

TARGETED MDROS IN INDIANA

CALEB COX
SENIOR MDRO EPIDEMIOLOGIST

04/15/2025

OUR MISSION:

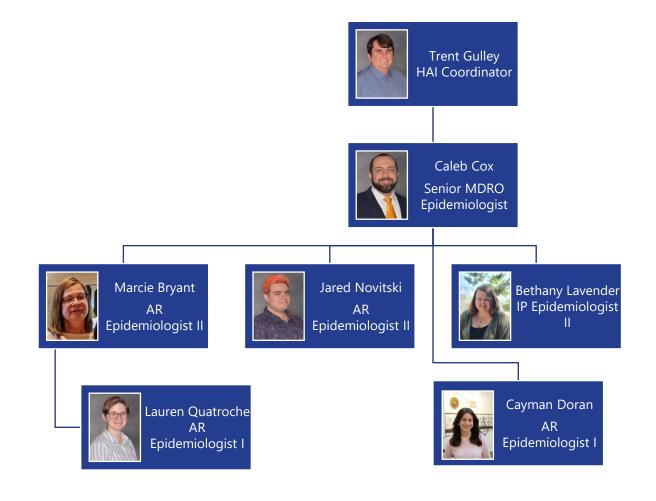
To promote, protect, and improve the health and safety of all Hoosiers.

OUR VISION:

Every Hoosier reaches optimal health regardless of where they live, learn, work, or play.



Antimicrobial Resistance Team





Communicable Disease Rule

Announcement: In March 2023, the Indiana Department of Health (IDOH) announced changes to Indiana's Communicable Disease (CD) Rule (410 IAC), including streamlined timeframes for reporting and updates to reportable diseases. Indiana Code Title 16 (Health 16-41-2-1) was amended in 2019. This amendment allows the IDOH to publish and update the list of reportable communicable diseases and control measures on the IDOH website. External documents have been created to house this information, which will allow for updates and changes to be made more easily in the future.

Communicable disease reporting changes went into effect April 1.

Key Updates

- The reporting timeframes have been streamlined to two options: immediately and within one working day. The number of immediately reportable diseases has decreased.
- The reportable disease list has been removed from the Indiana Communicable Disease (CD) Rule. Separate documents have been
 created for the reportable disease list for providers as well as a reportable result/pathogen list for laboratories. By listing this information
 in external documents, rather than within the rule, it will be easier to make changes and keep the information updated. Additionally,
 these documents will be more easily accessible and convenient for health care providers, hospitals, medical laboratories, and local
 health departments.

The following documents cover the changes in greater detail:

- Indiana Reportable Disease List for Healthcare Providers and Hospitals
- 2. Indiana Reportable Result/Pathogen List for Laboratories
- 3. Summary of Reporting Changes Document
- 4. Frequently Asked Questions Document
- 5. Local Health Department Webcast Slides





2025 Indiana Reportable Disease List for Healthcare Providers and Hospitals

410 IAC 1-2,5-75 & 76



REPORT IMMEDIATELY ON SUSPICION

Anthrax Botulism Cholera (Vibrio cholerae O1, O139, or toxigenic) Eastern equine encephalitis virus (EEEV) disease Hemolytic uremic syndrome (HUS), post-diarrheal Hepatitis, viral, Type B, pregnant woman (acute and chronic) or perinatally exposed infant Influenza A. Novel Measles (Rubeola) Melioidosis (Burkholderia pseudomallei) Meningococcal disease, invasive Middle East respiratory syndrome coronavirus (MERS-CoV) Plague Poliomyelitis

Rubella (German Measles) Rubella congenital syndrome SARS-associated coronavirus (SARS-CoV)

Smallpox (Variola infection) Tularemia

Viral hemorrhagic fever, filoviruses Ebola virus

Marburg virus Viral hemorrhagic fever, other

Crimean-Congo hemorrhagic fever virus Guanarito virus Junin virus Lassa virus Lujo virus Machupo virus Sabia virus

TO REPORT:

Step 2:

- Immediately Reportable: complete steps 1-2.
- · Within One Working Day: complete step 2

Step 1: Call **317-233-7125** 317-233-1325 (After hours)



- NBS users: Report conditions via Morbidity Report in NBS
- · Non-NBS users: Report with this form

REPORT WITHIN ONE WORKING DAY

Acquired Immunodeficiency Syndrome (AIDS) Acute Flaccid Myelitis (AFM) Anaplasmosis

Animal bite or exposure

Rabies, human

Arboviral disease or infection, domestic: California serogroup viruses,

Jamestown Canyon virus,

La Crosse virus, Powassan virus.

