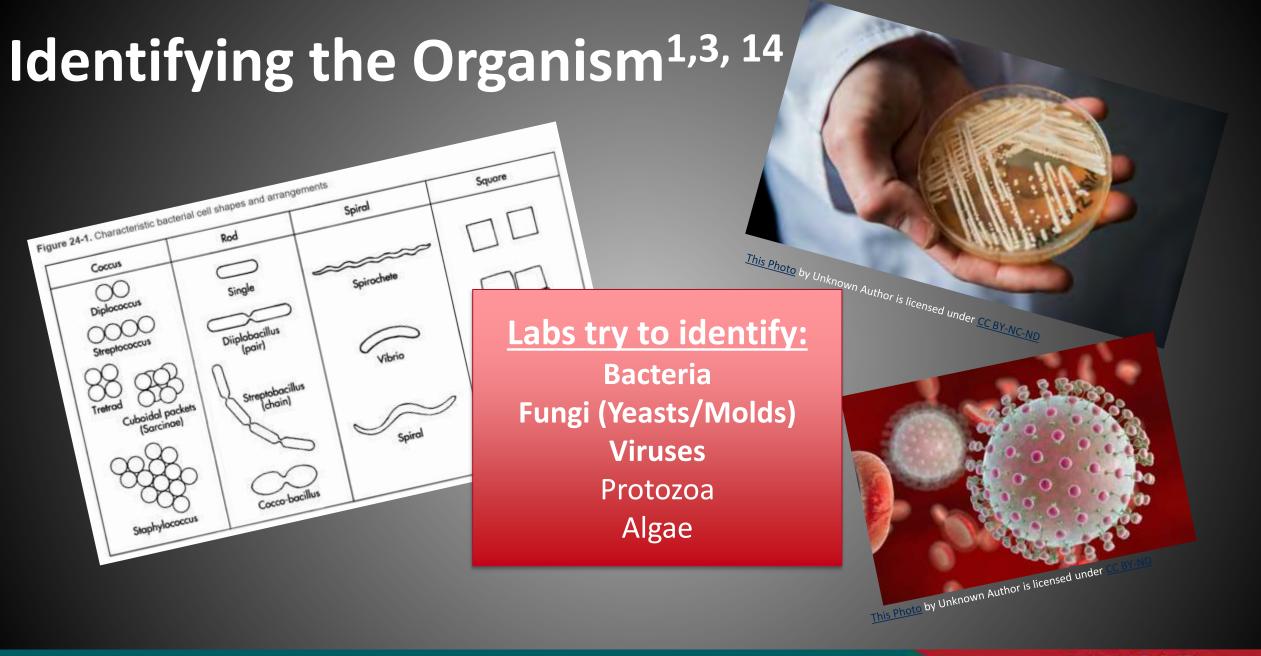
Susceptibility Testing



The Microbiology Lab Process⁶

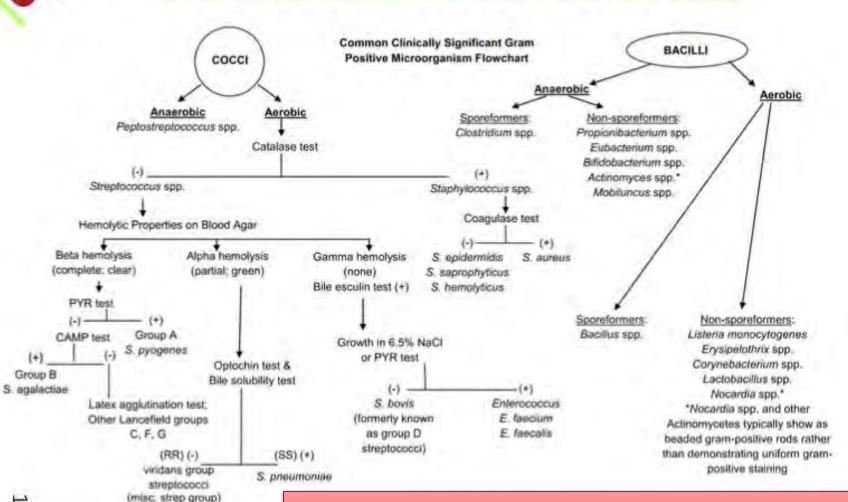








Organism Identification GRAM POSITIVE FLOWCHARTS



Identifying the Organism

Chart in the Workbook at page 17



Common Types of Cultures^{1,15}

- Blood
- Respiratory
- Urine
- Wound
- Stool





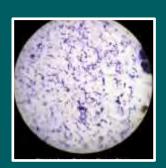


Common Methods Used for Identifying/Isolating the Organism^{1,14}



Microscope Slides

- Direct Examination
- Direct Wet Mount



Gram Stain



Acid-Fast Stain



Streak Plate



PCR



URINE CULTURE RESULTS

>100,000 col/ml Escherichia coli Note: Positive for extended spectrum beta-lactamase (ESBL)

Ampicillin	<=16	Resistant	
Cefazolin	>16	Resistant	
Cefepime	8	Resistant	
Ceftriaxone	>32	Resistant	
Cefuroxime	>16	Resistant	
Gentamicin	<=2	Susceptible	
Ciprofloxacin	>4	Resistant	
A STATE OF THE STA	<=1		
Meropenem	<=8	Susceptible	
Piperacillin/Tazobactam	<=0.5	Resistant	
Ertapenem	<=32	Susceptible	
Nitrofurantoin	<=0.5	Susceptible	
Trimethoprim/sulfamethoxazole	<=0.5/9.5	Susceptible	

Antibiotic Resistance¹

- Understanding antibiotic resistance patterns is crucial for selecting the most effective treatment
- Also impacts Infection
 Prevention and Control
 Measures to implement,
 such as transmission-based
 precautions.



Antibiotic Susceptibility Testing (AST)^{1,4}

Antibiotic	MIC	Interpretation
Aztreonam	8	S
Ceftriaxone	> 32	R
Cefepime	2	S
Ciprofloxacin	≤ 1	1

Next to each antibiotic is the susceptibility interpretation: S (sensitive), I (intermediate), or R (resistant), followed by the MIC in μg/mL

- Sensitive implies that the organism is inhibited by the serum concentration of the drug that is achieved using the usual dosage
- Intermediate implies that the organisms are inhibited only by the maximum recommended dosage
- Resistant implies that the organisms are resistant to the usually achievable serum drug levels.



