New Clonal Strain of Candida auris, Delhi, India

Anuradha Chowdhary, Cheshta Sharma, Shalini Duggal, Kshitij Agarwal, Anupam Prakash, Pradeep Kumar Singh, Sarika Jain, Shallu Kathuria, Harbans S. Randhawa, Ferry Hagen, and Jacques F. Meis

A new clonal strain of Candida auris is an emerging etiologic agent of fungemia in Delhi, India. In 12 patients in 2 hospitals, it was resistant to fluconazole and genotypically distinct from isolates from South Korea and Japan, as revealed by M13 and amplified fragment length polymorphism typing.

In 2009, yeast isolates from the external ear canal of a Japanese patient were identified as a new species, Candida auris, on the basis of sequence analysis of the nuclear rRNA gene. The species is phylogenetically related to C. haemulonii (1). In the same year, 15 isolates of C. auris were reported from otitis media patients in South Korea (2). Genotyping of these isolates revealed a clonal origin (3). In addition, in 3 patients, persistent fungemia caused by C. auris was resistant to fluconazole; 2 of these patients died (4). Fungemia caused by C. auris seems to be extremely rare, however. We report a series of fungemia cases caused by a new clonal strain of this emerging pathogen, involving 12 inpatients from 2 hospitals in Delhi, India.

The Study

We characterized 12 bloodstream C. auris isolates, originating from an equal number of patients, collected during 2009–2011 at 2 hospitals (a tertiary care general hospital and a pediatric center) in Delhi, India. The controls included reference strains of C. auris from South Korea (CBS12372, CBS12373) and Japan (JCM15448, DSMZ21092), C. haemulonii (CBS7801–7802, CBS5149, CBS5150), C. pseudohaemulonii (KCTC1787, CBS10004, JCM12453), and C. duobushaemulonii (CBS7798–7800, CBS9754).

The yeast isolates were oval without pseudohyphae and germ tube formation. They appeared pink on CHROMagar Candida medium (Difco, Becton Dickinson, Baltimore, MD, USA) and grew at 37°C and 42°C. VITEK2 (bioMérieux, Marcy l’Etoile, France) misidentified 10 isolates as C. haemulonii and 2 as C. famata. Similarly, API20C (bioMérieux) also misidentified them as C. sake. C. auris isolates from India assimilated N-acetylglucosamine (NAG) in contrast to the isolates from Japan and South Korea.

Molecular identification was done by sequencing internal transcribed spacer (ITS) and D1/D2 regions (5–7). ITS sequences (GenBank accession nos. KC669203–KC669205) of our isolates showed 100% homology with an unrelated C. auris isolate (GenBank accession no. HE797773) and 98% identity with isolates from Japan and South Korea (GenBank accession nos. AB375772 and EU884189). Also, large ribosomal subunit sequences (GenBank accession nos. KC669203–KC669204) of the India isolates showed 100% homology with an unrelated C. auris isolate (GenBank accession no. HE797774) and 98%–100% identity with isolates from Japan and South Korea (accession nos. AB375773 and EU881960). ITS and large ribosomal subunit sequences of the isolates in our study showed 85%–98% similarity with the closest C. haemulonii complex species. These isolates have been deposited at the CBS-KNAW Fungal Biodiversity Centre, Utrecht, the Netherlands (accession nos. CBS12766–CBS12777).

M13 PCR fingerprinting analysis was performed as described (8,9). The isolates from India had a distinct banding pattern in relation to isolates from Japan and South Korea. Except for 2 isolates (CBS12772, CBS12776), the C. auris isolates showed an identical banding pattern, suggesting a single genotype (Figure 1). The isolates’ genotypic diversity was also determined by using amplified fragment length polymorphism (AFLP) fingerprint analysis (10). In a dendrogram in which standard Pearson and UPGMA (unweighted pair group method with averages) settings were used, C. auris isolates clustered separately from C. haemulonii, C. duobushaemulonii, and C. pseudohaemulonii. The C. auris isolates from India showed a clonal origin and were genotypically distinct from the isolates from Japan and South Korea (Figure 2).

Antifungal susceptibility testing was done by using microbroth dilution according to Clinical Laboratory Standards Institute protocols (11). All isolates showed high geometric mean MICs of fluconazole (Pfizer, Groton, CT, USA) (28.5 mg/L), whereas isavuconazole (Basilea Pharmaceutica, Basel, Switzerland [now Astellas]), posaconazole (Schering-Plough, Kenilworth, NJ, USA [now Merck]), itraconazole (Lee Pharma, Hyderabad, India), and voriconazole (Pfizer) exhibited highly potent activity.
Amphotericin B (Sigma-Aldrich, St. Louis, MO, USA), flucytosine (Sigma-Aldrich), and the 3 echinocandins also showed excellent activity (Table). Clinical features of the infections and response to antifungal therapy of the 12 patients with fungemia are summarized in the online Technical Appendix Table (wwwnc.cdc.gov/EID/article/19/10/13-0393-Techapp1.pdf). Each patient had at least 3 predisposing risk factors. Seven had immunosuppressive conditions: 6 (50%) had diabetes mellitus, 5 (42%) had chronic kidney disease, 2 (17%) had undergone cancer chemotherapy, and 1 (8%) was infected with HIV. Of the 5 children, 3 were infants of low birthweight, 1 neonate had late onset sepsis, and one 10-year-old child had acute lymphoblastic leukemia. Having an indwelling urinary catheter was the predominant risk factor for 83% of patients, followed by receiving broad-spectrum antimicrobial drugs (75%), spending time in an intensive care unit (58%), having neutropenia (50%), and having a central venous catheter (CVC) (42%). Notably, 7 (58%) patients had breakthrough fungemia while receiving fluconazole (1–3 weeks), and persistent candidemia developed in 9 (75%) patients. In 8 of these 9 patients, candidemia cleared after antifungal therapy in 1–3 weeks. For the remaining 3 patients, blood culture reports were available post mortem. C. auris represented 5% of the annual candidemia cases in a pediatric hospital and 30% of annual candidemia cases in a tertiary care general hospital. Overall mortality rates as high as 50% were noted; 30% of deaths were attributed to critical illness.