St. Louis encephalitis virus,

West Nile virus, Western equine encephalitis virus

Arboviral disease or infection, imported:

Chikungunya virus, Dengue virus, Japanese encephalitis, Oropouche virus, Yellow fever,

Zika virus Babesiosis Brucellosis Campylobacteriosis Candida auris

Carbapenemase-Producing Organisms (CPO)

Chancroid Chlamydia trachomatis, genital infection Lymphogranuloma venerum

Coccidioidomycosis COVID-19-associated deaths (all ages) Cronobacter infection, invasive, infants

(younger than 1 year of age)

Cryptosporidiosis Cyclosporiasis Ehrlichiosis Escherichia coli (E. coli) infection (Shiga toxinproducing E. coli (STEC) including, but not limited to, E. coli O157 and other serogroups)

Giardiasis Gonorrhea

Disseminated gonococcal infection Haemophilus influenzae, invasive disease, (including antimicrobial susceptibility

testing)

Hansen's disease (leprosy)

Hantavirus infection (pulmonary and non-pulmonary), including, but not limited to: Sin Nombre virus,

Seoul virus Hepatitis, viral, Type A

Hepatitis, viral, Type B (acute and chronic) Hepatitis, viral, Type C (acute and chronic)

Hepatitis, viral, Type C, pregnant woman (acute or chronic) or perinatally exposed infant

Hepatitis, viral, Type Delta Hepatitis, viral, Type E

Hepatitis, viral, trype E Hepatitis, viral, unspecified

Histoplasmosis HIV infection

HIV infection, pregnant woman or perinatally exposed infant

Influenza-associated deaths (all ages) Latent tuberculosis infection (LTBI)

Legionellosis Leptospirosis

Listeriosis Lyme disease

Malaria Mpox (formerly known as Monkeypox) Multisystem Inflammatory Syndrome in adults (MIS-A) Multisystem Inflammatory Syndrome in

children (MIS-C)

Pandrug-resistant Organisms Pertussis (whooping cough) Psittacosis

Q Fever

Rabies, postexposure prophylaxis administration

Salmonellosis, non-typhoidal Shigellosis

Spotted fever rickettsiosis, including Rocky Mountain Spotted fever

Streptococcus pneumoniae, invasive disease (including antimicrobial

susceptibility testing)
Streptococcus, Group A, invasive disease
Syphilis

Tetanus

Toxic shock syndrome (streptococcal or staphylococcal)

Trichinellosis

Tuberculosis disease, reportable upon suspicion

Typhoid and paratyphoid fever, cases and carriers

Vancomycin-resistant Staphylococcus aureus (VRSA) and Vancomycin-intermediate Staphylococcus aureus (VISA)

Varicella (chickenpox) Vibriosis (non-cholera Vibrio infection) Yersiniosis, non-*pestis*

2025 Indiana Reportable Disease List for Healthcare Providers and Hospitals

410 IAC 1-2,5-75 & 76



IMMEDIATELY REPORTABLE OUTBREAKS

- Any disease required to be reported as listed above.
- 2. Newborns with diarrhea in hospitals or other institutions
- 3. Foodborne or waterborne diseases in addition to those specified above
- Streptococcal illnesses
- Conjunctivitis
- 6. Impetigo
- Clusters or suspected outbreaks of any disease associated with hospitals and healthcare facilities
- 8. Influenza-like illness
- 9. Viral meningitis
- 10. Unusual occurrence of disease
- Any disease (e.g. anthrax, plague, tularemia, Brucella species, smallpox, or botulism) or chemical illness considered a bioterrorism threat, importation, or laboratory release

OTHER REPORTABLE NON-COMMUNICABLE CONDITIONS AND DISEASES

- Report all blood lead results (capillary and venous) in children and adults within one week (410 IAC 29-3-1)
- Report confirmed cases of cancer occurring in residents diagnosed or treated in Indiana to the state cancer registry (410 IAC 21-1-2)

1/1/25

Reporting – Within One Working Day

How to report:

- Option 1: create morbidity report in NBS
 - o attach all documents to morbidity report
- Option 2: fax all documents to IDOH
 - o secure fax number: 317-234-2812

Documents to include when reporting:

- history and physical
- all relevant lab reports
- antimicrobial susceptibility testing (AST) report
- Confidential Report of Communicable Diseases (if faxing)



