

Research Brief

Susceptibility of *Candida auris* and *Candida albicans* to 21 germicides used in healthcare facilities

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Candida auris is an emerging fungal pathogen that is often resistant to major classes of antifungal drugs. It is considered a serious global health threat because it can cause severe infections with frequent mortality in more than a dozen countries. It can survive on healthcare environmental surfaces for at least 7 days and can cause outbreaks in healthcare facilities. Clearly, infection prevention strategies, such as surface disinfection, will be essential to controlling *Candida* transmission. Unfortunately, data on the activity of antiseptics and disinfectants used in healthcare to inactivate this pathogen are limited.^{1–5} In this study, we investigated 12 different disinfectants (ie, 8 low- and intermediate-level disinfectants in 2 dilutions of sodium hypochlorite and 5 high-level disinfectants/chemical sterilants) and 9 antiseptics commonly used in healthcare facilities for their antimicrobial activity against *C. auris* and *C. albicans*.

We used the disc-based quantitative carrier testing to evaluate the germicidal activity of multiple antiseptics and disinfectants against the emerging pathogen *C. auris*.^{6,7} We considered the carrier test to mimic disinfectant application on an inanimate surface in a clinical environment better than a suspension test commonly reported by manufacturers and the published literature. The *C. auris* isolate used was Antibiotic Resistance Bank no. 0385 from the Centers for Disease Control and Prevention. Based on tentative minimum inhibitory concentration (MIC) breakpoints,⁸ this isolate was resistant to fluconazole and was susceptible to anidulafungin, micafungin, caspofungin, and amphotericin B. To determine whether *C. auris* susceptibility to germicides was similar to that of other *Candida* species, we also tested *C. albicans* (ATCC strain no. 60193). In brief, 10 µL inoculum containing ~10⁴ with 5% fetal calf serum of *C. auris* or *C. albicans* was placed onto each stainless steel disc (1 cm diameter) and dried in a vacuum desiccator for 2 hours. After drying, each carrier was placed in a plastic vial with the inoculated side up. The dried inoculum was entirely covered by 50 µL of the test germicide for 1 minute at room temperature (~20°C). Then 9.95 mL eluent with neutralizer (Dey/Engley neutralizing

broth) was added into each carrier holder to dilute and neutralize the germicide. Serial dilutions of the eluates were filtered to evaluate the fungal viability and to achieve countable numbers. The membrane filters of appropriate serial dilutions were placed on sheep blood agar plates and incubated for 24–48 hours at 37°C, and the fungal were then counted. We performed 3 replicates for each organism and germicide. Also, 3 carrier controls were prepared during each experiment in the manner described above but without germicide exposure. Compared to mean carrier control counts, the log₁₀ reduction of the test organism for each germicide was calculated.⁹

The efficacy of germicides with active ingredient, product name, manufacturer, and classification against *C. auris* and *C. albicans* are provided in Table 1. Under the challenging test conditions (ie, 5% FCS and 1 minute exposure time), 12 of 22 tested disinfectants and antiseptics (55%) demonstrated at least a 3-log₁₀ reduction, and 16 (73%) demonstrated at least a 2-log₁₀ reduction for *C. auris*. Also, 14 of these 22 (64%) demonstrated at least a 3-log₁₀ reduction, and 17 (77%) demonstrated at least a 2-log₁₀ reduction for *C. albicans*. Of the 9 antiseptics, 7 (78%) did not demonstrate a 3-log₁₀ reduction against *C. auris*; these included 10% povidone-iodine, 0.5% triclosan, 1% chloroxylenol, 1% chlorhexidine gluconate (CHG) with 61% ethyl alcohol, 2% CHG, 4% CHG, and 3% hydrogen peroxide. Of 13 tested disinfectants (low-level disinfectants and high-level disinfectants), 10 (77%) demonstrated at least a 3-log₁₀ reduction at 1 minute, with 3 exceptions: (1) a 1:50 dilution of 5.25% sodium hypochlorite (~1,245 ppm chlorine; Chlorine Test Kit, Model CN-21P, Hach, Loveland, CO); (2) a diluted, water-based quaternary ammonium compound (QAC); and (3) a 0.55% ortho-phthalaldehyde. In general, the log₁₀ reductions for *C. auris* and *C. albicans* were similar: 12 of 13 (92%) within a 1 log₁₀ difference) for the 13 tested disinfectants. But 4 of 9 (44%) of the antiseptics had a >1 log₁₀ difference in susceptibility.