Conclusions

This study documents the emergence of a new clonal strain of C. auris as an etiologic agent of candidemia in India. Fungemia caused by C. auris is extremely rare, identified only by sequencing. Because diagnostic laboratories do not undertake molecular identification routinely, fungemia is likely to be much more prevalent than published reports indicate. Growth at 40°C may differentiate C. auris isolates, routinely misidentified as C. haemulonii by VITEK. C. haemulonii does not grow at 40°C, as does C. auris (12). Sequencing may confirm their identity.

Although Lee et al. proposed negative assimilation of NAG for differentiating C. auris from C. pseudohaemulonii, this proposal seems invalid because all isolates from India assimilated NAG (4). The phenotypic divergence of these isolates was also supported by M13 and AFLP typing. Notably, C. auris isolates from 2 hospitals in Delhi were clonal, suggesting interhospital transmission. Transmission of the same genotype was observed in intensive care, surgical, medical, oncologic, neonatal, and pediatric wards, which were mutually exclusive with respect to health care personnel. This finding is analogous to that of the clonal transmission of C. auris among 3 hospitals in South Korea (2). Prevalence of a unique clonal strain of C. auris in 2 hospitals in Delhi, 13.5 km apart, underscores the need for more comprehensive studies to determine potential occurrence of other endemic clones in various regions of India.

Because most (83%) of the patients in our study had an indwelling urinary catheter, the source of fungemia could be urogenital colonization. However, no urine samples were cultured. Five (42%) patients had a CVC in situ, but...
none received total parenteral nutrition. Although Candida spp. adhere to intravascular catheters, providing a potential nidus for infection, a previous study showed that C. auris does not form a biofilm (3). Catheter tip cultures were negative for 3 of the 5 patients who had a CVC. Although 57% of patients admitted to critical care units died, the role of C. auris candidemia in fatal cases could not be ascertained because no comparison was available with a population that did not have candidemia.

In conclusion, C. auris is an emerging yeast pathogen that is underreported because it is misidentified in routine diagnostic laboratories. The resistance of C. auris to fluconazole is particularly worrisome.

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Dr Chowdhary is an associate professor and head, Department of Medical Mycology, Vallabhbhai Patel Chest Institute, University of Delhi. Her research interest areas are clinical mycology, antifungal resistance, and ecology of pathogenic fungi.

Table. Results of in vitro antifungal susceptibility testing of 12 Candida auris isolates originating from fungemia patients in 2 hospitals, Delhi, India, 2009–2011

<table>
<thead>
<tr>
<th>MIC, mg/L*</th>
<th>Drug†</th>
<th>AMB</th>
<th>FLU</th>
<th>ITC</th>
<th>VRC</th>
<th>ISA</th>
<th>POS</th>
<th>FC</th>
<th>CAS</th>
<th>MFG</th>
<th>AFG</th>
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<tbody>
<tr>
<td>MIC&lt;sub&gt;50&lt;/sub&gt;</td>
<td>0.25</td>
<td>32</td>
<td>0.125</td>
<td>0.5</td>
<td>0.125</td>
<td>0.125</td>
<td>0.125</td>
<td>0.25</td>
<td>0.25</td>
<td>0.125</td>
<td>0.06</td>
</tr>
<tr>
<td>MIC&lt;sub&gt;90&lt;/sub&gt;</td>
<td>0.5</td>
<td>64</td>
<td>0.25</td>
<td>0.125</td>
<td>0.125</td>
<td>0.125</td>
<td>0.125</td>
<td>0.25</td>
<td>0.25</td>
<td>0.125</td>
<td>0.125</td>
</tr>
<tr>
<td>GM</td>
<td>0.33</td>
<td>28.5</td>
<td>0.15</td>
<td>0.39</td>
<td>0.10</td>
<td>0.12</td>
<td>0.125</td>
<td>0.25</td>
<td>0.23</td>
<td>0.072</td>
<td>0.26</td>
</tr>
<tr>
<td>Range</td>
<td>0.25–1</td>
<td>16–64</td>
<td>0.125–0.25</td>
<td>0.125–1</td>
<td>&lt;0.015–0.25</td>
<td>0.06–0.25</td>
<td>0.125</td>
<td>0.125–0.25</td>
<td>0.06–0.125</td>
<td>0.125–0.5</td>
<td></td>
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*MIC<sub>50</sub>, 50% minimum inhibitory concentration; MIC<sub>90</sub>, 90% minimum inhibitory concentration; GM, geometric mean of MICs.
†AMB, amphotericin B; FLU, fluconazole; ITC, itraconazole; VRC, voriconazole; ISA, isavuconazole; POS, posaconazole; FC, flucytosine; CAS, caspofungin; MFG, micafungin; AFG, anidulafungin.

Figure 2. Dendrogram of amplified fragment length polymorphism analysis of Candida auris isolates from India, Japan, and South Korea and members of C. haemulonii complex. It was constructed by using UPGMA (unweighted pair group method with averages) in combination with the Pearson correlation coefficient and was restricted to fragments of 60–400 bp. Scale bar indicates the percentage similarity.
References


Address for correspondence: Anuradha Chowdhary, Department of Medical Mycology, Vallabhbhai Patel Chest Institute, University of Delhi, Delhi 110 007, India; email: dranuradha@hotmail.com