Acronyms

- CRE Carbapenem-resistant *Enterobacterales*
- CP-CRE Carbapenemase-producing carbapenem resistant Enterobacterales
- CRPA Carbapenem-resistant Pseudomonas aeruginosa
- CRAB Carbapenem-resistant *Acinetobacter baumannii*
- CPO Carbapenemase-producing ₀rganism
- MDRO Multidrug-resistant organism
- CDC- Centers for Disease Control and Prevention
- NBS- National Electronic Disease Surveillance System (NEDSS) base system



Carbapenemase Producing Organisms

- Carbapenemase-producing Enterobacterales, Pseudomonas aeruginosa, and Acinetobacter baumannii from any site OR
- Isolates of *Enterobacterales, Pseudomonas aeruginosa*, and *Acinetobacter baumannii* that are resistant to carbapenems but have not been tested for Carbapenemase production

Condition Name in NBS:

- Carbapenemase-Producing Organisms (CPO), Clinical
- Carbapenemase-Producing Organisms (CPO), Screening

Reporting Timeframe:

Within One Working Day

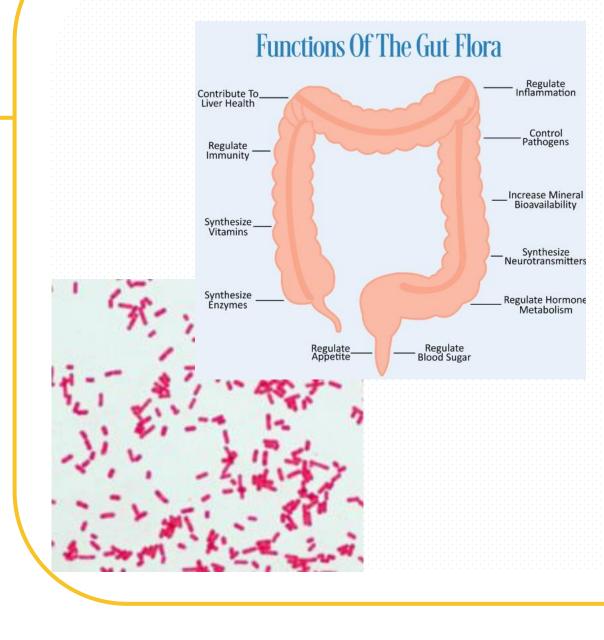


Enterobacterales

Enterobacterales

- A category of bacteria that live in the GI tract of humans
- These bacteria are all gram-negative rods
- Examples
 - E. coli
 - Klebsiella sp.
 - *Enterobacter* sp.
 - Serratia sp.
 - Proteus sp.
 - ° Citrobacter sp.

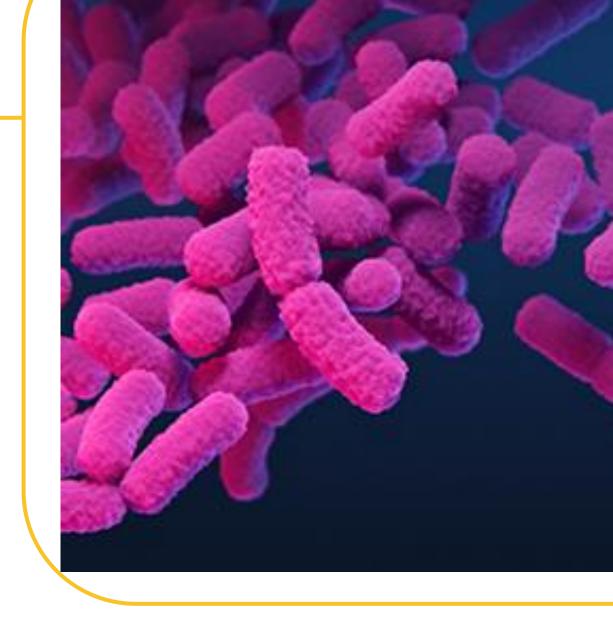




CRE

Definition: Any *Enterobacterales* that are not susceptible (i.e. intermediate or resistant) to a carbapenem antibiotic

- Enterobacterales can be resistant to carbapenems through several resistance mechanisms
- Carbapenemase production is currently the most concerning





CRPA

- Pseudomonas is commonly found in the soil and water.
- Pseudomonas aeruginosa can cause infections in the blood, lungs, or other parts of the body after surgery.
- Can colonize a patient without causing infections.
- Can be intrinsically resistant to Ertapenem





CRAB

- Acinetobacter is commonly found in soil and water.
- Acinetobacter baumanii can cause infections in the blood, urinary tract, lungs, and in wounds.
- Can colonize a patient without causing infections.





