

# *Candida auris*

## An Emerging MDR Fungal Threat in Healthcare

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# Learning Objectives

## 1. Describe the history

Explain how and when *C. auris* emerged globally and why it is considered a novel pathogen

## 2. Understand the threat

Identify why *C. auris* is uniquely dangerous in healthcare settings compared to other *Candida* species

## 3. Recognize patient impact

Describe mortality rates, risk factors, and the patient populations most affected

## 4. Navigate antifungal resistance

Understand resistance patterns and the rationale for first-line treatment choices

## 5. Apply treatment strategies

Select appropriate antifungal therapy and understand when to consult infectious disease

## 6. Implement infection control

Apply CDC-recommended measures to prevent nosocomial transmission in your facility

# What Is Candida auris?

## A novel yeast pathogen

*C. auris* is a species of ascomycetous fungus that grows as a budding yeast. It rarely forms hyphae or pseudohyphae. Named from the Latin *auris* (ear), it was first isolated from an ear canal.

## Genetically distinct

More closely related to *C. haemulonii* than to the common *C. albicans*. It belongs to the CTG clade (translates the CTG codon as serine rather than leucine).

## Unique biology

Grows at 42°C · Tolerates high salt concentrations · Thrives on skin · Can switch between White and Brown cell states with different resistance and virulence properties

**CRITICAL**

WHO Priority

**URGENT  
THREAT**

CDC Classification

**42°C**

Max Growth Temp

**5**

Clades

# History & Global Emergence

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- 1996** Earliest known isolate — retrospectively identified from a Korean pediatric bloodstream infection (unrecognized at the time)
- 2009** First formal description: isolated from ear canal of a 70-year-old Japanese woman at Tokyo Metropolitan Geriatric Hospital (Sato et al.)
- 2011** First confirmed invasive infections reported from South Korea; isolates initially misidentified as *C. haemulonii* and *Rhodotorula glutinis*
- 2012–2015** Simultaneous emergence on 3 continents confirmed by whole-genome sequencing (South Asia, Africa, South America) — each clade arising independently
- 2016** CDC issues first clinical alert to US healthcare facilities; first US cases confirmed. *C. auris* reported in >35 countries by 2019
- 2022** WHO designates *C. auris* a Critical Priority Pathogen — one of only 19 fungal pathogens on the first-ever fungal priority list
- 2023** CDC reports 4,514 new US clinical cases — year-over-year increases continuing since first detection in 2013. Clade I now predominant in the US

# Why We Care

Five features that make *C. auris* uniquely dangerous



## Multidrug Resistance

Resistant to  $\geq 1$  antifungal class in most isolates; pan-resistant to all three classes in growing number of cases



## Diagnostic Evasion

Misidentified by standard biochemical methods (API 20C, Vitek 2, Phoenix) as *C. haemulonii* or other species; requires MALDI-TOF or molecular testing



## Silent Colonization

Patients may be asymptotically colonized on skin and body for months to years, serving as reservoirs for ongoing transmission



## Environmental Persistence

Survives on hospital surfaces for weeks to months; resistant to many common disinfectants; found on high- and low-touch surfaces



## High Mortality

30–60% mortality in invasive infections; nearly 50% of infected patients died within 90 days in early US outbreak reports

# Patient Impact & Risk Factors

**30–60%**

Mortality in  
invasive infections

**~90%**

US isolates resistant  
to fluconazole

**>19 days**

Median time from  
admission to infection

**73%**

Patients with central  
venous catheter

## Patient Risk Factors

ICU admission / critical illness

Central venous catheter (CVC)

Recent surgery or invasive procedures

Prolonged broad-spectrum antibiotic use

Prior antifungal therapy (especially fluconazole)

Diabetes mellitus (present in ~41% of cases)

Immunocompromised state (malignancy, transplant, HIV)

Mechanical ventilation

Long-term care facility or hospital exposure

International healthcare travel or transfer

# Diagnosis: The Identification Challenge

 Standard laboratory methods can MISIDENTIFY *C. auris* as other, less dangerous species — a critical barrier to timely treatment and infection control

## ✘ Conventional Methods (FAIL)

- API 20C AUX → *C. haemulonii*
- Vitek 2 → *C. haemulonii* or *C. famata*
- Phoenix → *C. haemulonii*
- MicroScan → *C. guilliermondii*
- BD BBL Crystal → *C. lusitanae*
- Sensititre YeastOne → misID

## ✔ Accurate Methods

- MALDI-TOF mass spectrometry (Bruker Biotyper or Vitek MS) — requires updated database
- Sequencing of ITS region / D1/D2 of 26S rDNA
- *C. auris*-specific PCR panels
- Chromogenic selective media (*C. auris* CHROMID)
- Salt Sabouraud Dulcitol broth at 42°C (screening)