There is no standard level of germicidal efficacy for environmental surfaces, but most of the disinfectants tested that demonstrated at least a 3-log₁₀ reduction are likely to be clinically effective against *C. auris* when used appropriately. Some of the Environmental Protection Agency (EPA)-registered disinfectants used in this study have an EPA registration claim longer than the 1 minute used in this study. All of the FDA-cleared high-level disinfectants have a

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Table 1. Germicidal Activity Against *Candida auris* and *Candida albicans* Using a Quantitative Carrier Test Method

Germicide name	Manufacturer, Location	Active Ingredient	Formulation Tested	Classification	<i>C. auris</i> ^a	<i>C. albicans</i> ^a
Purell Advanced instant hand sanitizer	GOJO, Akron, OH	70% ethanol	Undiluted	Antiseptic	4.0	2.5
Betadine solution	Purdue Products, Stamford, CT	10% povidone-iodine/1% iodine	Undiluted	Antiseptic	2.5	2.3
Medicated Soft 'N Sure	Steris, St. Louis, MO	0.5% triclosan	Undiluted	Antiseptic/Handwash	1.4	1.7
Soft Care Defend	Diversey, Charlotte, NC	1% chloroxylenol	Undiluted	Antiseptic/Handwash	2.8	3.9
Avagard	3M, St Paul, MN	1% chlorhexidine gluconate solution, 61% ethyl alcohol	Undiluted	Antiseptic/Surgical hand scrub	2.0	1.9
Scrub-Stat 2%	Ecolab, St Paul, MN	2% chlorhexidine gluconate solution	Undiluted	Antiseptic/Surgical hand scrub/handwash	1.6	2.8
Scrub-Stat 4%	Ecolab, St Paul, MN	4% chlorhexidine gluconate solution	Undiluted	Antiseptic/Surgical hand scrub/handwash	1.9	3.5
Isopropyl rubbing alcohol 70% USP	Medichoice, Mechanicsville, VA	70% isopropyl alcohol	Undiluted	Antiseptic/Disinfectant	3.8	4.1
Solution of hydrogen peroxide 3% USP	Medichoice, Mechanicsville, VA	3% hydrogen peroxide	Undiluted	Antiseptic	1.4	1.8
Austin's A-1 bleach 1:10	James Austin Co, Mars, PA	5.25% sodium hypochlorite (~6,100–6,700 ppm)	1:10 dilution	Disinfectant	4.1	4.0
Austin's A-1 bleach 1:50	James Austin Co, Mars, PA	5.25% sodium hypochlorite (~1,245 ppm)	1:50 dilution	Disinfectant	1.6	1.5
Vesphene IIse	Steris, St Louis, MO	9.09% o-phenylphenol, 7.66% p-tertiary amylphenol	1:128 dilution	Disinfectant	4.1	3.6
Hydrogen peroxide cleaner disinfectant	Clorox, Oakland, CA	1.4% hydrogen peroxide	Undiluted	Disinfectant	4.1	4.1
Lysol disinfectant spray	Reckitt Benckiser, Parsippany, NJ	58% ethanol, 0.1% QAC ^b	Undiluted	Disinfectant	3.8	4.1
A-456 II disinfectant cleaner	Ecolab, St Paul, MN	21.7% QAC ^c	1:256 dilution	Disinfectant	1.7	1.5
Super Sani-Cloth wipe	PDI, Orangeburg, NY	55% isopropyl alcohol, 0.5% QAC ^d	Undiluted ^f	Disinfectant	3.9	4.1
Prime Sani-Cloth wipe	PDI, Orangeburg, NY	28.7% isopropyl alcohol, 27.3% ethyl alcohol, 0.61% QAC ^e	Undiluted ^f	Disinfectant	4.1	4.1
S40 sterilant concentrate S4000	Steris, Mentor, OH	35% peracetic acid	0.20%	High-level disinfectant/Chemical sterilant	4.1	4.1
Cidex-OPA	Advanced Steril Prod, Irvine, CA	0.55% ortho-phthalaldehyde	Undiluted	High-level disinfectant	2.3	3.8
Cidex	Advanced Steril Prod, Irvine, CA	2.4% glutaraldehyde	Undiluted	High-level disinfectant/Chemical sterilant	4.1	4.1
Oxycide	Ecolab, St Paul, MN	27.5% hydrogen peroxide, 5.8% peroxyacetic acid	1:43 dilution	High-level disinfectant/Chemical sterilant	4.1	4.1
Revital-Ox Resert	Steris, Mentor, OH	2% accelerated hydrogen peroxide	Undiluted	High-level disinfectant	4.1	4.1