CPO

Screening definition:

 Organisms cultured specifically for screening from any body site that are found to produce a carbapenemase enzyme

Clinical definition:

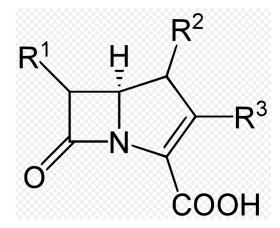
 Organisms cultured during routine care from any body site that are found to produce a carbapenemase enzyme



Carbapenem vs carbapenemase

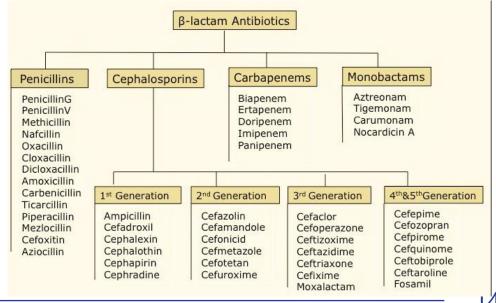
Carbapenem

- Class of broad-spectrum antibiotics
- Treatment is typically reserved for severe or highly resistant infections only
- Includes
 - Meropenem
 - Ertapenem
 - Doripenem
 - Imipenem



Carbapenemase

- An enzyme that can break down carbapenems (and all beta-lactams)
- Includes: KPC, VIM, OXA-48, IMP, NDM





Carbapenamase Producing Organisms (CPO) Data

2023: 405

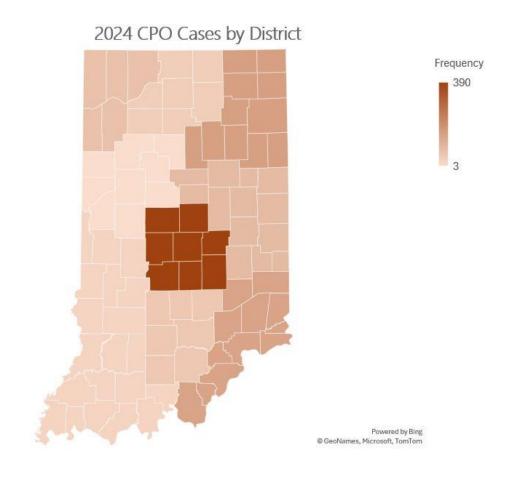
• Clinical: 328 (81.0%)

• Screening: 77 (19.0%)

2024: 440

• Clinical: 399 (90.7%)

• Screening: 41 (9.3%)





Indiana 2024 CPO Cases

Organism:

- Klebsiella spp.: 105
- E. coli: 46
- Enterobacter spp: 23
- Other: 77
- Acinetobacter spp: 170
- Pseudomonas spp: 8

Mechanism:

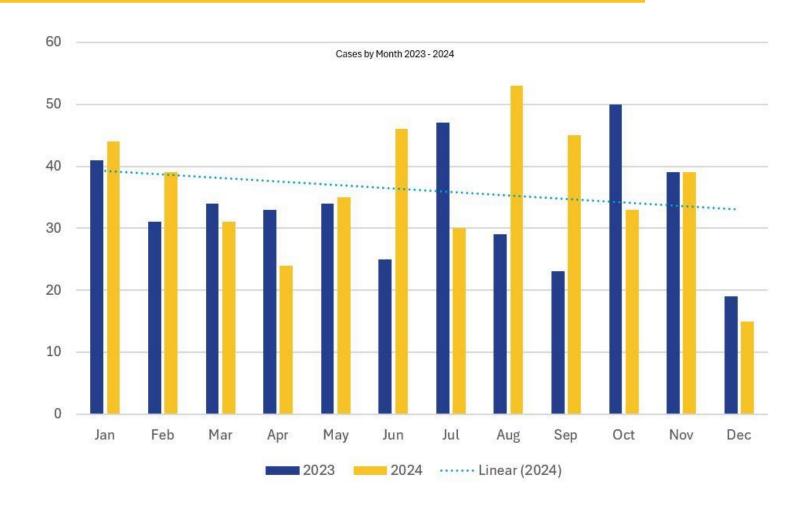
- KPC: 163
- VIM: 30
- NDM: 37
- IMP: 7
- OXA-48: 16