Minimum Inhibitory Concentration (MIC)^{1,4}

Measure of drug activity = minimum inhibitory concentration (MIC)

 Breakpoints established by the U.S. Clinical and Laboratory Standards Institutes (CLSI)

Result	MIC	Clinical Correlation
Susceptible	≤ the defined susceptibility breakpoint	high likelihood of therapeutic success
Intermediate or Indeterminate	Intermediate value	therapeutic effect uncertain
Resistant	> the defined susceptibility breakpoint	high likelihood of therapeutic failure



What is CLSI?2,12

- Clinical and Laboratory Standards Institute
- Establishes the standards used by all labs to do susceptibility testing
- Annually updated
- Developed by physicians, microbiologists, pharmacists, and industry



Aminoglycoside Breakpoints	for Enterobacterales and A	Pseudomonas aeruginosa
----------------------------	----------------------------	------------------------

Table 1. Current CLSI Aminoglycoside MIC Breakpoints

	Antimicrobial	Interpretive Categories and MIC Breakpoints, µg/mL						
Organism Group	Agent	5	500		R			
Enterobacterales	Gentamicin	≤2	-	4^	≥8			
	Tobramycin	≤2	-	4^	≥8			
	Amikacin	≤ 4	-	8^	≥16			
P. aeruginosa	Gentamicin	4	4	2	- 4			
	Tobramycin	≤1		2^	≥4			
	Amikacin (U) ^b	≤ 16	3	32^	≥ 64			

Abbreviations: I, intermediate; MIC, minimal inhibitory concentration; R, resistant; S, susceptible; SDD, susceptible-dose dependent; U, urine. Symbol: ^, designation for agents that have the potential to concentrate in the urine.

Example from CLSI's rationale document MR 16 "Aminoglycoside Breakpoints from Enterobacterales and Pseudomonas Aeruginosa", published March 2025



^{*}Last reviewed June 2022; first published in CLSI M100-Ed33.

^{*} Report only on organisms isolated from the urinary tract.

AST Methods Done in the Lab¹

Manual Methods

- Disk diffusion
- MIC
- Gradient strips

Automated Methods

- Vitek 2
- Phoenix
- Microscan
- Sensititre



Resistance Genes^{9,13}

	Common Antibiotic Resistance Genes					
Gene	Indicate resistance to:	Organisms most affected:				
mecA mecA/C MREJ	methicillin and other applicable beta-lactam antibiotics	Staphylococcus species				
vanA/B	vancomycin	Enterococcus faecalis and Enterococcus faecium (and occasionally other organisms)				
CTM-M	·	Several types of Gram-negative bacteria can produce these enzymes				
mcr-1	colistin	Enterobacterales				
KPC						
IMP		Carbapenemase Gene found most often associated				
VIM	Carbapenems such as ertapenem,	with Carbapenem Resistant Enterobacterales (CRE), Pseudomonas Aeruginosa, and Acinetobacter				
NDM		species.				
OXA-48-like		,				



Active Infection vs. Colonization^{1,16}

	Active Infection	Colonization
Definition	Infection occurs when an organism invades a body site and causes signs and symptoms of disease	When an organism is found in the body, but it is NOT causing any symptoms or disease.
Treatment	Treatment is based on case-by-case basis	Treatment is often not required
Infectiousness	Can spread to others	Can develop infections and spread to others
Identification	Usually identified as part of standard clinical process.	Usually identified through screening
Transmission Based Precautions for LTCF	Contact Precautions	Enhanced Barrier Precautions



Multidrug-Resistant Organisms (MDRO) Tiers for Nebraska⁸

Tier	Definition of Included Organisms and Mechanisms	Examples (not all inclusive) of organisms/mechanisms for Nebraska	Transmission-Based Precautions Recommendations
Tier 1	Never (or very rarely) been identified in the United States and for which experience is extremely limited	Novel Carbapenemases	Contact precautions until otherwise recommended by HAI/AR team
Tier 2	Primarily associated with healthcare settings and are not commonly identified in the region (i.e., not been previously identified in the region or have been limited to sporadic cases or small outbreaks), corresponding to "not detected" or "limited to moderate spread" epidemiologic stages. No current treatment options exist (pan not-susceptible) and potential to spread more widely.	Pan-resistant organisms* Candida auris Carbapenemase (e.g., KPC, NDM, OXA-48, VIM, IMP) producing organisms (CPO) Enterobacterales Pseudomonas aeruginosa Acinetobacter Baumannii	Contact Precautions Long-term Care Facilities (LTCF): Enhanced barrier precautions (EBP) recommended for colonized resident(s)**
Tier 3	Include MDROs targeted by the facility or region for epidemiologic importance that have been identified frequently across a region, indicating advanced spread, but are not considered endemic	 Extended spectrum beta-lactamase (ESBL) producing organisms Carbapenem-resistant Enterobacterales (CRE) Carbapenem-Resistant Pseudomonas aeruginosa (CRPA) 	Contact Precautions Long-term Care Facilities (LTCF): Enhanced barrier precautions (EBP) considered for colonized resident(s)**
Tier 4	Endemic in a region and have been targeted by public health for their clinical significance and potential to spread rapidly	 Methicillin-resistant Staphylococcus aureus (MRSA) Vancomycin-Resistant Enterococci (VRE) 	Contact precautions per facility risk assessment Long-term Care Facilities (LTCF): Enhanced barrier precautions (EBP) considered for colonized resident(s)**

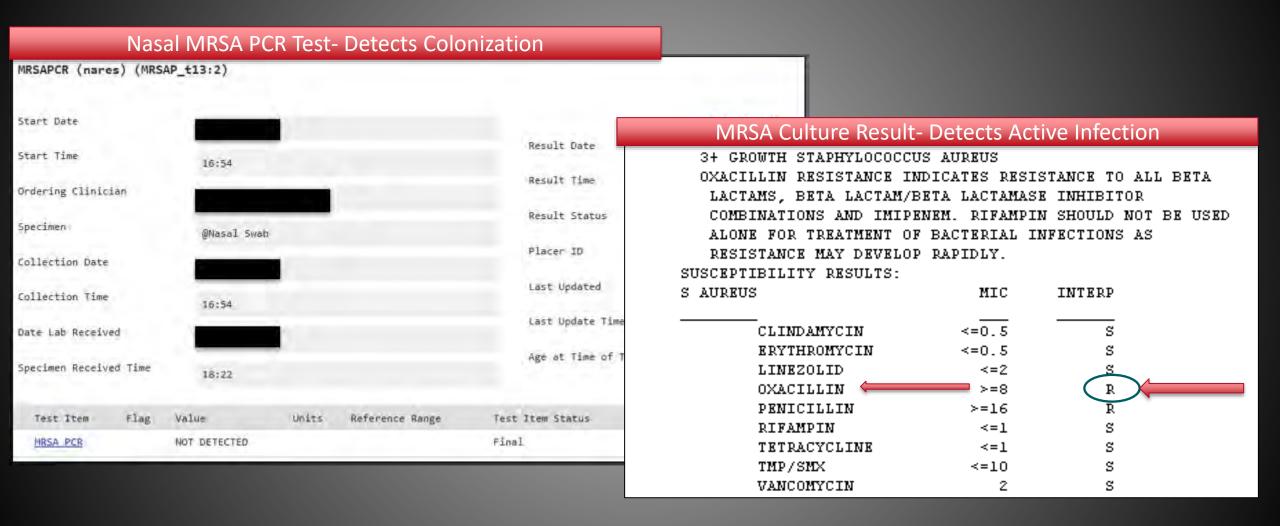
^{*} Contact tracing and colonization screening may not be indicated for these organisms

