



Candida auris: **An Enigmatic Fungal Pathogen of Global Concern**

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Disclosure



Rebecca is employed by Diversey—A Solenis Company. The company pays her expenses to attend this meeting & create educational content (salary). Diversey has had no input into this presentation from a commercial interest.

Objectives

01 Distinguish *Candida auris* (CA) from other *Candida* species

02 Discuss the epidemiology and current prevalence of CA

03 Understand why CA is an emerging threat

04 Review recent CA outbreaks

05 Summarize CA outbreak IPC recommendations



01

**Background:
Candida
species/auris**





Laboratory Basics

Fungi have two basic growth forms—molds and yeasts:

- Molds generally have a “fuzzy” appearance on lab media
- Yeasts appear more like a bacterial colony on media: soft, opaque cream color, 1 – 3 mm in diameter



Mold



Yeast

Candida species

Most common cause of fungal infections worldwide:

- *C. albicans* – most common Candida pathogen
- *C. parapsilosis*
- *C. glabrata*
- *C. tropicalis*
- *C. guilliermondii*
- *C. dubliniensis*

Generally called:
non-albicans Candida,
or Candida species- not
C. albicans





Candida species as a cause of HAIs

Top 15 HAI Pathogens Reported to NHSN, Adults, 2018-2021

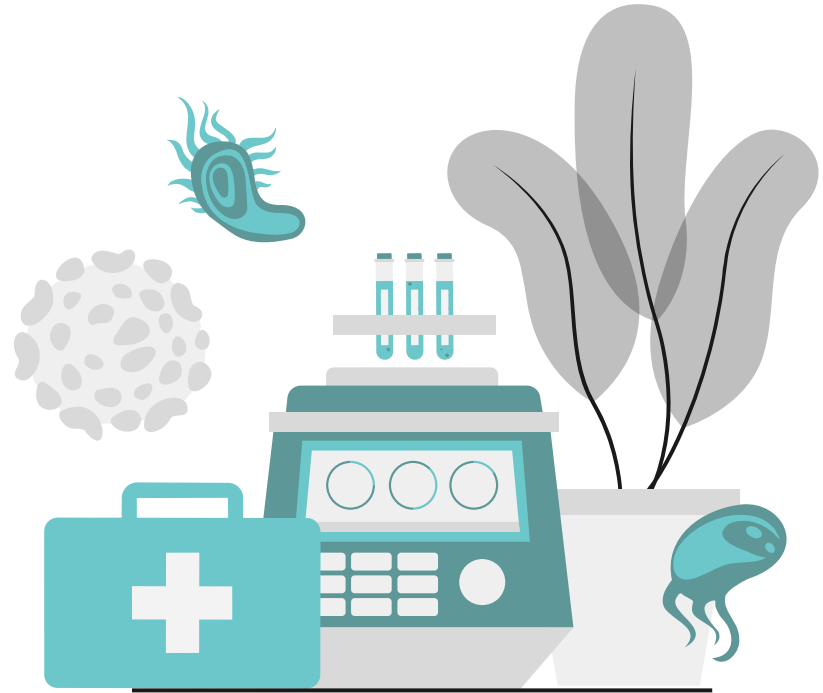
When analyzed to the genus level, Candida species ranked #6 (7.5%)



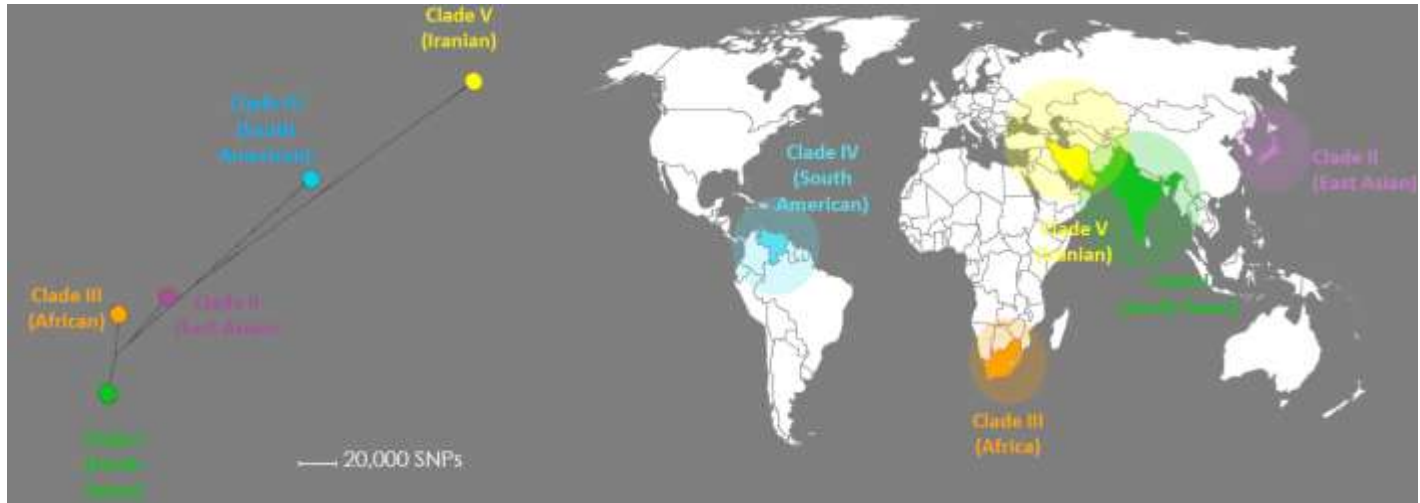
Pathogen	# Pathogens	% Pathogens	Rank
<i>Escherichia coli</i>	73,556	16.2	1
<i>Staphylococcus aureus</i>	51,131	11.3	2
<i>Enterococcus faecalis</i> ²	39,129	8.6	3
Select <i>Klebsiella</i> spp.	38,496	8.5	4
<i>Pseudomonas aeruginosa</i>	36,004	7.9	5
Coagulase-negative staphylococci	32,276	7.1	6
<i>Enterobacter</i> spp.	18,431	4.1	7
<i>Enterococcus faecium</i> ²	16,904	3.7	8
<i>Candida albicans</i> ²	16,458	3.6	9
<i>Proteus</i> spp.	13,953	3.1	10
<i>Bacteroides</i> spp.	11,602	2.6	11
Viridans group streptococci	9,962	2.2	12
Other <i>Candida</i> spp. ²	9,803	2.2	13
Other <i>Enterococcus</i> spp. ²	9,091	2.0	14
<i>Candida glabrata</i> ²	7,622	1.7	15
Other pathogen	68,522	15.1	
Total	452,940	100.0	