Acinetobacter only

- OXA-23: 58
- OXA-24/40: 109
- OXA-58: 0



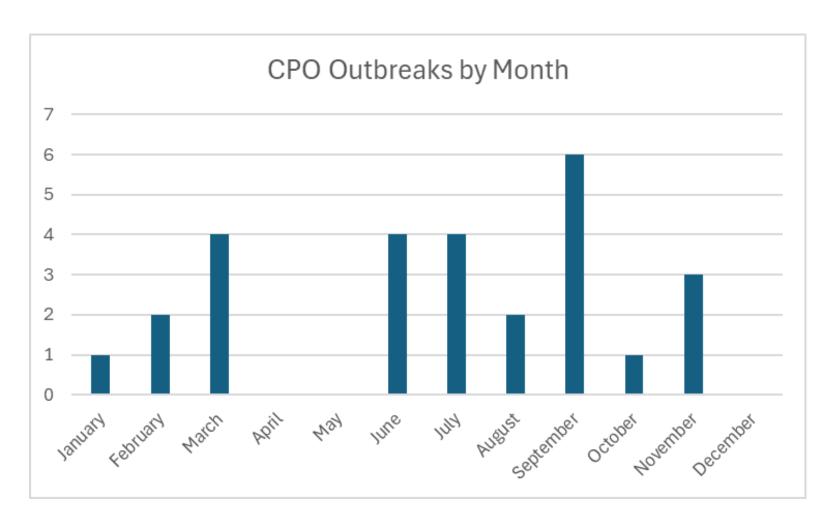
Carbapenamase Producing Organisms (CPO) Trends





CPO Outbreaks

- Total Outbreaks: 28
 - CRAB OXA-24/40: 12
 - CRAB OXA-23: 5
 - *E. coli* NDM: 2
 - K. pnuemoniae KPC: 2
 - CP-CRPA mCIM+/PCR Neg: 1
 - MRSA: 1
 - Multiple MDROs: 1
 - Multiple CP-CRE KPC: 1
 - S. marcescens KPC: 1
 - No MDROs: 1





Candida auris (C. auris)

- Candida auris clinical isolates representing both invasive (e.g., blood and CSF) and noninvasive sources (e.g., urine, wound, and respiratory tract), OR
- Colonized cases diagnosed via a skin or axilla/groin swab used to screen for C. auris,

Condition Name in NBS:

- Candida auris, clinical
- Candida auris, colonized

Reporting Timeframe:

 Within One Working Day



Background

- Candida auris (C. auris) is an emerging multidrug-resistant yeast that can cause invasive infections associated with high mortality.
- C. auris can persist on surfaces and medical equipment, spread between patients, and lead to outbreaks in healthcare settings.
- Approximately 40% of the U.S. clinical cases have been bloodstream infections; the rest involved non-sterile sites such as urine, respiratory tract, and wounds.
- C. auris can colonize the skin, nares, and other body sites. Patients can be colonized with C. auris for long periods of time, even after successful treatment of infection.
- Although patients colonized with *C. auris* in non-sterile sites may not need medical treatment, they can be a source of transmission to other individuals.

*Source CORHA Candida auris https://www.corha.org/wp-content/uploads/2021/08/Candida-auris-Recommendations-for-Healthcare-Outbreak-Response.pdf



Background

- Risk of infection or colonization with *C. auris* is greatest among people:
 - With extensive healthcare exposures, especially in long-term care facilities providing ventilator care
 - Infected or colonized with another multidrug-resistant organism, especially carbapenemase-producing organisms (CPOs)
 - With invasive medical devices such as central venous catheters and with tracheostomy, or gastrostomy tubes



C. auris clades

- Clade A group of biological taxa (such as species) that includes all descendants of one common ancestor
- Originally referred to by their geographic origins.
 - South Asian (Clade I)
 - East Asian (Clade II)
 - African (Clade III)
 - South American (Clade IV)
- Clade V isolates from Iran
- Novel Clade VI isolated in Bangladesh, first published June 2024



Clades, Indiana, and whole genome sequencing (WGS)

- WGS is needed to determine to which clade a C. auris isolate belongs
- Due to lack of testing capacity, WGS for C. auris previously took many months
 - Had to be done by special request to the CDC
- The Wisconsin State Laboratory of Hygiene (WSLH) now has the capability to sequence and analyze C. auris isolates. Routine testing includes:
 - The first isolate that initially diagnoses a patient with *C. auris*
 - All isolates that exhibit echinocandin resistance
 - Any isolates upon special request