^{**}Contact precautions for acute/active infections or uncontained drainage/secretions

Examples of Microbiology Results and Interpretation



MRSA Infection Versus Colonization





CP- CRE Culture Result- Typically Detects Active Infection

Resistant E. coli

Escherichia coli (Isolate 1), 20,000 cfu/mL

*Carbapenemase detected. Notified

OF

Results reported to the State Health Department.

Sensitivity Analysis Isolate 1

Amikacin	<=16 5
Ampicillin	>16 R
Ampicillin/Sulbactam	>16/8 R
Aztreonam	>16 R
Cefazolin	>16 R
Cefepime	>16 R
Cefoxitin	>16 R
Ceftazidime	>16 R
Ceftazidime/Avibactam	>16 R
Ceftriaxone	>32 R
Ciprofloxacin	>2 R
Ertapenem	>1 R
Gentamicin	<=4 S
Imipenem	>4 R
Levofloxacin	>4 R
Meropenem	>8 R
Nitrofurantoin	<=32 S
Pipercillin/Tazobactam	>64 R
Tetracycline	<=4 5
Tobramycin	<=4 5
Trimethoprim/Sulfamethoxaz	>2/38
The state of the s	

S = Susceptible I = Intermediate R = Resistant

R* = Resistant, Predicted

ESBL = Extended Spectrum Beta-Lactamase

Blac = Beta-Lactamase Positive

N/R = Not reported, contact micro lab if testing is indicated

CP-CRE Infection Versus Colonization

Carbapenemase Screening Tes					
Test Item	Flag	Value	Units	Reference Range	Test Item Status
SPECIMEN SOURCE XXX		Rectal specimen submitted.			Final
CARBAPENEMASE RESISTANCE GENES ISLT PCR	Normal	Not Detected		NotDetected	Corrected
blaNDM Islt/Spm Ql	Normal	Not Detected		NotDetected	Final
blaKPC Islt/Spm Ql	Normal	Not Detected		NotDetected	Final
Test Item	Flag	Value	Units	Reference Range	Test Item Status
SPECIMEN SOURCE XXX		Rectal specimen submitted.			Final
CARBAPENEMASE RESISTANCE GENES ISLT PCR	Abnorma:	1 DETECTED		NotDetected	Corrected
blaNDM Islt/Spm Q1	Normal	Not Detected		NotDetected	Final
blaKPC Islt/Spm Q1	Normal	Not Detected		NotDetected	Final



CP-CRE Infection Versus Colonization

CP-CRE Culture Result- Typically Detects Active Infection

Test Item	Flag	Value	Units	Reference Range	Test Item Status	Comments	Sensitivities
SPECIMEN SOURCE XXX		Rectum			Final		
ANNOTATION COMMENT		None			Final		
MICROORGANISM/AGENT XXX	Abnormal	Organism submitted identified as: Klebsiella oxytoca / Raoultella ornithinolytica This isolate is positive for a KPC carbapenemase and may be clinically resistant to all beta-lactam antibiotics. Infectious Disease consult recommended.			Corrected		
Report Status					Corrected		
ORGANISM		Klebsiella oxytoca / Raoultella ornithinolytica This isolate is positive for a KPC carbapenemase and may be clinically resistant to all beta-lactam antibiotics. Infectious Disease consult recommended.			Corrected	Organism submitted identified as: Klebsiella oxytoca / Raoultella ornithinolytica This isolate is positive for a KPC carbapenemase and may be clinically resistant to all beta-lactam antibiotics. Infectious Disease consult recommended.	2
BACTERIAL SUSC PNL ISLT MIC		MIC			Final		
AMPICILLIN SUSC ISLT		>16			Final		R
AMPICILLIN+SULBAC SUSC ISLT		>16/8			Final		R
AZTREONAM SUSC ISLT		>16			Final		R
CEFAZOLIN SUSC ISLT		>16			Final		R
CEFEPIME ISLT MIC		>26			Final		R