CA Background

- First described in 2009 in an ear isolate in Tokyo, Japan but the earliest known case was retrospectively identified in South Korea and dated back to 1996
- 5 distinct clades
 - Clade: A taxonomic group of organisms classified together on the basis of homologous features traced to a common ancestor – (dictionary.com)



CA Clades & Worldwide Distribution



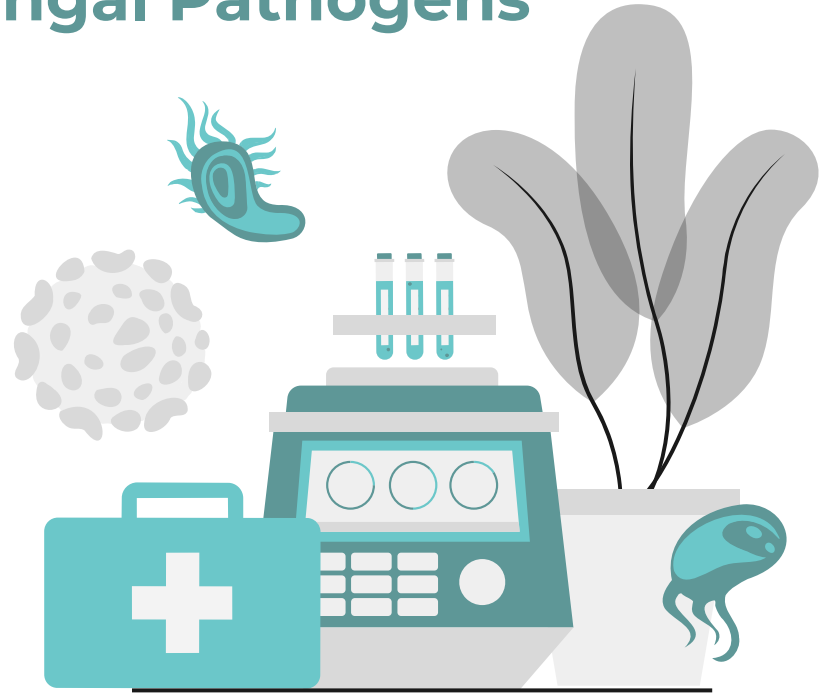
Around the world, CA has been reported in at least 50 countries on 6 continents



Global Risk of Fungal Pathogens

WHO: CA is under-recognized global threat compounded by rapid spread, emergence of antifungal resistance and limited access to quality diagnostics and treatment.

Critical group	
	<i>Cryptococcus neoformans</i>
	<i>Candida auris</i>
	<i>Aspergillus fumigatus</i>
	<i>Candida albicans</i>



WHO fungal priority pathogens list to guide research, development and public health action: 2022

COVID-19 Impacts on 18 Antimicrobial-Resistant Bacteria and Fungi Threat Estimates

The following table summarizes the latest national death and infection estimates for 18 antimicrobial-resistant bacteria and fungi. The pathogens are listed in three categories—urgent, serious, and concerning—based on level of concern to human health identified in 2019.

	Resistant Pathogen	2017 Threat Estimate	2018 Threat Estimate	2019 Threat Estimate	2017-2019 Change	2020 Threat Estimate and 2019-2020 Change
URGENT	Carbapenem-resistant <i>Acinetobacter</i>	8,500 cases 700 deaths	6,300 cases 500 deaths	6,000 cases 500 deaths	Stable*	7,500 cases 700 deaths Overall: 35% Increase* Hospital-onset: 78% Increase*
	Antifungal-resistant <i>Candida auris</i>	171 clinical cases†	329 clinical cases	466 clinical cases	Increase	754 cases Overall: 60% Increase
	<i>Clostridioides difficile</i>	237,000 infections 12,800 deaths	221,200 infections 12,600 deaths	202,600 infections 11,500 deaths	Decrease	Data delayed due to COVID-19 pandemic
	Carbapenem-resistant Enterobacterales	13,100 cases 1,100 deaths	10,300 cases 900 deaths	11,900 cases 1,000 deaths	Decrease*	12,700 cases 1,100 deaths Overall: Stable* Hospital-onset: 35% Increase*
	Drug-resistant <i>Neisseria gonorrhoeae</i>	550,000 infections	804,000 infections	942,000 infections	Increase	Data unavailable due to COVID-19 pandemic
SERIOUS	Drug-resistant <i>Campylobacter</i>	448,400 infections 70 deaths	630,810 infections	725,210 infections	Increase	Data delayed due to COVID-19 pandemic 26% of infections were resistant, a 10% decrease
	Antifungal-resistant <i>Candida</i>	34,800 cases 1,700 deaths	27,000 cases 1,300 deaths	26,600 cases 1,300 deaths	Decrease*	28,100 cases 1,400 deaths Overall: 12% Increase* Hospital-onset: 26% Increase*
	ESBL-producing Enterobacterales	197,400 cases 9,100 deaths	174,100 cases 8,100 deaths	194,400 cases 9,000 deaths	Increase*	197,500 cases 9,300 deaths Overall: 10% Increase* Hospital-onset: 32% Increase*
	Vancomycin-resistant Enterococcus	54,500 cases 5,400 deaths	46,800 cases 4,700 deaths	47,000 cases 4,700 deaths	Stable*	50,300 cases 5,000 deaths Overall: 16% Increase* Hospital-onset: 14% Increase*

CDC, 2022 Special Report: Covid-19, US Impact on Antimicrobial Resistance

60%
overall CA
Increase
from 2019
to 2020!

02

Epidemiology & Prevalence



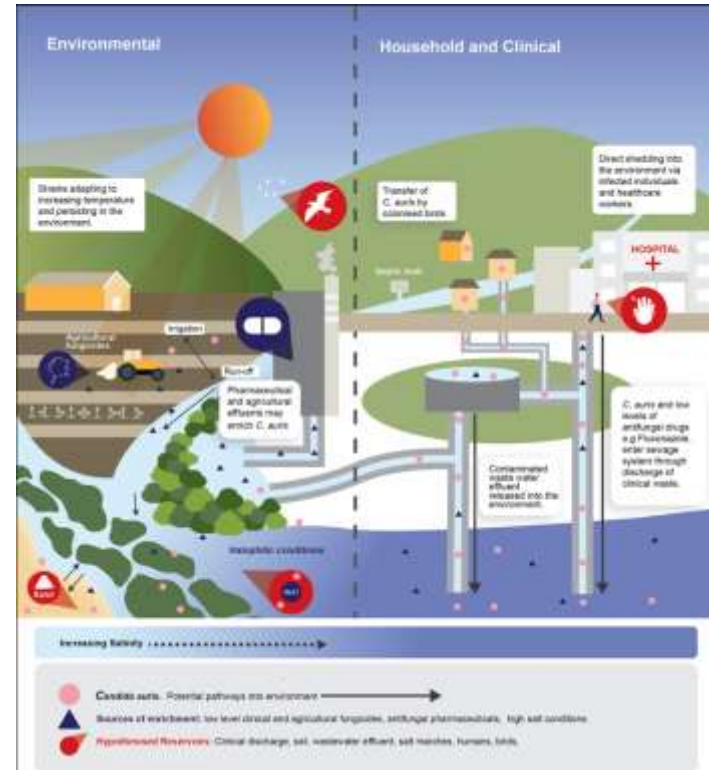


Define “emerging” infectious diseases/pathogen

The National Institute of Allergy and Infectious Diseases (NIAID) defines “emerging infectious diseases/pathogens” as those “that have newly appeared in a population or have existed but are rapidly increasing in incidence or geographic range.”