Candida auris (C. auris) Data

2023: 279

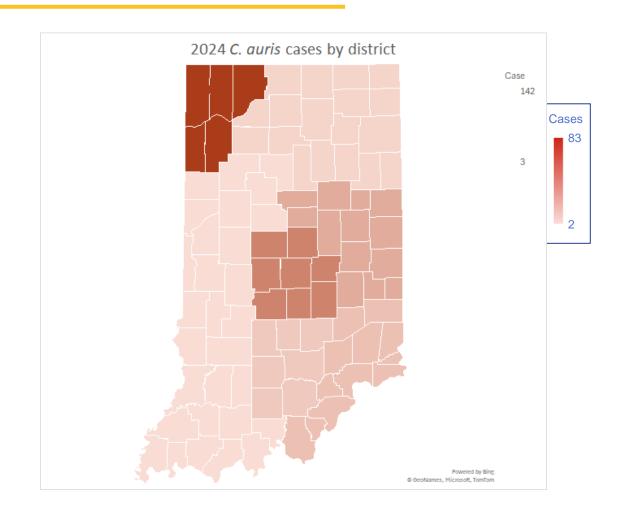
• Clinical: 105 (37.6%)

• Screening: 174 (62.3%)

2024: 362

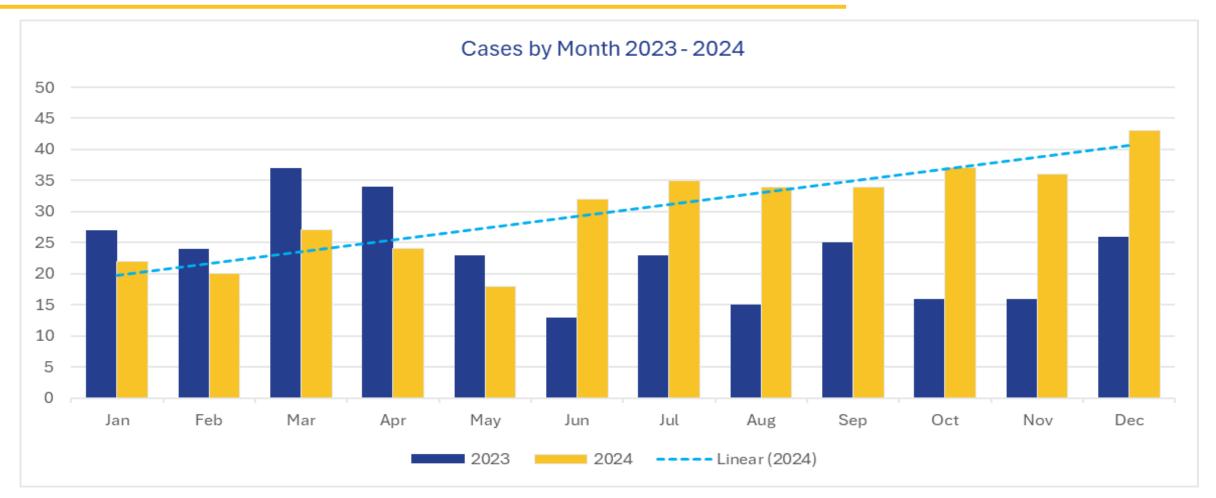
• Clinical: 166 (45.8%)

• Screening: 196 (54.1%)





Candida auris (C. auris) Trends

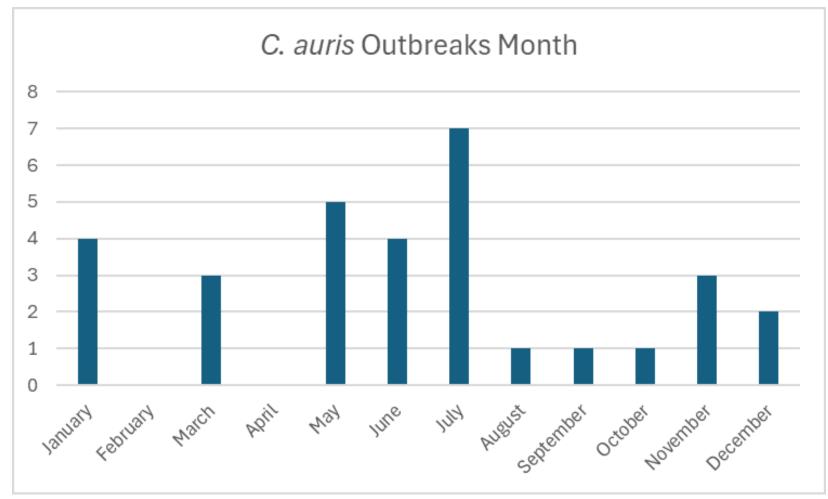




Candida auris Outbreaks

Total: 32

- Two responses due to echinocandin resistance
- One response due to organ donor involvement





United States *C. auris* Antifungal Resistance

Amphotericin B	echinocandins	Fluconazole
30 %	2 %	90 %

Please note that a finding of an elevated minimum inhibitory concentration (MIC) for an antifungal drug should not necessarily preclude its use. This is especially true if the use of other antifungal drugs for the patient has been ineffective.