Candida Auris Infection Versus Colonization

C. Auris Colonization Culture Result-Detects Colonization

Candida auris Colonization Culture

Final Report

4/28/2025

Healthcare Facility of Origin Name

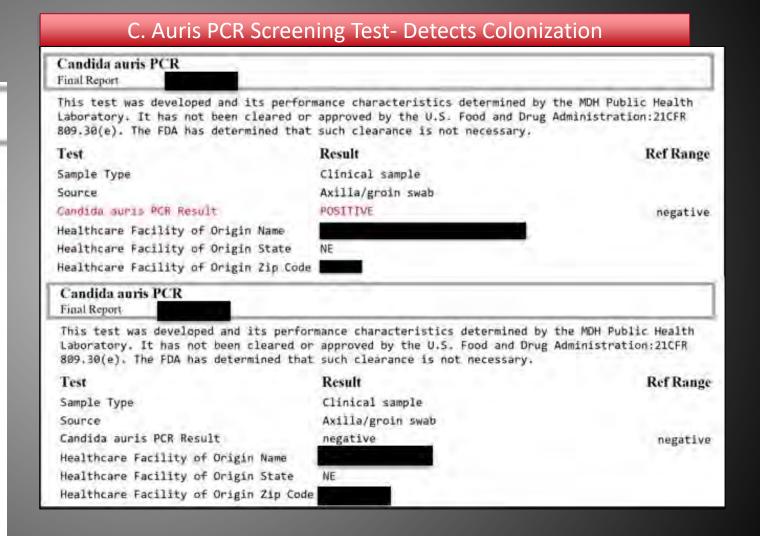
Healthcare Facility of Origin State NE

Healthcare Facility of Origin Zip Code

Sample Type Clinical sample

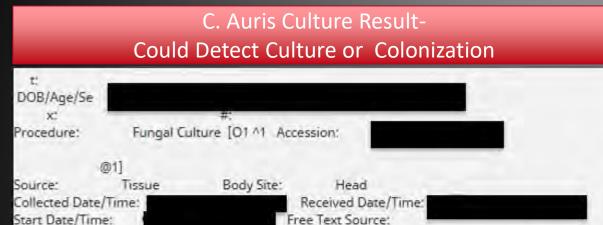
Source Axilla/groin swab

Culture Result Candida auris





Candida Auris Infection Versus Colonization

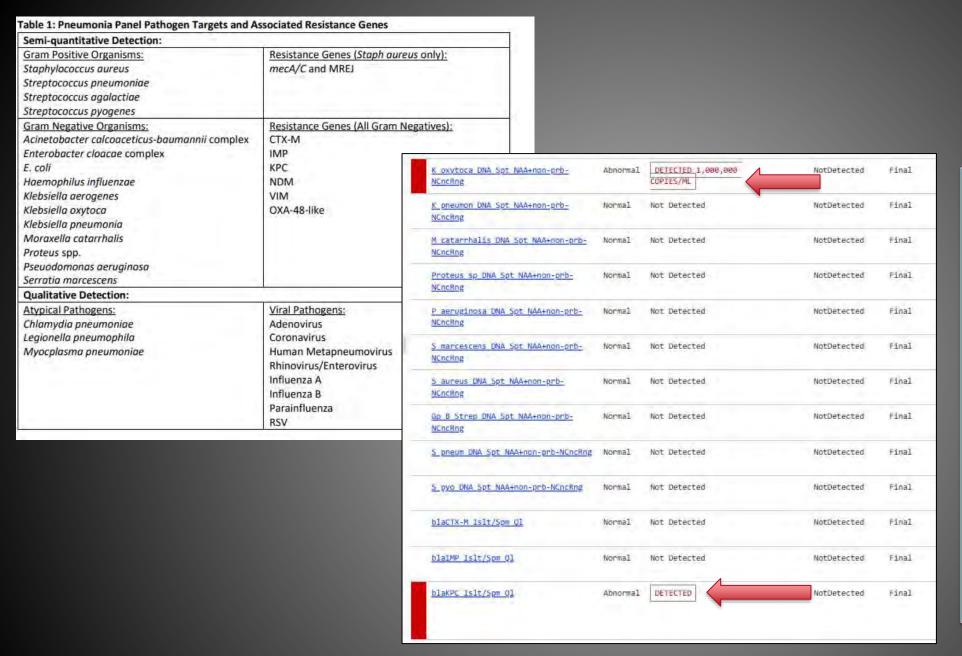


FINAL REPORTS

Verified Date/Time/Personne Rare Candida auris isolated.

Further investigation is needed; likely need to consider if the patient has active infection symptoms

	C. Auris Culture Result-
	Could Detect Culture or Colonization
DOB/Age/Se x: Procedure:	#: Fungal Culture [O1 ^1 Accession:
Source: Collected Date/T Start Date/Time: ***FINAL REPO Final Report [] Verified Date/Time	Free Text Source:
Moderate Candi With rare except	eomyces glabratus (prev Candida glabrata) isolated. da auris isolated. ions, Candida species are not thought to cause pulmonary infections and its epresents oropharyngeal colonization/contamination.



Multiplex Nucleic Acid Tests¹³

Multiplexed nucleic acid tests, which can detect several different bacteria, viruses and antibiotic resistance genes, can be done as quickly as one hour.

Nebraska Medicine

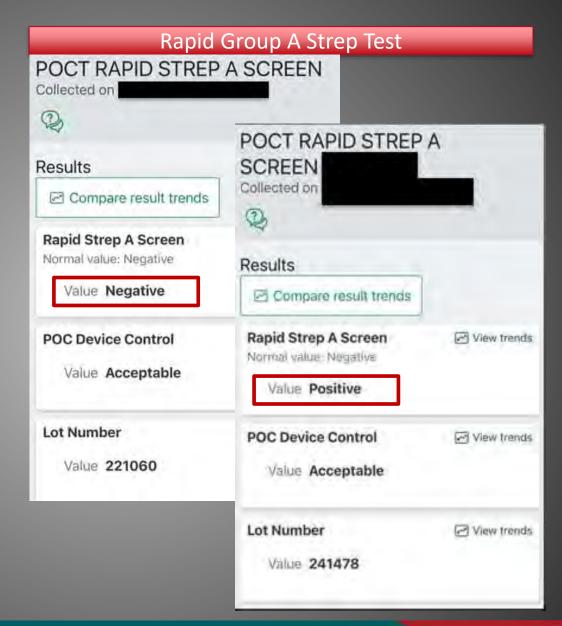
"Guidance on Use of
the Pneumonia Panel
for Respiratory
Infections



Other Examples

COVID/Influenza Combined PCR







Blood culture aerobic and anaerobic

Status: Final result Visible to patient: This result is not viewable by the patient. Next appt. None

Newer results are available. Click to view them now.

Source

Additional Information Culture Result 2wk ago

Blood, Peripheral Draw

None

Gram Stain result: Gram Positive Cocci in Clusters in Aerobic Bottle Only.

Time to detection: 18,22 hours

Methicillin Resistant Staphylococcus aureus (The Infectious Diseases Service may be consulted regarding treatment options for patients colonized or infected with

methicillin-resistant Staphylococcus aureus.)

Micro Report Status Organism 09/17/2014 Final

Methicillin Resistant Staphylococcus aureus

Resulting Agency Culture & Susceptibility

METHICILLIN RESISTANT STAPHYLOCOCCUS AUREUS

Antibiotic	Sanstiuta	MIC	Method	Status
Clindamycin	Resistant	>4	MIC	Final
Daptomycin	Susceptible	<=0.5	MIC	Final
Erythromycin	Resistant	>4	MIC	Final
Gentamicin	Susceptible	<=4	MIC	Final
Levofloxacin	Resistant	>4	MIC	Final
Linezolid	Susceptible	2	MIC	Final
Oxacillin	Resistant	>2 Susceptibility to Oxacillin can be used to predict susceptibility to Cefazolin	MIC	Final
Penicillin	Resistant	>8	MIC	Final
Rifampin	Susceptible	<=1	MIC	Final
Tetracycline	Resistant	>8	MIC	Final
Trimethoprim-Sulfa.	Susceptible	<=0.5/9.5	MIC	Final
Vancomycin	Susceptible	1	MIC	Final
Comments METHICILLIN RESIST	ANT STAPHYLOCOCCUS AUREUS			

Methicillin Resistant Staphylococcus aureus (The Infectious Diseases Service may be consulted regarding treatment options for patients colonized or infected with methicillin-resistant Staphylococcus aureus.)

