

论著

卡泊芬净、米卡芬净对念珠菌体外药物敏感性的动态研究

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摘要: 目的 动态研究卡泊芬净、米卡芬净体外对念珠菌的药物敏感性。方法 参照CLSI公布的M-27A方案微量液体稀释法分别测定卡泊芬净、米卡芬净、氟康唑对85株念珠菌的体外敏感性,并连续7 d观测结果。结果 48 h卡泊芬净对白念珠菌、光滑念珠菌及其他念珠菌MIC₅₀、MIC₉₀中位数分别为0.030μg/mL、0.030μg/mL、0.060μg/mL、0.125μg/mL、0.125μg/mL、0.500μg/mL。48 h米卡芬净对白念珠菌、光滑念珠菌及其他念珠菌MIC₅₀、MIC₉₀中位数分别为0.030μg/mL、0.030μg/mL、0.060μg/mL、0.060μg/mL、0.250μg/mL、0.500μg/mL。48 h氟康唑对白念珠菌、光滑念珠菌及其他念珠菌MIC₈₀、MIC₁₀₀中位数分别为2μg/mL、128μg/mL、64μg/mL、128μg/mL、2μg/mL、32μg/mL。85株念珠菌中未见对3种药物同时耐药的菌株。卡泊芬净组白念珠菌MIC₅₀、MIC₉₀ 24 h后不再升高;光滑念珠菌MIC₅₀ 72 h后不再升高,MIC₉₀ 120 h后不再升高;其他念珠菌组MIC₅₀ 168 h、MIC₉₀ 96 h后不再升高。米卡芬净组白念珠菌、光滑念珠菌MIC₅₀、MIC₉₀ 24 h后不再升高;其他念珠菌MIC₅₀、MIC₉₀在72 h后不再升高。结论 卡泊芬净、米卡芬净对念珠菌属有较好的抗菌作用,其中对白念珠菌、光滑念珠菌作用更强,且MICs随着作用时间延长而升高并存在药物特异性和念珠菌种属特异性。

关键词: 动态 卡泊芬净 米卡芬净 氟康唑 念珠菌 体外药敏

Dynamic study on the susceptibilities of caspofungin and micafungin to *Candida* species *in vitro*

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Abstract: Objective To assess the susceptibilities of *Candida* species to caspofungin and micafungin. Methods Susceptibilities of *Candida* isolates to caspofungin, micafungin and fluconazole were determined by microdilution method based on CLSI M27-A2 for consecutive seven days. Results After 48 hours, the medians of MIC₅₀ and MIC₉₀ of *C. albicans*, *C. glabrata*, and other *Candida* spp. to caspofungin were 0.03 and 0.03 μg/mL, 0.06 and 0.125 μg/mL, 0.125 and 0.5 μg/mL, respectively. The medians of MIC₅₀ and MIC₉₀ of *C. albicans*, *C. glabrata*, and other *Candida* species to micafungin were 0.03 μg/mL and 0.03 μg/mL, 0.06 μg/mL and 0.06 μg/mL, 0.25 μg/mL and 0.5 μg/mL, respectively. The medians of MIC₈₀ and MIC₁₀₀ of *C. albicans*, *C. glabrata*, and other *Candida* species to fluconazole were 2 μg/mL and 128 μg/mL, 64 μg/mL and 128 μg/mL, 2 μg/mL and 32 μg/mL, respectively. None of the 85 isolates of *Candida* species showed cross-resistance to caspofungin, micafungin or fluconazole. To caspofungin, there was no change on MIC₅₀ or MIC₉₀ of *C. albicans* after 24 hours. No increase on MIC₅₀ and MIC₉₀ of *C. glabrata* was observed after 72 hours and 120 hours. Neither did increase on MIC₅₀ and MIC₉₀ of other *Candida* species after 168 hours and 96 hours. To micafungin, there was no change on MIC₅₀ and MIC₉₀ of *C. albicans* and *C. glabrata* after 24 hours and no increase on MIC₅₀ and MIC₉₀ of other *Candida* species after 72 hours. Conclusions Caspofungin and micafungin may have great antifungal activities to all candida species, especially to *C. albicans* and *C. glabrata*. The MICs would increase with the exposure time, which show specificity to drug and *Candida* species.

Keywords: dynamic caspofungin micafungin fluconazole *Candida* species *in vitro* susceptibility

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