Based on these MIC breakpoints, many isolates are resistant to multiple classes of drugs. Some U.S. *C. auris* isolates have been found to be resistant to all three classes of antifungal drugs. CDC has received reports of pan-resistance found in other countries as well.

^{*}For treatment information go to: https://www.cdc.gov/candida-auris/hcp/clinical-care/index.html



^{*}Source https://www.cdc.gov/candida-auris/hcp/laboratories/antifungal-susceptibility-testing.html

Indiana C. auris Antifungal Resistance thru 2023

YEAR	Micafungin	Anidulafungin	Caspofungin	Fluconazole	Amphotericin B
Breakpoint	<u>></u> 4	<u>></u> 4	<u>></u> 2	<u>></u> 32	MIC <u>></u> 2 ETEST <u>></u> 1.5
2020 N=34	0	0	0	74.47 %	0
2021 N=77	0	0	0	71.0 %	0
2022 N-145	0.69 %	0.69 %	0.7 %	70.83 %	2.78 %
2023 N=180	0.56 %	0.56 %	0.68 %	83.33 %	7.78 %



For surveillance purposes only

C. auris reporting form



Reporting Facility: _____

Address:

Recommendations

Candida auris Reporting Form

Please submit one report per patient per admission within one working day. Attach all laboratory results including antibiotic susceptibility test results. Fax form with the Confidential Report Form to Indiana Department of Health (317)-234-2812 or upload to NBS Morbidity Report.

Reporter Name: _____

Phone Number:

Patient name: DOB: Address:		NBS ID:			
		Phone:			
aboratory Information ***Attac	th all laboratory reports o	and antibiotic susceptibility testin	g results. ***		
Organism:		Collection date:			
Specimen site:		☐ Clinical culture ☐ Colonization culture			
Clinical information ***Attach al	l history and physical repor	ts available. ***			
Admission date:		Discharge date:			
From:		To:	To:		
□ Transfer form used upon ad	mission	☐ Transfer form used upon discharge			
Contact precautions start date:		Roommates: ☐ Yes ☐ No Dates:			
Were bleach cleaning products	T				
Invasive devices at time of	Invasive procedures	History of MDROs	Recent travel histo		
specimen collection	in past 6 months:	☐ MRSA ☐ VRSA	☐ Yes ☐ No		
☐ Central venus line		□ VRE □ ESBL	Where:		
☐ Mechanical vent		□ CRE			
☐ Tracheostomy		☐ Drug-resistant PA	When:		
☐ Urinary catheter		☐ Drug-resistant AB			
☐ Wound VAC					
Hospitalized in the last 3	Resident of a long-	Antibiotic use in past 30 days	Treatment		
months in acute care	term care facility?	Antibiotic use in past 30 days	Antibiotic:		
hospital or long-term care	□ Yes □ No	range of the	Paradione.		
facility?	Facility name:	Start date:	Start date:		
□ Yes □ No	,	Stop date:	Stop date:		
Facility name:					
Preexisting Conditions: No	ne 🗆 Unknown 🗆 Diabet	es mellitus Heart failure/CHF	Emphysema/COPD		
☐ Chronic renal insufficiency/o	thronic renal disease 🗆 O	besity Acute/Chronic respiratory	failure		
□ Peri/Hemi/Quadriplegia □	Wound/Ulcer/Abscess □	Chronic/Recurring UTI ☐ Cancer/I	Malignancy		
☐ Other:					





Candida auris Reporting Form

Please submit one report per patient per admission within one working day. Attach all laboratory results including antibiotic susceptibility test results. Fax form with the Confidential Report Form to Indiana Department of Health (317)-234-2812 or upload to NBS Morbidity Report.

We recommend placing the patient in enhanced barrier contact precautions (if applicable).

We recommend the use of an approved cleaning product from EPA List P.

We recommend flagging the patient chart in case the patient is readmitted to limit transmission.

We recommend utilizing a transfer form if patient is transferred.

If the patient had a roommate, we have a concern of transmission. Screening may be recommended.

If you would like additional resources, please visit the HAI/AR Website.