Where Did It Come From?

- Prior to being recognized as a human pathogen, CA likely existed as a plant saprophyte in wetlands
- CA has the unique ability to grow in:
 - Higher temperatures and levels of humidity
 - High salinity
- First environmental isolates came from a sandy beach and a salt marsh wetland in the Andaman Islands, India and from an estuary in Colombia
- Indicates that CA existed as an environmental fungus

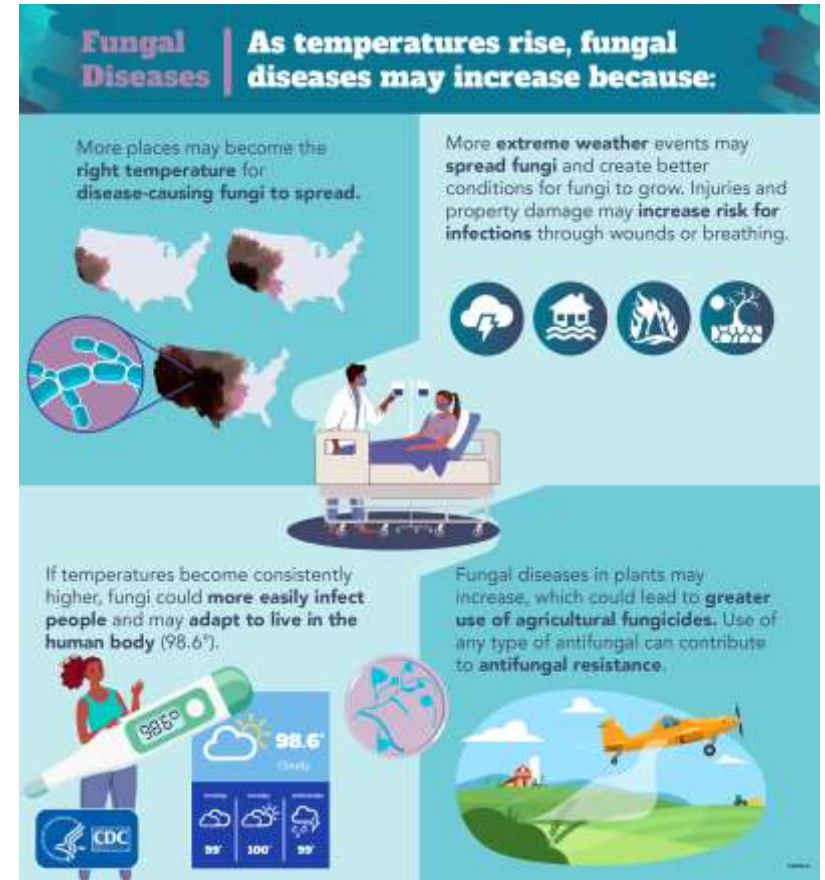


Global Warming Hypothesis

Increase in global warming led to simultaneous emergence of thermo-tolerant CA in different geographical locations

Other factors include:

- Global human migration
- Poor hygiene
- High population densities
- Use of fungicides in agriculture (contributes to antifungal resistance)



Colonization

Commonly reported sites of colonization include:

- Skin, especially groin and axilla areas
- Mucosal surfaces of the gastrointestinal and genitourinary tract
- Respiratory tract (oropharynx, nose)
- Ear (named for the latin word for ear “auris”)
- Tips of central venous catheters

In areas of high CA prevalence, colonization rates of 2.5%-33.9% have been reported

* Contributes to its ability to spread easily in health care settings



Infections



Individuals colonized or infected with CA frequently have comorbidities such as:

- Diabetes
- Bloodstream infection and multi-organ failure
- Pulmonary diseases/pneumonia
- Chronic or acute kidney failure
- Immunosuppressive conditions
- Solid tumour/malignancies
- Cardiovascular/hypertension
- Chronic otitis/media
- Gastrointestinal disease

Nearly 10% of CA-colonized patients develop infection, particularly those in the ICU setting (on mechanical ventilation and with invasive devices)

Overall crude mortality rate of 30-60%



Infections



Non-invasive CA infections include:

- Respiratory tract
- Urinary tract
- Wound infections
- Skin abscesses (often related to catheters)
- Otitis externa

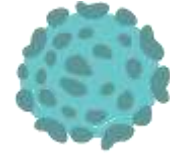
Invasive CA infections include:

- Bloodstream infection
- Pericarditis
- Myocarditis
- Meningitis
- Osteomyelitis

* CA is thermotolerant- grows optimally at 98°F, but it can remain viable at 107°F. This gives CA the ability to cause invasive infections and tolerate fever

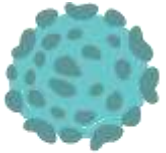


Infections



Epidemiology of Candida infections is changing:

- Candida albicans is the most common cause of candidemia and invasive candidiasis
- >50% of all infections are now caused by non-albicans species
- Likely due to the increasing use of antifungal drugs for prophylaxis and treatment which has resulted in the selection of Candida species with reduced susceptibility to antifungal drugs



*CA has been linked to been link to major outbreaks of invasive infection in healthcare facilities around the world

CA Prevalence in Europe



RED- CA outbreak countries with inter-facility spreading or endemicity (Spain, Italy, Greece, UK)

YELLOW- Sporadic outbreaks with or with none or limited inter-facility spreading

BLUE- Sporadic locally acquired cases or an unknown or imported origin



Prevalence in the US



2017

2017



Number of *C. auris* clinical cases through December 31, 2022

In 2017, there were 173 clinical cases and 272 screening cases

- 0 clinical cases and at least 1 screening case
- 1 to 10
- 11 to 50
- 51 to 100
- 101 to 500
- 501 to 1000
- 1001 or more

