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Prevalence and antifungal susceptibility profiles of *Candida glabrata*, *Candida parapsilosis* and their close-related species in oral candidiasis

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Abstract

Objective: To evaluate the importance of *Candida glabrata*, *Candida parapsilosis* and their close-related species, *Candida bracarensis*, *Candida nivariensis*, *Candida metapsilosis* and *Candida orthopsilosis* in patients with oral candidiasis and, to determine the in vitro activities of antifungal drugs currently used for the treatment.

Methods: One hundred fourteen isolates of *C. glabrata* and 97 of *C. parapsilosis*, previously identified by conventional mycological methods, were analysed by molecular techniques. In vitro antifungal susceptibility to fluconazole, itraconazole, miconazole, and nystatin was evaluated by CLSI M44-A2 disk diffusion test, and by CLSI M27-A3 microdilution for fluconazole.

Results: All *C. glabrata* isolates were identified as *C. glabrata sensu stricto*, 93 out of 97 *C. parapsilosis* isolates as *C. parapsilosis sensu stricto*, three as *C. orthopsilosis* and one as *C. metapsilosis*. *Candida glabrata* was mainly isolated in mixed cultures but *C. parapsilosis* complex was more frequent in pure culture. *Candida metapsilosis* and *C. orthopsilosis* were isolated as pure culture and both species were susceptible to all antifungal agents tested. Most *C. glabrata* isolates were susceptible to miconazole and nystatin, but resistant to fluconazole and itraconazole. Azole cross resistance was also observed. *Candida parapsilosis* isolates were susceptible to fluconazole although azole cross resistance to miconazole and itraconazole was observed.

Conclusion: This study highlights the importance of accurate identification and antifungal susceptibility testing of oral *Candida* isolates in order to have an in-depth understanding of the role of *C. glabrata* and *C. parapsilosis* in oral candidiasis.

Keywords: Antifungal susceptibility; *Candida glabrata*; *Candida parapsilosis* complex; Oral candidiasis.

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