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Abstract (oral session)

**Susceptibility testing of *Aspergillus section Flavi* over a 21-year period in a general hospital**

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**Background:** We studied *Aspergillus section Flavi* is an occasional causative agent of disseminated infection among immunocompromised patients. We evaluated the epidemiology and antifungal susceptibility of clinical *Aspergillus section Flavi*. **Methods:** We recorded 195 isolates of *Aspergillus section Flavi* strains between 1988 and 2008. Charts were reviewed for clinical data. Minimum inhibitory concentrations (MICs) were determined using the CLSI M38-A2 method with amphotericin B (AMB), itraconazole (IZ), voriconazole (VZ), posaconazole (POS), terbinafine (TB), caspofungin (CAS), and micafungin (MF). Modal MICs and epidemiological cut-off values (ECVs, encompassing >95% of isolates) were obtained by defining wild-type distributions. **Results:** We divided the study into 2 periods: 1988-98 (40 isolates/36 patients) and 1999-2008 (155 isolates/116 patients). Nineteen patients had invasive aspergillosis (IA) caused by *Aspergillus section Flavi*. The underlying conditions were chronic obstructive pulmonary disease (9), cancer (7), organ transplantation (7), and surgery (1). *Aspergillus section Flavi* was isolated from the respiratory tract [93], ear [60], wounds [23], and other sites [19]). The range/mode of MICs and MECs in µg/ml was as follows: AMB, 2-4/2; IZ, 0.125-1/0.5; VZ, 0.125-2/1; POS, 0.06-0.5/0.5; TB, 0.03-64/0.03; CAS, 0.06->16/0.06; and MF, 0.03->16/0.06. The ECVs (% of isolates covered) were as follows: AMB 4, (100%); IZ, 1 (100%); VZ, 1 (95.6%); POS, 0.5 (100%); TB, 0.125 (97.3%); CAS, 0.06 (93.1%); and MF, 0.06 (93.1%). **Conclusions:** During the study period, the number of patients with IA caused by *Aspergillus section Flavi* increased in our institution. Few isolates showed in vitro resistance to candins, but their susceptibilities were good to amphotericin, azoles, and terbinafine.