Recent 12 months



Number of *C. auris* clinical cases through December 31, 2022

In the most recent 12 months, there were 2,377 clinical cases and 5,754 screening cases (January 2022 - December 2022)

- 0 clinical cases and at least 1 screening case
- 1 to 10
- 11 to 30
- 31 to 100
- 101 to 500
- 501 to 1000
- 1001 or more

Prevalence in the US

Cases:

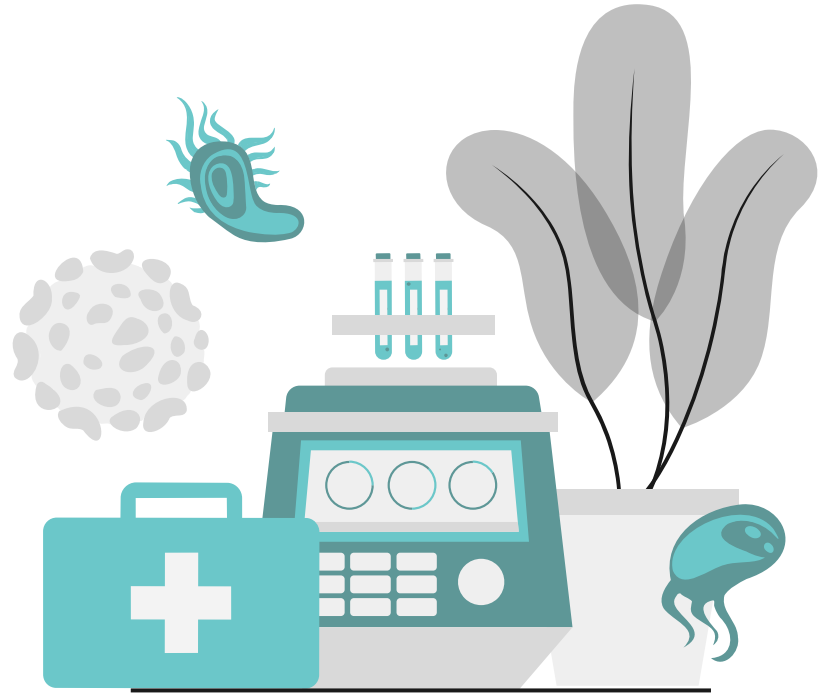
2018-2019: 44% increase

2019-2020: 59% increase

2020-2021: 95% increase

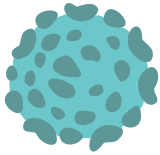
In 2022, Nevada was the most affected state followed by California, Florida and New York

CA has been detected in over 50% of American states



Candida auris is a nationally notifiable condition and is reportable in many states. Laboratories that identify cases of CA should report cases immediately to the state or local health department.

CA is on North Dakota's Mandatory Reportable Infectious Conditions list!



NORTH Dakota Health & Human Services		Mandatory Reportable Infectious Conditions	
Be. Laboratory		If highlighted red, report immediately: 800-472-2180 or 701-328-2378	
Report all other conditions within one business day			
Acute Flaccid Myelitis	Hepatitis D	Respiratory Panel Results ³	
Alpha-gal Syndrome	Hepatitis E	Respiratory Syncytial Virus ³	
Anaplasmosis	HIV/AIDS infection ²	• Pediatric deaths	
Anthrax ♦♦	Influenza ³	Rocky Mountain spotted fever	
Arboviral infection (other)	• Pediatric deaths ♦	Rubella ♦	
Babesiosis	• Suspect novel, PCR influenza A unsubtypeable ♦	Salmonellosis ♦	
Botulism ♦♦	Jamestown Canyon virus disease	SARS-CoV-2 ¹	
Brucellosis ♦♦	Laboratory incidents with possible release of category A agents or novel influenza virus ♦	• Pediatric deaths	
Campylobacteriosis	La Crosse encephalitis	Scabies outbreaks in institutions	
Candida auris ♦	Legionellosis	Shigellosis ♦	
Carbapenem-resistant organisms	Leptospirosis	Smallpox ♦♦	
• Enterobacteriales ♦		Staphylococcus aureus	
• Pseudomonas aeruginosa ♦		• Vancomycin-resistant and intermediate resistant (VRSA and VISA) – any site ♦	
• Acinetobacter baumannii ♦			

<https://www.cdc.gov/fungal/fungal-disease-reporting-table.html>

Case Definitions 2023

National Notifiable Diseases Surveillance System (NNDSS)



Screening Case

Detection of *C. auris* in a specimen from a swab obtained for the purpose of colonization screening using either culture or validated culture-independent test (e.g., nucleic acid amplification test [NAAT])



Clinical Case

Detection of *C. auris* in a clinical specimen obtained during the normal course of care for diagnostic or treatment purposes using either culture or a validated culture-independent test (e.g., NAAT)

03

Why is *Candida auris* an emerging threat?



Why Are We Concerned?



Morbidity and Mortality



**Potential to Cause
Outbreaks in
Healthcare Settings**



**Laboratory
Identification**



Disinfection Challenges



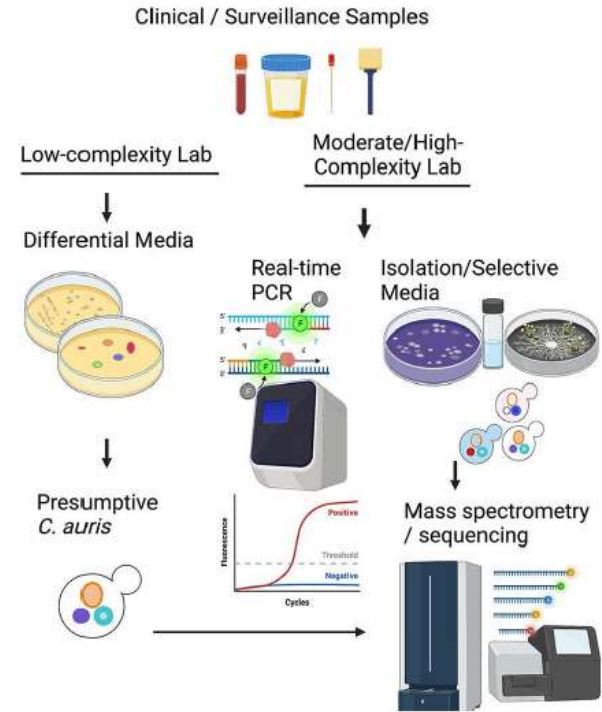
**Resistance to
Antifungals**

Laboratory Identification

Cultures based methods take several days to result and lack sensitivity (50% of invasive candida infections are culture negative)

In clinical laboratories that rely on biochemical-based testing for yeast identification, up to 90% of CA isolates can be misidentified as other *Candida* species or other yeasts