## ⚡ Clinical Pearl

- Alert your lab if *C. auris* is suspected
- Bloodstream infections are most common presentation
- Antifungal susceptibility testing required for ALL clinical isolates
- Report ALL cases to state or local health department
- Contact DOH or CDC if confirmation needed

# Antifungal Resistance

A three-class problem — and it's getting worse

## AZOLES (Fluconazole, Voriconazole)

~90%

of US isolates resistant

### Mechanism:

ERG11 gene mutations; upregulation of efflux pumps (CDR1/CDR2/MDR1)

**✗ Do NOT use fluconazole for *C. auris* treatment**

## ECHINOCANDINS (Micafungin, Caspofungin, Anidulafungin)

5–10%

of US isolates resistant (increasing)

### Mechanism:

FKS1/FKS2 hot-spot mutations; reports of pan-resistant strains are rising

**⚠ First-line treatment — but resistance is emerging**

## POLYENES (Amphotericin B)

~17%

of isolates resistant globally

### Mechanism:

ERG3/ERG6/ERG11 mutations disrupting ergosterol biosynthesis

**⚠ Reserve for echinocandin failure; monitor renal function**



**PAN-RESISTANT *C. auris* (resistant to all 3 classes) is confirmed in multiple US facilities. Consider investigational agents (rezafungin, fosmanogepix, ibrexafungerp).**

# Treatment Approach

## STEP 1 — FIRST-LINE: Echinocandin (adults & children >2 months)

Micafungin 100 mg IV daily | Caspofungin 70 mg IV load → 50 mg IV daily | Anidulafungin 200 mg IV load → 100 mg IV daily

### No Response After 5 Days / Persistent Fungemia

Switch to Liposomal Amphotericin B 5 mg/kg IV daily  
Consider combination therapy; consult Infectious Disease

### Pan-Resistant Isolates: Investigational Agents

Rezafungin · Ibrexafungerp (Brexafemme) · Fosmanogepix · Olorofim | Contact company expanded access programs | Report to CDC

## Key Treatment Pearls

- ★ Susceptibility testing MANDATORY for all clinical isolates — do not rely on species alone
- ★ Patients remain colonized long-term (possibly indefinitely) even after successful treatment — maintain infection control measures
- ★ Do NOT treat colonization without signs/symptoms of active infection
- ★ Consult Infectious Disease for all confirmed *C. auris* infections | Report all cases to your state/local health department

# Infection Control: Stopping Nosocomial Spread

## Hand Hygiene

Soap & water preferred (*C. auris* may persist on hands after alcohol gel alone). Follow WHO 5-moments. Gloves do not replace hand hygiene.

## Patient Isolation

Single-patient room required. Contact precautions (gown + gloves) for all care. Maintain for entire inpatient stay; deisolation criteria rarely met.

## Environmental Disinfection

Use EPA-approved List P disinfectants. Standard quaternary ammonium compounds are INEFFECTIVE. Chlorine-based products ( $\geq 1,000$  ppm), UV-C, and accelerated hydrogen peroxide have demonstrated efficacy.

## Screening & Surveillance

Screen close contacts of newly identified cases. Composite skin swabs (axilla + groin) per CDC SOP. Report immediately to state/local public health. Track contacts across facility transfers.

## Equipment & Transfer

Dedicate non-critical equipment to *C. auris* patients if possible. Communicate *C. auris* status at ALL patient transitions (transfer, discharge, referral). Receiving facilities must be notified.

## Staff Education

Train all staff who may enter room. Audit PPE compliance. Engage pharmacy in antifungal stewardship. Multidisciplinary approach essential for outbreak control.

# Future Outlook

Where is this heading — and what should rural healthcare prepare for?



## Cases Will Continue to Rise

CDC data shows uninterrupted year-over-year case increases since 2016. Rising from scattered cases to endemic in some US regions. Rural facilities will encounter this.

## New Clades May Emerge

Clade V confirmed in Iran. Genomic surveillance is detecting new subclades. The pattern of simultaneous independent emergence on multiple continents is not fully understood.

## Pipeline Drugs Are Coming

Rezafungin (weekly dosing), ibrexafungerp, fosmanogepix, and olorofim represent new classes and mechanisms. FDA approved ibrexafungerp; others in Phase III trials for *C. auris*.

## Climate Change Connection

*C. auris* uniquely grows at 42°C. Casadevall et al. hypothesize its emergence is linked to global warming enabling thermal adaptation. This may drive further fungal pathogen emergence.

## Diagnostics Will Improve

Rapid molecular panels (PCR) increasingly available. MALDI-TOF databases being updated routinely. Point-of-care testing under development — critical for rural hospitals.