Pandrug-Resistant Organisms

- Pandrug-resistant Organisms
- Any isolates that shows intermediate or resistant to all antimicrobials tested

Condition Name in NBS:

Pandrugresistant Organisms

Reporting Timeframe:

 Within One Working Day



VISA/VRSA

Vancomycin-resistant *Staphylococcus aureus* (VRSA) and Vancomycin-intermediate *Staphylococcus aureus* (VISA) from any site

- Vancomycin Resistant: MIC ≥ 16
- Vancomycin Intermediate: MIC 4-8

Condition Name in NBS:

- VRSA
- S. aureus, vancomycin intermediate susc (VISA)

Reporting Timeframe:

 Within One Working Day



VISA, VRSA, and Pan-Drug Resistant Organisms

- VISA: All time- 3 Confirmed Cases
- VRSA: All time- 0 Confirmed Cases
- Pan-drug Resistant: 2023-12 confirmed cases. 2024- 9 confirmed cases



Infection Control Assessment and Response (ICAR)

- What if multiple individuals who test positive for an MDRO were at the same facility?
 - Confirmation that all individuals were at the facility would occur
- IDOH would work with the facility and suggest an onsite visit
 - A team from IDOH would meet with stakeholders at the facility
 - An ICAR tool would be completed by the infection prevention team at IDOH
 - Usually takes two to three hours:
 - Roundtable discussion with leadership, facility infection preventionist (IP), and housekeeping manager
 - Involves walking through the facility and observing staff performing their routine work where the cases reside



ICAR tool and IP scope

"ICAR tools are used to systematically assess a healthcare facility's infection prevention and control (IPC) practices and guide quality improvement activities (e.g., by addressing identified gaps)."- CDC

We use the Infection Control Assessment and Response (ICAR) tools to work with facilities across the state of Indiana to enhance infection prevention practices and improve outcomes for patients, staff, and residents

- Long-term Cares and Assisted Livings
- Long-term Care Acute Hospitals (LTACHs)
- Acute Care Hospitals (ACH)
- Department of Corrections (DOC)/Prisons' infirmaries
- Dental facilities
- Dialysis facilities



Outbreak ICAR: Targeted MDRO

Targeted MDRO Outbreak in a healthcare setting is identified.

- *C. auris* outbreak- two or more *Candida auris* cases with an epidemiological link.
- CPO outbreak- two or more CPOs of the same organism and mechanism occur in a 4-week period in patients who are epidemiologically linked or determined to be genetically related by laboratory testing.

IDOH offers an ICAR

- Assessment is educational.
 - Discussion with facility infection prevention personnel.
 - Review of facility policy and procedure.
 - Direct observations.



Benefits of an ICAR

- Potential identification of gaps in infection prevention practices that should be corrected
- Performance of the ICAR is collaborative and in no way punitive
- Increased awareness by:
 - Staff regarding the target organism
 - Staff regarding IP practices
 - Leadership regarding need for resources



The point prevalence survey (PPS)

- A key tool for public health
- Per CDC recommendations:
 - Should be used if evidence of transmission in a facility
 - Every patient on a given unit or floor where transmission is suspected should be screened
- IDOH facilitates supplies through our AR (antimicrobial resistance) lab network reference facility

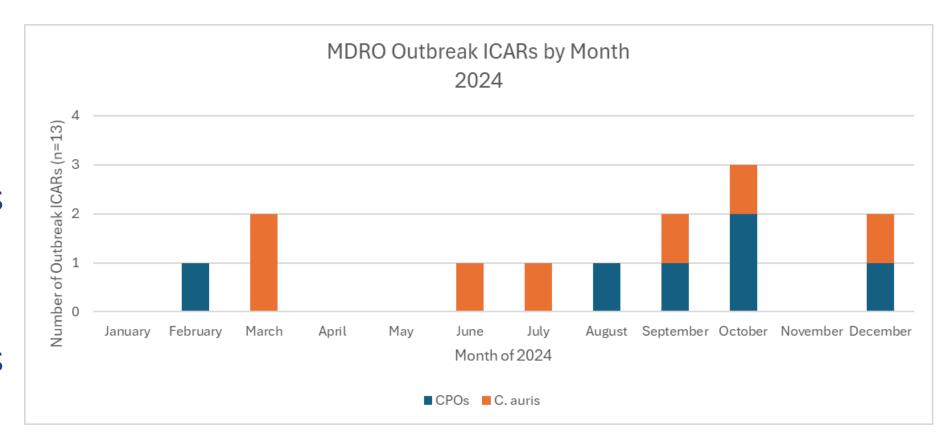


ICARs in 2024

• Total: 13

CPO: 6
 outbreak ICARs

• *C. auris*: 7 outbreak ICARs





Questions?

Caleb Cox, MPH
Senior MDRO Epidemiologist
calcox@health.in.gov