Matrix-Assisted Laser Desorption Ionization-Time of Flight Mass Spectrometry (MALDI-TOF MS) systems are widely used for the rapid and accurate identification of bacteria, mycobacteria and certain fungal pathogens in the clinical microbiology laboratory-
*had to add isolates from the 4 major clades to their databases



Laboratory Identification

GOLD STANDARD

DNA sequence analysis- this determines the exact sequence of nucleotides or bases in a DNA molecule (D1/D2 and ITS sequencing)

Other methods such as whole genome sequencing (WGS) can further delineate local clusters to inform the source of transmission

Molecular sequencing of ribosomal DNA loci further enables clade differentiation



<https://www.pacb.com/blog/the-evolution-of-dna-sequencing-tools/>

Antibiotic Resistance Laboratory Network (ARLabnetwork)



Purpose is to close the gap between local lab capabilities and the data needed to combat antimicrobial resistance by providing:

- Comprehensive lab capacity and infrastructure for antimicrobial-resistant pathogens
- Cutting-edge technology, like DNA sequencing
- Data to drive response and prevent infections





Anti-Fungal Susceptibility Worldwide



3 main classes of FDA approved antifungals:
Azoles, Polyenes, Echinocandins

Echinocandins are the first-line therapy for invasive Candida infections and most CA infections.

*CA is often resistant to at least one class of antifungals

Azoles- 87-100% resistance to fluconazole, susceptibility to other azoles vary
Polyene- moderate resistance to Amphotericin B (8-35%)

4% of CA isolates are **pan-resistant** (reported in U.S, India, South America)

Resistance rates in different countries and different health care settings vary considerably and are mainly due to different clade distributions in different settings





Anti-Fungal Susceptibility Worldwide



Features	Clade I (South Asian)	Clade II (East Asian)	Clade III (South African)	Clade IV (South American)	Clade V (Iranian)
Antifungal Resistance Profile	Resistant to fluconazole, echinocandins, amphotericin B Pan-resistance identified in some strains ³⁰	Usually susceptible to antifungal drugs ²⁴	Resistant to fluconazole Cross-resistant to echinocandins, amphotericin B Pan-resistance identified in some strains ³⁰	Resistant to fluconazole Cross-resistant to echinocandins, amphotericin B Pan-resistance identified in some strains ³⁰ Note: the first isolates in Ontario were all clade IV and all were pan-susceptible	Resistant to fluconazole ³¹





Anti-Fungal Susceptibility United States



90% of CA isolates resistant to fluconazole

30% of CA isolates resistant to amphotericin B

2-5% of CA isolates resistant to echinocandins

*In the New York-New Jersey area where 55% of all U.S. isolates occur:
99.8% fluconazole-resistant
50% amphotericin B-resistant

AR Lab Network reported pan-resistant strains:
4 cases before 2020
6 cases in 2020
7 cases in 2021
Identified in Texas, NYC, District of Columbia



Notes from the Field

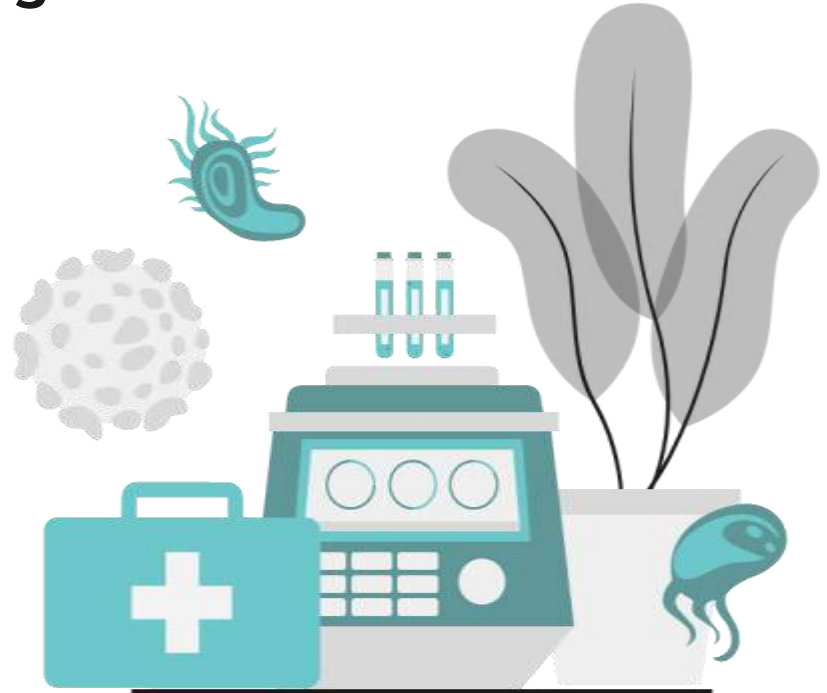
Transmission of Pan-Resistant and Echinocandin-Resistant *Candida auris* in Health Care Facilities — Texas and the District of Columbia, January–April 2021

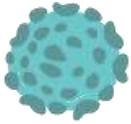
Meghan Lyman, MD¹; Kaitlin Forsberg, MPH¹; Jacqueline Reuben, MHS²; Thi Dang, MPH³; Rebecca Free, MD¹; Emma E. Seagle, MPH¹; D. Joseph Sexton, PhD¹; Elizabeth Soda, MD⁴; Heather Jones, DNP⁴; Daryl Hawkins, MSN²; Adonna Anderson, MSN²; Julie Bassett, MPH³; Shawn R. Lockhart, PhD¹; Erynnaya Merengwa, MD, DrPH³; Preetha Iyengar, MD²; Brendan R. Jackson, MD¹; Tom Chiller, MD¹

Public Health Ontario 2023

Factors Affecting Spread in Health Care Settings

- Commonly reported sites of colonization include the skin (especially the groin and axilla areas)- Continuous carriage for more than a year after initial isolation of *C. auris* has been documented
- CA remains viable on surfaces for a prolonged period of time and is shown to survive for up to 4 weeks despite surface decontamination and remain viable on inanimate surfaces for several months
- CA can withstand many common hospital disinfectants





Cleaning and Disinfection



Quaternary ammonium chemistries (QACs) don't work, but are the most widely used in healthcare by EVS (Han 2021).

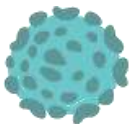
Not all “fungicidal” disinfectants are effective against CA- check!

Important to adhere to manufacturers instructions for dilution and contact time to ensure efficacy

Reference “List P” for products with EPA approved claims against CA

The screenshot shows the EPA website header with the logo and navigation menu. Below the header, the page title is 'List P: Antimicrobial Products Registered with EPA for Claims Against Candida Auris'. Underneath, there is a section titled 'On this page:' followed by a list of links: 'Products on List P', 'How to use List P products effectively', 'How to check if a product is on List P', and 'Additional Resources'.

