Evaluation of fungal diagnostic stains and prolonged sputum culture in cystic fibrosis fungal disease


Objectives: Standard culture may fail to detect fungal pathogens. Reliance upon this technique alone amidst clinical decline or antifungal therapy can make clinical decision-making and disease-monitoring challenging. This pilot study aims to further evaluate the CF fungal airways burden in recognition of this unmet clinical need.

Methods: 16 adult CF patients with a spectrum of fungal disease provided spontaneous sputum samples at a single timepoint to be examined by 3 different techniques: (1) Prolonged fungal cultures (Sabouraud’s Dextrose Agar + chloramphenicol and Dermasel Agar incubated at 37°C 48 h then 30°C and interpreted at Day 7, D14, D21, D28). (2) Microbiology stain: Calcofluor White (CFW). (3) Cytology stain: Grocott’s Methenamine Silver (GMS).

Results: Prolonged fungal culture (>D7) yielded a positivity rate of 57% compared to 35% of samples at <D7. However CFW stain revealed fungal elements eg. hyphae, in 75% of samples. GMS stain revealed fungal findings in 73%. Of the fungal culture-negative samples, stains detected fungal elements in up to 50% (CFW 33%, GMS 50%).

Conclusion: Whilst the sample size in this preliminary cohort is small, the clinical utility for additional evaluation of fungal disease by a panel of laboratory techniques available to most hospital pathology departments is highlighted. Of note, Scedosporium apiospermum and Exophiala dermatitidis, two emerging fungal pathogens required >D7 culture. GMS stain a had greater detection rate for fungal elements compared to CFW in the fungal culture-negative samples. The evaluation of these techniques in monitoring disease over time and for early detection of fungal infection forms the focus of the next stage of this study.

Epidemiological study of Achromobacter xylosoxidans in a reference center for CF treatment in Brazil

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Chronic lung infection by A. xylosoxidans may lead to rapid clinical deterioration in cystic fibrosis patients (CFP). A. xylosoxidans is intrinsically resistant to cephalothin, cefoxitin, cefotaxime, aztreonam and aminoglycosides. The pattern of transmission of A. xylosoxidans from patient to patient is not fully understood. The purpose of this study was to evaluate the frequency of infection by A. xylosoxidans in CFP patients transitioning to adult care in Brazil, the antimicrobial susceptibility and possible clonal similarity between the strain, correlating results with clinical data. During September 2011/2013, isolates were identified by PCR and biochemical tests. The MIC of antimicrobial activity was performed in accordance with CLSI. The macrorestriction profile was analyzed by PFGE (XhoI). From 106 patients, 75% showed two or more cultures with A. xylosoxidans. Their mean age was 18 years, BMI was around 19 and the Shwachman score was 50. However, lung function was very low with the means of FVC = 1.76 L and FEV1 = 1.06 L, and one girl died. Although controversial, two brothers showed the same macrorestriction profile and three other patients, without kinship, showed equal pulsotype. All of these patients showed colonization for other bacteria including P. aeruginosa, S. aureus and Bcc. A. xylosoxidans was 100% sensitive to imipenem, 81.8% to meropenem, 72.7% to piperacillin-tazobactam, 63.6% to ceftazidime, 54.5% to polymyxin B and 9% to ciprofloxacin. In conclusion, our group of patients with A. xylosoxidans showed a very compromised lung function and low Shwachman score without nutritional impairment, whereas bacteria showed a similar sensibility profile to literature.