Note. QAC, quaternary ammonium compound.

^aValues are shown in mean log₁₀ reductions under a test condition of 10⁴ test organisms with 5% fetal calf serum and 1 minute contact time.

^bQAC: alkyl (C₁₄ 50%, C₁₂ 40%, C₁₆ 10%) dimethyl benzyl ammonium saccharinate 0.1%.

^cQAC: octyl decyl dimethyl ammonium chloride 6.51%; dioctyl dimethyl ammonium chloride 2.604%; didecyl dimethyl ammonium chloride 3.906%; alkyl (50% C₁₄, 40% C₁₂, 10% C₁₆) dimethyl benzyl ammonium chloride 8.68%

^dQAC: n-alkyl (C₁₂ 68%, C₁₄ 32%) dimethyl ethylbenzyl ammonium chlorides 0.25%; n-Alkyl (C₁₄ 60%, C₁₆ 30%, C₁₂ 5%, C₁₈ 5%) dimethyl benzyl ammonium chlorides 0.25%.

^eQAC: didecyl dimethyl ammonium chloride 0.61%.

^fExtract from cloth.

registration claim >1 minute (eg, 8–45 minutes). In summary, with the exception of a water-based QAC and a 1:50 dilution of sodium hypochlorite, our data demonstrate that most disinfectants (10 of 13, 77%) used in healthcare facilities are effective (>3-log₁₀ reduction) against *C. auris*. Importantly, water-based QACs, which are commonly used for surface disinfectants, had limited activity and therefore should not be used for disinfection of environmental surfaces or

noncritical patient equipment in rooms housing patients with *C. auris*.¹⁰ In contrast, 7 of 9 antiseptics (78%) did not achieve a 3-log₁₀ reduction of *C. auris* in 1 minute.

Because *C. auris* can persist on surfaces in healthcare environments, cleaning and disinfecting the patient care environment (daily and discharge/terminal cleaning) with effective products is essential. The CDC has recommended the use of EPA-registered

hospital-grade disinfectants effective against *Clostridium difficile* spores (primarily chlorine-based products).⁸ Our data demonstrate that several other commonly used surface disinfectants (ie, a phenolic, 1.4% improved hydrogen peroxide, and alcohol-quaternary ammonium compounds) are as effective against *C. auris* as chlorine-based products. Other infection prevention strategies to minimize the contribution of the environment to *C. auris* transmission are “no-touch” room decontamination technologies¹⁰ and improved thoroughness of cleaning/disinfecting environmental surfaces using thoroughness indicators (eg, fluorescent markers). Further studies are needed to evaluate other test surfaces (eg, polymer) and to identify infection prevention strategies that prevent contaminated surfaces from being a source of acquisition by patients of this globally emerging pathogen.

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Conflicts of interest. Dr Rutala is a consultant for PDI and Advanced Sterilization Products. Dr Weber is a consultant for PDI. The other authors report no conflicts of interest relevant to this article.

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