<https://www.epa.gov/pesticide-registration/list-p-antimicrobial-products-registered-epa-claims-against-candida-auris>



Cleaning and Disinfection

Patient Care Environment and Equipment



Perform thorough daily and terminal cleaning and disinfection of the patient's room and other areas they received care (i.e. physiotherapy, imaging etc.)

Dedicate equipment and supplies (e.g., stethoscopes, glucometers, temperature probes, blood pressure cuffs), and other shared equipment (e.g., ventilators, Hoyer lifts, physical therapy equipment) to CA positive patients

Clean and disinfect all supplies and equipment immediately after use and dispose of unused, disposable supplies after patient discharge.

*** A List P product should be used by BOTH:**

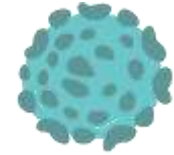
Environmental Services



Clinical Staff



Cleaning and Disinfection

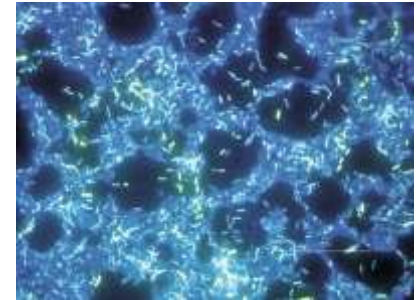


Most CA strains have shown the ability to form biofilms:

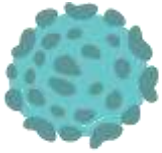
Biofilms develop when microorganisms adhere to a surface embedded in an extracellular polymeric substance (EPS) matrix. The matrix contains polysaccharides, proteins, lipids, enzymes, extracellular DNA and water.

Biofilms prolong the survival of microorganisms and renders them tolerant to normal hospital cleaning and disinfection procedures.

Biofilms are traditionally associated with wet or damp surfaces (such as drains) but they can also form on inanimate dry surfaces.



*Currently there is no standard method for testing efficacy of detergents and disinfectants against biofilm formed on dry surfaces.



Cleaning and Disinfection

Biofilms can develop on or within indwelling medical devices such as:

- central venous catheters and needleless connectors
- endotracheal tubes
- intrauterine devices
- mechanical heart valves and pacemakers
- dialysis catheters
- prosthetic joints
- urinary catheters

Compared to other *Candida* species, CA can persist in a viable form on dried or moist surfaces for several weeks longer than other *Candida* species





04

What Can We Learn From Recent Outbreaks?

CA Cluster in Paediatric Patients in Acute Care Hospital in Nevada



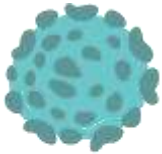
ACH had 31 adult cases and then identified 3 pediatric cases between December 2021 and April 2022

Pediatric cases:

- Under 6 months
- Born at ACH
- Severely ill and had invasive BSIs
- Had overlapping stays in the unit in neighboring rooms

CDC review identified:

- Cleaning and disinfection (C&D) policy of “who uses it cleans it” but staff could not state this was the expectation
- EVS staff not always moving from clean to dirty, potential contamination of the EVS cart and misuse of PPE
- Echocardiogram (which was wheeled between patient rooms) performed on all 3 pediatric cases and 66.7% of adult cases BEFORE testing positive



CDC Recommendations



- Ensure use of a disinfectant with claims against CA
- Assign clear C&D responsibility
- Train staff on how to perform adequate C&D
- Implement C&D audit and feedback system
- Improve hand hygiene compliance
- Refresher training on correct PPE use
- Education on best practices for C&D workflow
- Ensure all Candida isolates are speciated to identify CA
- Point prevalence surveys on all units with suspected transmission
- Targeted admission screening



CA Outbreak in Northern Italy 2019-2021



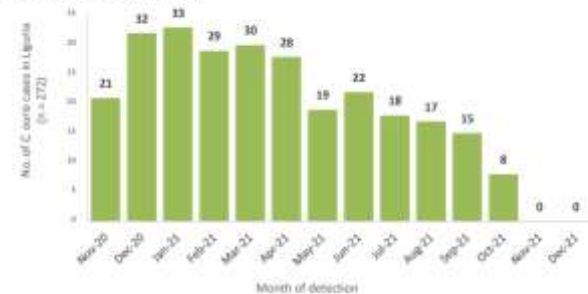
First CA case in ACH was identified in July 2019 and sporadic cases continued to occur with a total of 277 across 8 healthcare facilities as of this report

CA was detected in a Covid-19 ICU in February 2020 and cases continued to increase to October 2021

4 healthcare facilities received patients with CA from this ACH

Risk of spread within Italy was deemed to be “HIGH”

Figure 1. Epidemiological curve of *C. auris* cases in Liguria, northern Italy, from November 2020 to December 2021 (n = 272*)



European Centre for Disease Prevention and Control
Rapid Risk Assessment February 2022



ECDC Recommendations



Good standard infection prevention and control including environmental C&D, adequate C&D of reusable medical equipment, adequate microbiology laboratory capacity, sufficient capacity for isolation of patients in healthcare facilities

In addition, early, robust action when a case is identified:

- Prompt communication to IP and clinical team
- Detailed investigation of all cases- case review, contact tracing, consider point prevalence based on risk assessment
- Contact precautions and flagging of patients in case of readmission
- Single use equipment if possible or dedicated equipment for length of stay
- Use of a disinfectant effective against CA
- Raising awareness by providing education to all healthcare personnel
- Regular active surveillance cultures on wards



CA Outbreak in a Covid-19 Specialty Care Unit- Florida July-August 2020



Hospital's Covid-19 unit consisted of 5 wings on 4 floors

In July 2020, the Florida Department of Health reported 4 Covid-19 positive patients with CA- 3 with bloodstream infections and 1 with urinary tract infection

Amongst 67 patients screened during point prevalence surveys, 35 (52%) were CA positive and 6 (17%) subsequently had positive clinical samples

Joint investigation by Florida Department of Health and CDC revealed:

- Staff wearing multiple layers of PPE- First layer worn at all times (extended use), second layer donned on entry to patient rooms
- Clean equipment handled using inner layer of PPE including gloves which may be contaminated
- Multiple opportunities for contamination of the inner layer of PPE during doffing and through direct contact with the patient environment
- Shared patient equipment not always disinfected between uses
- Missed opportunities for hand hygiene



Florida Department of Health and CDC Recommendations



Multiple layers of PPE is not recommended. Use one layer of PPE at a time for the care of Covid-19 patients