## Stewardship Is Essential

Antifungal stewardship programs will become standard of care. Pharmacists and ID-trained clinicians in rural settings are the frontline. Regional networks are forming.

# Key Takeaways & Resources

*C. auris* is a real and growing threat — it has reached all US regions and every rural facility should have a response plan.

Standard lab methods fail to identify it — know your lab's capabilities; update MALDI-TOF databases and establish ID pathways.

Echinocandins are first-line — but resistance is rising. All clinical isolates require susceptibility testing. Call ID.

Infection control is your most powerful tool — contact precautions, correct disinfectants (EPA List P), and prompt public health notification are essential.

Colonized patients remain colonized — infection control measures continue for the entire hospitalization, regardless of treatment response.

Questions? Consult your infectious disease team or state health department when *C. auris* is suspected or confirmed in your facility.

## Key Resources

### ▶ CDC *C. auris* Hub

[cdc.gov/candida-auris](https://cdc.gov/candida-auris)

### ▶ IDSA Candidiasis Guidelines

[idsociety.org](https://idsociety.org) (2016 + updates)

### ▶ WHO Fungal Priority List

[who.int](https://who.int), 2022

### ▶ MMWR *C. auris* Reports

[cdc.gov/mmwr](https://cdc.gov/mmwr)

### ▶ EPA List P Disinfectants

[epa.gov/pesticide-registration](https://epa.gov/pesticide-registration)

### ▶ APHL *C. auris* Guidance

[aphl.org](https://aphl.org)

### ▶ State Health Dept (report cases)

Contact your state HD

# References

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1. Satoh K, et al. *Candida auris* sp. nov., a novel ascomycetous yeast isolated from the external ear canal of an inpatient in a Japanese hospital. *Microbiol Immunol*. 2009;53(1):41-44.
2. Lockhart SR, et al. Simultaneous emergence of multidrug-resistant *Candida auris* on 3 continents confirmed by whole-genome sequencing. *Clin Infect Dis*. 2017;64(2):134-140.
3. Cortegiani A, et al. Epidemiology, clinical characteristics, resistance, and treatment of infections by *Candida auris*. *J Intensive Care*. 2018;6:69.
4. CDC. Clinical Treatment of *C. auris* Infections. Updated April 24, 2024. [cdc.gov/candida-auris](https://www.cdc.gov/candida-auris)
5. CDC. Infection Control Guidance for *C. auris*. Updated April 24, 2024. [cdc.gov/candida-auris](https://www.cdc.gov/candida-auris)
6. Chowdhary A, et al. Multidrug-resistant *Candida auris*: new kid on the block in hospital-associated infections? *PLOS Pathogens*. 2017;13(4):e1006290.
7. WHO. Fungal Priority Pathogens List to Guide Research, Development and Public Health Action. Geneva: WHO; 2022.
8. Jeffery-Smith A, Taori SK, Schelenz S, Jeffery K, Johnson EM, Borman A; Manuel R, Brown CS. *Candida auris*: a Review of the Literature. *Clin Microbiol Rev*. 2017 Nov 15;31(1):e00029-17. doi: 10.1128/CMR.00029-17. PMID: 29142078; PMCID: PMC5740969.
9. Rutala WA, et al. Susceptibility of *Candida auris* and *Candida albicans* to 21 germicides. *Infect Control Hosp Epidemiol*. 2019;40(3):380-382.
10. Casadevall A, Kontoyiannis DP, Robert V. On the Emergence of *Candida auris*: Climate Change, Azoles, Swamps, and Birds. *mBio*. 2019 Jul 23;10(4):e01397-19. doi: 10.1128/mBio.01397-19. PMID: 31337723; PMCID: PMC6650554.
11. Chowdhary A, Jain K, Chauhan N. *Candida auris* Genetics and Emergence. *Annu Rev Microbiol*. 2023 Sep 15;77:583-602. doi: 10.1146/annurev-micro-032521-015858. Epub 2023 Jul 5. PMID: 37406342; PMCID: PMC12962553.
12. CDC. Tracking *C. auris*: US Case Counts. 2024. [cdc.gov/candida-auris/tracking](https://www.cdc.gov/candida-auris/tracking)
13. Cornely OA, Sprute R, Bassetti M, et al. Global guideline for the diagnosis and management of candidiasis: an initiative of the ECMM in cooperation with ISHAM and ASM. *Lancet Infect Dis*. 2025 May;25(5):e280-e293. doi: 10.1016/S1473-3099(24)00749-7. Epub 2025 Feb 13.
14. Sexton DJ, et al. Evaluation of nine surface disinfectants against *Candida auris*. *Infect Control Hosp Epidemiol*. 2020.