Blood culture¹⁵

A blood culture can help diagnose bloodstream infections caused by bacteria or fungi.



Negative Blood Culture

Report Status

BLOOD CULTURE (BLDCU_t13) Start Date Order Details Trend Result Date Start Time 04:55 Result Time Ordering Clinician Result Status Corrected Specimen BLPH Placer ID Collection Date Last Updated Collection Time 05:47 Last Update Time Date Lab Received Age at Time of Test Specimen Received Time 05:53 Sensitivities Test Item Flag Value Units Reference Range Test Item Status Other Flags Collection Date Comments Blood, Peripheral Draw Final SPECIMEN SOURCE XXX ANNOTATION COMMENT IMP None Final MICROORGANISM/AGENT XXX No Growth at 5 Days Corrected

Final



04/24/2025 Final

Urine Cultures Examples

Procedure: Culture Urine [^2 @2] Accession:
Source: Urine Body Site:
Collected Date/Time: Received Date/Time:
Start Date/Time: ree Text Source:

FINAL REPORTS
Final Report []
Verified Date/Time/Personnel:
Normal urogenital flora isolated.

Example of a urine culture that grew normal urogenital flora

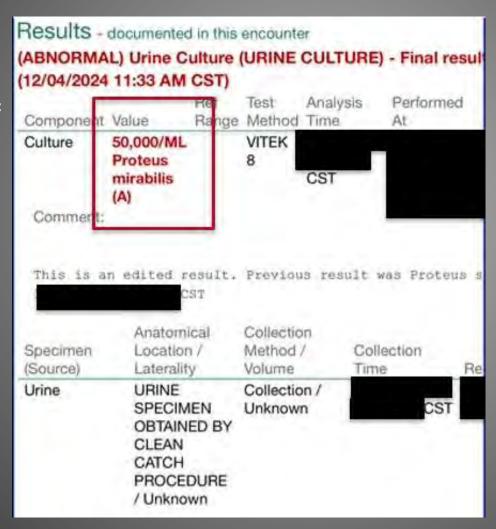
Cultures are often be ordered as a reflex test following an abnormal urinalysis.

			UA Dipstick
Procedure			Reference Range
UA Color	Yellow '1		[Light Yellow]
UA Clarity	Cloudy' ^{^1}		[Clear]
UA Glucose	2+**	mg/dL	[Negative]
UA Ketones	Trace`"	-1991	[Negative]
UA Spec Grav	1.015"	. 167 . 17	[1.005-1.030]
UA Blood	Negative 1		[Negative]
UA pH	7.0'1		[5.0-8.5]
UA Protein	Negative 1		[Negative]
UA Urobilinogen	1.0′1	EU/mL	[0.2]
UA Nitrite	Negative 1		[Negative]
UA Leuk Est	3+**1	mcL	[Negative]
UA Bilirubin	Negative 1		[Negative]
Micro?	Yes "1		
		<u> </u>	A Microscopic
ocedure	Result	Units	Reference Range

Urine Culture Examples¹⁵

A urine culture may show the presence of specific bacteria, such as E. coli or Proteus Mirabilis.

The results may also include the number of bacteria per milliliter (CFU/ml)



Sensitive example:

Sensitivity Analysis

Escherichia coli (Isolate 1), >100,000 cfu/mL

Isolate 1

Serisitivity Arialysis	130late 1
Amikacin	<=16 S
Ampicillin	<=8 S
Ampicillin/Sulbactam	<=8/4 S
Aztreonam	<=4 S
Cefazolin	<=2 S
Cefepime	<=2 S
Cefoxitin	<=8 S
Ceftazidime	<=1 S
Ceftriaxone	<=1 S
Ciprofloxacin Enterobact	. S
Ertapenem	<=0.5 S
Gentamicin	<=4 S
Imipenem	<=1 S
Levofloxacin Enterobact	S
Meropenem	<=1 S
Nitrofurantoin	<=32 S
Pipercillin/Tazobactam	<=16 S
Tetracycline	<=4 S
Tobramycin	<=4 S
Trimethoprim/Sulfametl	oxaz <=2/38 S

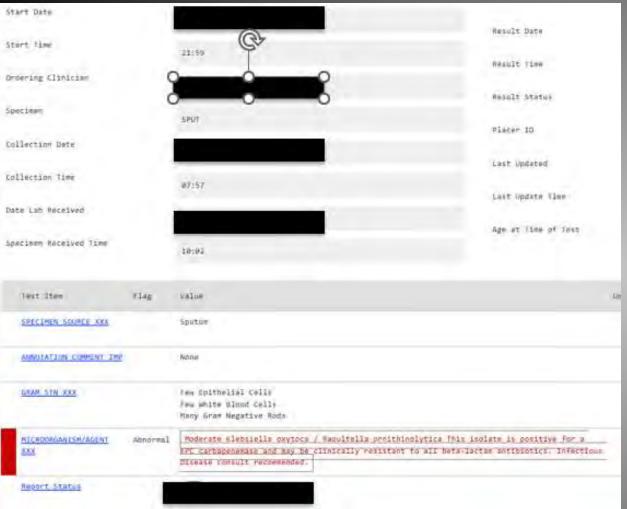
S = Susceptible I = Intermediate R = Resistant R* = Resistant, Predicted