Shared equipment must be appropriately cleaned and disinfected between patients

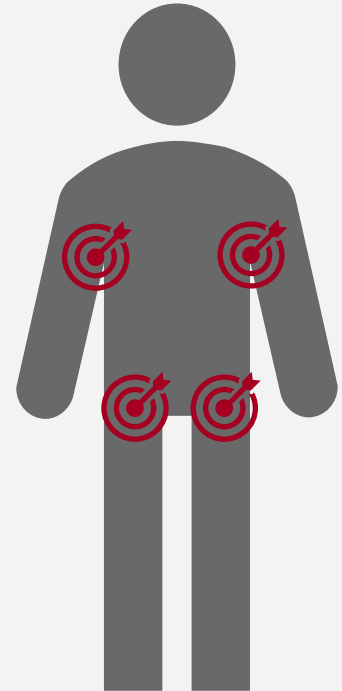
Clean patient equipment should not be handled wearing potentially contaminated PPE

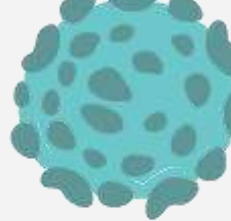
Implement strategies to increase adherence to appropriate hand hygiene



Positive Correlation between *Candida auris* Skin Colonization Burden and Environmental Contamination in Ventilator-Capable Skilled Nursing Facility (vSNF) in Chicago, Illinois (Sexton et al 2021)

- 70-bed facility in Chicago Illinois
 - First CA case was identified by point prevalence in March 2017
 - **In 18 months (Sept 2018), CA colonization climbed to 71%!**
- Study sampled bilateral axillary/inguinal swabs on all residents





Study Findings vSNF Chicago: *Candida auris* Positive Environmental Cultures



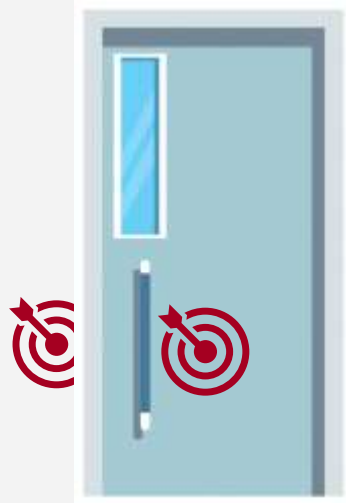
Bedrails

Left

Right

81%

78%



Door handles

Inner

Outer

58%

25%



Windowsills

75%

Resident & Environmental CA Heatmap!

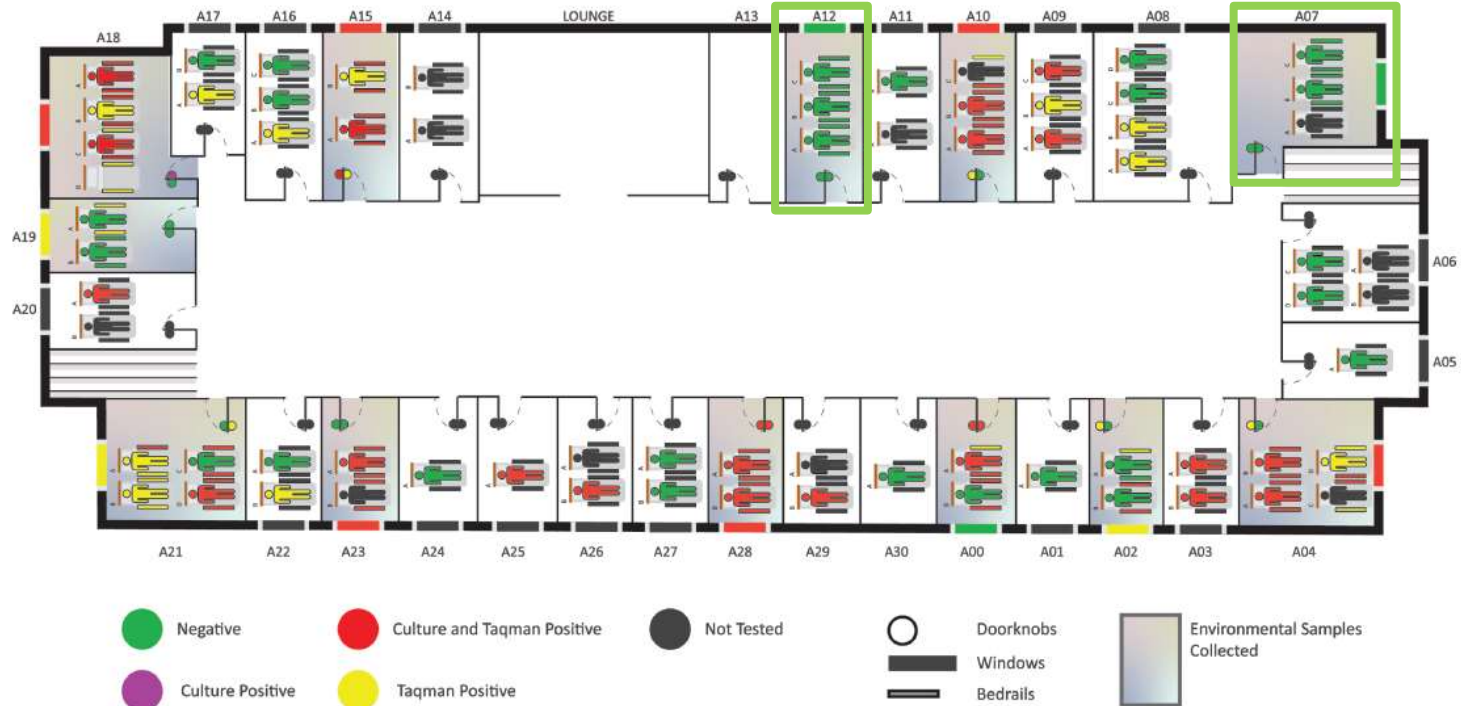


Figure 1. Facility map with culture-based and qPCR results for residents and associated environmental surfaces. The specific organization of beds within a room may differ from the image.

Study Findings vSNF Chicago



Colonized residents can have **high CA burden on their skin**, which was positively related with **contamination** of their surrounding healthcare **environment**.