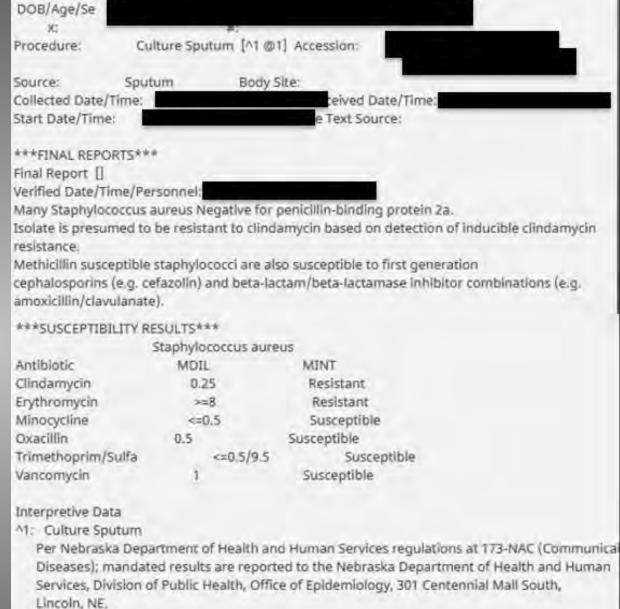
ESBL = Extended Spectrum Beta-Lactamase

Blac = Beta-Lactamase Positive

N/R = Not reported, contact micro lab if testing is indicated

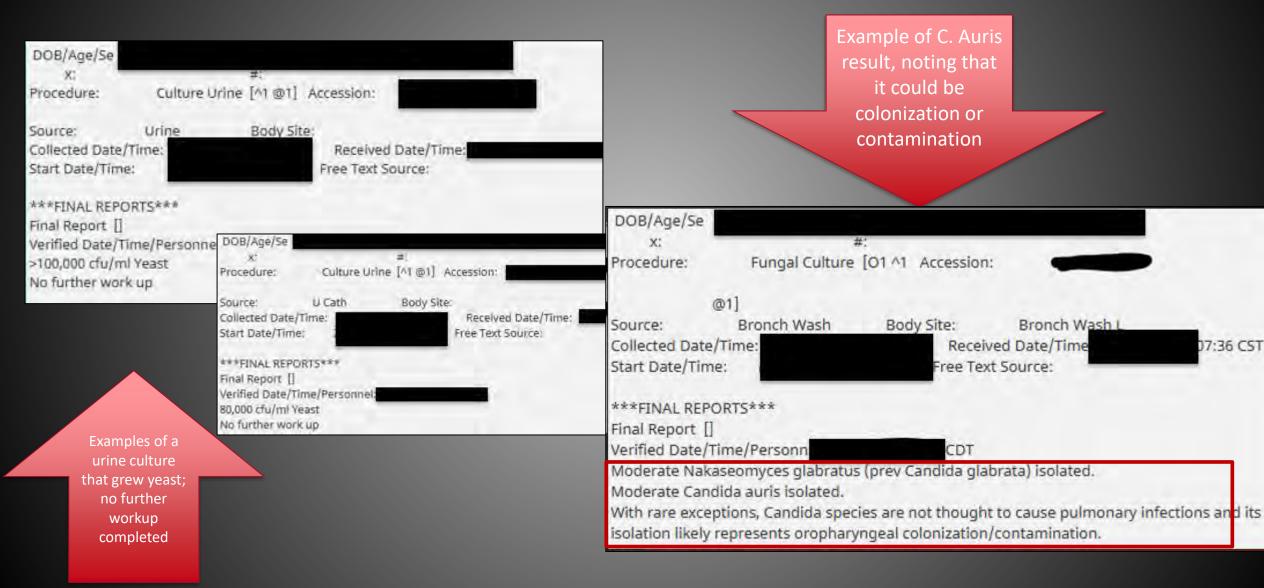
Sputum Culture Examples







Yeast/Fungal Cultures Examples



07:36 CST

Lab Report Examples: ESBL+ Escherichia Coli

Resistance Example 1:

Culture Result::

>100,000 col/ml Escherichia col POSITIVE for Extended Spectrum Beta-Lactamase (ESBL) This is an extended spectrum Beta-lactamase (ESBL) producing strain which is clinically resistant to cephalosporins and aztreonam. Although isolates that produce ESBLs may be susceptible or intermediate to Piperacillin/Tazobactam, clinical efficacy has not been documented. The Infectious Disease Service may be consulted.

ORGANISM:

Escherichia coli

>100,000 col/ml Escherichia coli POSITIVE for Extended Spectrum Beta-Lactamase (ESBL) This is an extended spectrum Beta-lactamase (ESBL) producing strain which is clinically resistant to cephalosporins and aztreonam. Although isolates that produce ESBLs may be susceptible or intermediate to Piperacillin/Tazobactam, clinical efficacy has not been documented. The Infectious Disease Service may be consulted.

METHOD:

Tigecycline:

Minocycline:

MIC

Amikacin: <=16 Susceptible >16 Resistant Ampicillin: Amp-Sulbactam: 8/4 Resistant Aztreonam: 16 Resistant Cefazolin: >16 Resistant 8 Resistant Cefepime: Cefotaxime: >32 Resistant Ceftazidime: 16 Resistant Ceftriaxone: >32 Resistant Cefuroxime* >16 Resistant <=2 Susceptible Gentamicin: Levofloxacin: >4 Resistant <=1 Susceptible Meropenem: Pip/Tazo: <=8 Resistant Tobramycin: <=2 Susceptible Trimethoprim-Sulfa: <=0.5/9.5 Susceptible Amoxacillin/Clavu anic Acid: <=8/4 Resistant Cefoxitin: <=8 Resistant Ertapenem: <=0.5 Susceptible Nitrofurantoin: <=32 Susceptible Tetracycline: >8 Resistant

<=2 Susceptible

8 Intermediate

Resistance Example 2:

Bacteria Ur Cult

ESBL Escherichia coli

Final

>100000 CFU/mL Escherichia coli ESBL

This organism has been confirmed as an ESBL producer.



Factors to Consider During Interpretation



URIME CULE, VOIDED COLLECT DATE AND TIMES

SPECIMEN DESCRIPTION: SPECIAL REQUESTS:

INCHATE DI

Manager Manager Manager In B

MIC

1 4

1 >16

1 >16

1 < 41

1.28

(>2/88 ss - seceptius

\$ # 4800/DAY

BRADES - MOST MANUFLABILIE

Ucina Noné

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BALLAND HULTATERSKERT ON " ALH

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R | \$3

R 1 99

COLONY COUNT:

>100,000 col/ar

CULTURE:

DRUG

Ampicillia

Coffrience

Coftazidimo

Cefotagino

Cotazolin

Cafuroxide

Gentamicio

ted penem

Karopentia

Tobrosycia

TMP/SHX

Ertapenem

Amox/Clavelana) 16/8

Cigcofloxacia | >2

Hitroforantoin (<-32

Levellonacin | >4

Pio/Tazobactual <=18

Rycherichia coli This organism has been confirmed an an ESEL producer.

The testing includes pathogens (bacterial, parasitic and viral) commonly known to cause gastroenteritis. Note that not all individuals with positive findings will present with symptoms. Many factors, including the health of the individual (such as immune health, digestive function, and microbiome balance), the transient nature of most pathogens, and the presence and expression of virulence factors, all contribute to pathogen virulence and individual symptoms.

Reporting¹¹

Clear and Accurate

BACTERIAL PATHOGENS	Result	Reference
Campylobacter	<di>40</di>	< 1.00e3
C. difficile Toxin A	6.77e8 High	4 1.00e3
C. difficile Toxin B	2.65e6 High:	< 1.00e3
Enterohemorrhagic E. coli	* A	< 1.00e3
E. coli O157		< 1.00e3
Enteroinvasive E. coli/Shigella	AND NO.	< 1.00e3
Enterotoxigenic E. coli LT/ST	₹ ∱⊲	< 1.00e3
Shiga-like Toxin E. coli stx1	-dl	< 1.00e3
Shiga-like Toxin E. coll stx2	-di	< 1.00e3
Salmonella 🧳 🕽	*dl	< 1.00e4
Vibrio cholerae	<di< td=""><td>< 1.00e5</td></di<>	< 1.00e5
Yersinia enterocolitica	×dI	< 1.00e5
PARASITIC PATHOGENS		
Cryptosporidium	<di -<="" td=""><td>< 1.00e8</td></di>	< 1.00e8
Entamoeba histolytica	<dl< td=""><td>< 1.00e4</td></dl<>	< 1.00e4
Giardia	<0.	< 5.00e3
VIRAL PATHOGENS		
Adenovirus 40/41	<dl< td=""><td>< 1,00e10</td></dl<>	< 1,00e10
Norovirus GI/II	<@	< 1.00e7

Essential for several reasons, including:

- Patient care
- Patient safety
- Effective treatment

Examples of unclear/hard to read lab results

CONSTRUCT - BRICK TO FOR BY PRODUCES UNLESS CRIMINATE ROYAL



Intrinsic Resistance⁵

- Some bacteria have inherent resistance to certain antibiotics
 - Example: Pseudomonas and ceftriaxone
- Results of antibiotics that are intrinsically resistant will often be suppressed in the results

Note that Ceftriaxone is not an antibiotic reported for this Pseudomonas Aeruginosa result.

Pseudomonas Aeruginosa AST Results		
	Pseudoi	monas aeruginosa
		MIC
Aztreonam	<=4	Susceptible
Cefepime	<=2	Susceptible
Ceftazidime	4	Susceptible
Ceftolozane/tazobactam	<=2	Susceptible
Levofloxacin	<=0.5	Susceptible
Meropenem	<=1	Susceptible
Piperacillin + Tazobactam	<=8	Susceptible
Tobramycin	<=2	Susceptible ¹



Enterobacterales		Antimicrobial Agent									
Organisms	Ampicillin	Amoxicillin-clavulanic acid	Ampicillin-sulbactam	Ticarcillin	Cephalosporin I: Cefazolin, Cenhalothin	Cephamycins: Cefoxitin, Cefotetan	Cephalosporin II: Cefuroxime	Imipenem	Tetracyclines and Tigecycline	Nitrofurantoin	Polymyxin B/Colistín
Citrobacter freundii	R	R	R	35	R	R	R				
Citrobacter koseri	R			R							Ĩ
Klebsiella aerogenes (prev.Enterobacter)	R	R	R		R	R	R			70	
Enterobacter cloacae complex	R	R	R		R	R	R				
Escherichia coli	1	T	here is no	intrinsic	resistanc	e to beta	-lactams	in thi	s organ	nism	
Escherichia hermannii	R			R							
Hafnia alvei	R	R	R		R	R					
Klebsiella pneumoniae	R			R							
Morganella morganii	R	R	النبيا		R		R	*	R	R	R
Proteus mirabilis	No	intrinsi	c resistanc	e to penio	illins and c	ephalospo	rins	*	R	R	R
Proteus penneri	R				R		R	*	R	R	R
Proteus vulgaris	R				R		R	*	R	R	R
Providencia rettgeri	R	R)ı = ı'		R			*	R	R	R
Providencia stuartii	R	R			R				R	R	R
Salmonella and Shigella spp	No intrinsic resistance to beta-lactams in these organisms 1 ST AND 2 ^{NO} generation cephalosporins may appear active in vitro, but are not effective clinical										
Serratia marcescens	R	R	R		R	R	R			R	R
Yersinia enterocolitica	R	R	$\mu = \mu$	R	R						

Note: Cephalosporins III, cefepime, aztreonam, ticarcillin-clavulanate, piperacillin-tazobactam, and the carbapenems are not listed, because there is no intrinsic resistance in Enterobacteiaceae.

Intrinsic Resistance^{5,10}

Example of organisms with their respective intrinsic antimicrobial resistance from LMH Health
Antimicrobial and Clinical
Microbiology Guidebook,
9th Edition, January 2023.

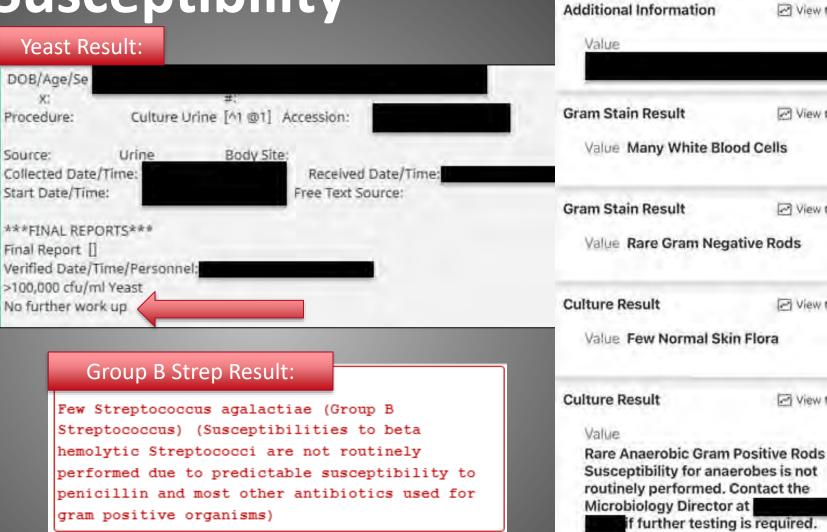


Predictable Susceptibility¹¹

Some bacteria, like those causing common don't typically undergo routine susceptibility testing because their susceptibility to common antibiotics is often predictable OR because they are commonly considered to be contaminants.