3 patients who screened negative had bedrails that tested positive:

- 1 patient was previously positive for CA
- The other 2 patients were in rooms which were previously occupied by a CA positive patient 1-2 months before the study

These findings underscore the importance of:

- Hand hygiene
- Transmission-based precautions
- **Frequent environmental disinfection with EPA List P disinfectants**



05

Summary of Outbreak Recommendations

Summary of Outbreak Recommendations

Refer to CDC Infection Prevention and Control for *Candida auris*
<https://www.cdc.gov/fungal/candida-auris/c-auris-infection-control.html>

HEALTHCARE ENVIRONMENT:

- Assign clear C&D responsibility
- Train staff on how to effectively perform C&D including best practices for workflow- who cleans what and how?
- Implement C&D monitoring system (ATP, fluorescent marking, direct observation) with timely feedback
- Use a disinfectant effective against CA (List P)

PATIENT MANAGEMENT

- Single use equipment if possible or dedicated equipment for length of stay
- Manage patient on Contact precautions and flag chart in case of readmission



Summary of Outbreak Recommendations

STAFF

- Raise awareness by providing education to all healthcare personnel
- Prompt communication of case to IP and clinical team
- Implement strategies to increase adherence to appropriate hand hygiene, PPE use and C&D of multi-use equipment

Communication of Interfacility Transfers

Healthcare facilities should be informed if an incoming patient has ever:

1. Tested positive for CA, with or without symptoms.
2. Was exposed to another patient with C. auris.
3. Was in a facility where an outbreak was occurring

Inter-facility Infection Control Transfer Form

This form must be filled out for transfer to accepting facility with information communicated prior to or with transfer.
Please attach copies of latest culture reports with susceptibilities if available.

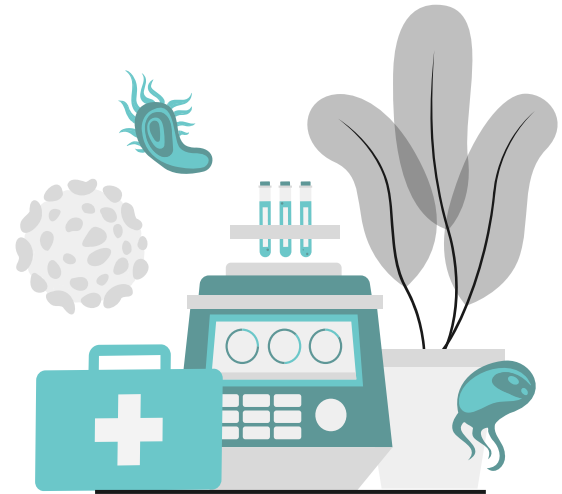
Sending Healthcare Facility:			
Patient/Resident Last Name	First Name	Date of Birth	Medical Record Number
Name/Address of Sending Facility		Sending Unit	Sending Facility Phone
Sending Facility Contacts	Contact Name	Phone	E-mail
Transferring RN/Unit			
Transferring physician			
Case Manager/Admin/SW			
Infection Preventionist			
Does the person* currently have an infection, colonization OR a history of positive culture of a multidrug-resistant organism (MDRO) or other potentially transmissible infectious organism?		Colonization or History (Check if YES)	Active Infection on Treatment (Check if YES)
Methicillin-resistant Staphylococcus aureus (MRSA)		<input type="checkbox"/> Yes	<input type="checkbox"/> Yes
Vancomycin-resistant Enterococcus (VRE)		<input type="checkbox"/> Yes	<input type="checkbox"/> Yes
Clostridioides difficile		<input type="checkbox"/> Yes	<input type="checkbox"/> Yes
Acinetobacter, multidrug-resistant		<input type="checkbox"/> Yes	<input type="checkbox"/> Yes
Enterobacteriaceae (e.g., E. coli, Klebsiella, Proteus) producing Extended Spectrum Beta-Lactamase (ESBL)		<input type="checkbox"/> Yes	<input type="checkbox"/> Yes
Carbapenem-resistant Enterobacteriaceae (CRE)		<input type="checkbox"/> Yes	<input type="checkbox"/> Yes
Pseudomonas aeruginosa, multidrug-resistant		<input type="checkbox"/> Yes	<input type="checkbox"/> Yes
Candida auris		<input type="checkbox"/> Yes	<input type="checkbox"/> Yes
Other, specify (e.g., list, scabies, norovirus, influenza):		<input type="checkbox"/> Yes	<input type="checkbox"/> Yes
Does the person* currently have any of the following? (Check here <input type="checkbox"/> if none apply)			
<input type="checkbox"/> Cough or requires suctioning	<input type="checkbox"/> Central line/PICC (Approx. date inserted: <input type="text"/>)		
<input type="checkbox"/> Diarrhea	<input type="checkbox"/> Hemodialysis catheter		
<input type="checkbox"/> Vomiting	<input type="checkbox"/> Urinary catheter (Approx. date inserted: <input type="text"/>)		
<input type="checkbox"/> Incontinence of urine or stool	<input type="checkbox"/> Suprapubic catheter		
<input type="checkbox"/> Open wounds or wounds requiring dressing change	<input type="checkbox"/> Percutaneous gastrostomy tube		
<input type="checkbox"/> Drainage (source): <input type="text"/>	<input type="checkbox"/> Tracheostomy		

<https://www.cdc.gov/hai/pdfs/toolkits/Interfacility-IC-Transfer-Form-508.pdf>

Summary of Recommendations

SURVEILLANCE AND TESTING:

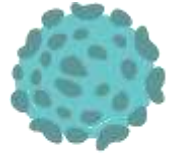
- Ensure all Candida isolates are speciated to identify CA.
- Perform point prevalence surveys on all units with suspected transmission.
- Consider targeted admission screening based on local epidemiology.
- Take early, robust action when a case is identified.
- Detailed investigation of all cases- case review, contact tracing, consider point prevalence based on risk assessment



In Summary

The rise and geographic spread of CA is concerning because of the:

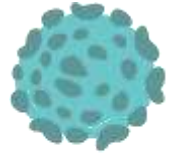
- Extent of resistance to antifungal treatments
- Ability to withstand common healthcare disinfectants
- High mortality rates of invasive infections
- Ability to cause prolonged outbreaks in health care settings
- Difficulty to identify in an accurate and timely manner in the laboratory



In Summary

What's needed:

- Surveillance & **early detection** of cases to mitigate transmission (**increased screening!**)
- Increased **laboratory capacity** for identification, cluster analysis, & antifungal susceptibility testing
- **New antifungal** agents
- Improved **disinfectants** with achievable contact times on **EPA's List P**, with **increased cleaning & disinfection during outbreaks**
- Strict adherence to infection prevention & control policies and procedures



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Thanks!

Questions?

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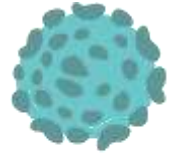
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Public Health Ontario Focus on Candida Auris, June 2023

<file:///C:/Users/dv117988/OneDrive%20-%20Solenis%20LLC/Documents/Candida%20auris/candida-auris%20PHO%202023.pdf>