Examples:

- Streptococcus pyogenes (Group A Strep)
- Candida albicans
- Streptococcus agalactiae (Group B Strep)
- **Anaerobic Gram-Positive** Rods





Anaerobic Gram-Positive

Rods Result:

✓ View trends

✓ View trends

✓ View trends

P View trends

View trends

Chart in the Workbook on page 18

CONTAMINANTS **VS PATHOGENS**

BLOOD

Normally sterile

Pathogens - any organism Isolated

Likely Contaminants

Coagulase-negative staphylococci Alpha-hemolytic streptoceco Bacillus son Corynebacterium spp. (except C Jelkelum) Propionibacterium acnes NOTE: Must take into consideration flow many cultures were drawn versus how many are positive and what the organism is

TISSUE AND BODY FLUIDS

Should be sterile

Pathogens = any organism isolated; use judgement to evaluate the possibility of normal flora being present in relation to the source of the

Normal Flora

Eve/Ear Coagulase-negative **staphylococcs** non-heimiolytic streptococci alpha-hemolytic streptococci Dightherolds Skin Coagulase-negative staphylococci Propionibacterium acnes. dightheroids alpha-hemolytic streptococci Bacillus spp.

GENTTAL

Pathogens

B-hemiolytic strephocacci Listeria spp. Gardnerella vaginalis Predominant numbers of 5. Predominant numbers of yearst

Neisseria gonorrhoeae

Normal flora

Staphylococcus app. Lactobacillus spp. Dishtheroids Enterococcus spp. Streptococcus app. Gram-negative rods Anaerobes Yearst

DEINE

Should be sterile

Pathogens.

Enterobateriaceae Enteracoccus spp. Pseudomorus spp. and other non-fermenters

group B Streptococcus (Streptococcus agalactiae) S. aureus and S. saprophyticus

Likely Contaminants

Diontherolds coagulase-negative staphylococci alpha-hemolytic streptococci Lactobacillus spp.

Bacillus spp. NOTE: significance of organism is determined by colony count and symptoms

GASTROINTESTINAL TRACT

Pathogens

Salmonella spp. Shigella sop-Campylobacter Jejuni E. call 0157:H7 Aeromonas/Flesiomonas spp. Yersinia enterocolitica Vibrio app. Clastridium difficile (toxin) 5. aureus (in the context of enterotaxin food poisoning) Helicobacter pylori (antigen)

Normal Flora

Enterobacteriaceae Staphylococcus spp. Streptococcus spp. Enterocaccus spp. Pseudomenas spp. Anaerobes Yearsh

RESPIRATORY TRACT

Pathogens

Group A Streptococcus-(Streptococcus pyogenes) Streptococcus pneumoniae Predominant S. aureus H. Influenzae Neisseria meningitidis/gonorrhoeae Predominant Enterobacteriaceae Predominant Pseudomonas spp. and other non-fermenters Carynebacterium diphtheriae Bordetella pertussis Legionella pneumophila Mycobacterium spp. Nocardia spp. Predominant Moraxella catarrhalis

Normal Flora

Staphylococcus spp.(CDag. negative) alpha-hemolytic streptococci Gram-negative rods B-hemolytic streptococci other than group A Neisseria spp. Enteroceccus spp. Corynebacterium spp. Bacillus spp. Yeast Anaerobes Haemophilus spp: Micrococcus spp. Stomatococcus spp. NOTE: amount of organism

present, source of culture, and

patient age may determine

significance as a pathogen.

Pathogen-Specific Considerations¹

Normal flora versus probable pathogen

- Example- staph epi on the skin
- Example- candida in the lungs

Specimen Contamination

Example-Bacillus spp. on a blood culture

https://www.unmc.edu/intmed/_documents/id/asp/clinicmicro-id_guidebook.pdf



Specimen Contamination^{1,7}

Contamination of Specimens (no more than 2 potential pathogens per culture)

 3 or more indicate a poorly collected specimen

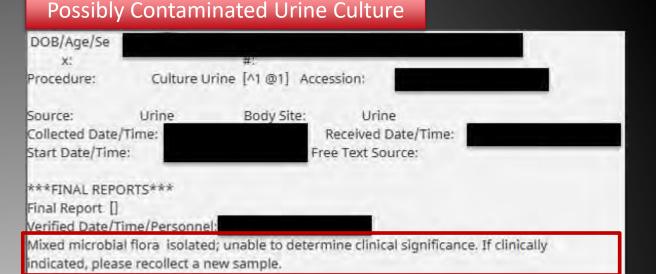
Possibly Contaminated Blood Culture

See Direct Blood Pathogen Identification Panel for rapid identification panel.

Staphylococcus species coagulase negative Probable contaminant, clinical significance of 1 positive out of 2 sets is unclear. If further workup is required, please contact Microbiology

Possibly Contaminated Urine Culture

Test Item	Flag	Value	Units	Reference Range	Test Item Status
SPECIMEN SOURCE XXX		Urine Clean Catch			Final
ANNOTATION COMMENT IMP		None Reflexed from S52625			Final
MICROORGANISM/AGENT		Three or more colony types indicate contamination. If clinically indicated, please submit a new specimen.	4		
Report Status					Final





Summary¹

- Interpreting microbiology results requires a combination of technical expertise and clinical judgment.
- It is important to accurately identify the microorganisms, understand their susceptibility patterns, and consider the patient's overall clinical picture.
- The clinical microbiology laboratory is an important partner in the practice of infection prevention.

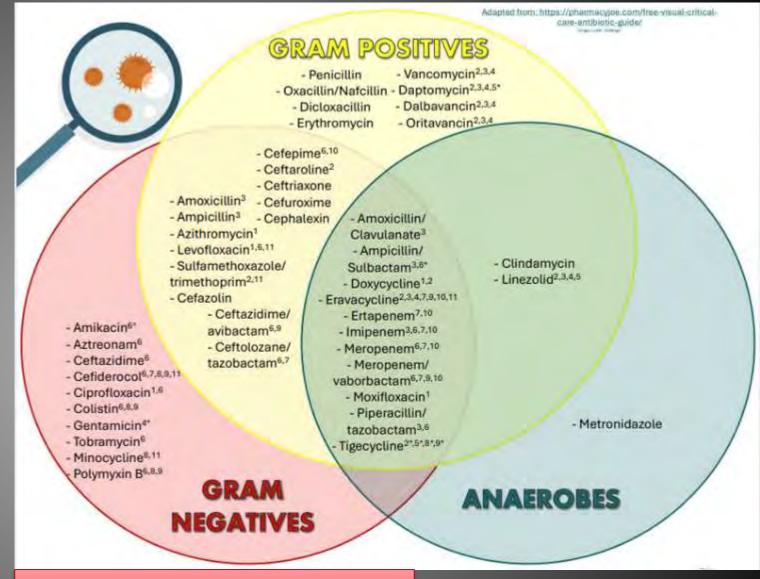


Figure in the Workbook on page 19



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