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Title: Cystic fibrosis: *Achromobacter xylosoxidans* colonized patients have more severe respiratory disease

Dr. Benoît 1714 Godbert benoit.godbert@hp-metz.fr MD ¹ and Dr. Amandine 1715 Briault doc_am_br@yahoo.fr ². ¹ Service De Pneumologie, Hôpitaux Privés De Metz, Metz, France and ² Service De Maladies Infectieuses, CHU De Nancy, Vandoeuvre-les-Nancy, France .

Body: Background: The consequence of *Achromobacter xylosoxidans* colonization on the respiratory function of cystic fibrosis (CF) patients is not well known. Methods: Case-control study including six *A. xylosoxidans*-colonized CF patients and 11 matched uncolonized controls. FEV₁, number of courses of antibiotics (nCA) and number of hospital admissions for respiratory reasons (nHAR) were compared in the two groups from T-3 (3 years before colonization) to T+3 (3 years after colonization). Results: nCA and nHAR were significantly higher from T-3 to T0, at T0 (colonization) and from T0 to T+3 in *A. xylosoxidans*-colonized patients than in controls ($p < 0.05$ for all parameters). Median FEV₁ at T0 ($p = 0.05$) and from T0 to T+3 ($p = 0.03$) was lower and decrease in FEV₁ from T-3 to T+3 was faster ($p = 0.05$) in *A. xylosoxidans*-colonized patients. Moreover, broncho-pulmonary aspergillosis occurred more often in *A. xylosoxidans*-colonized patients ($p = 0.003$). When patients were compared to themselves over the periods T-3 to T0 against T0 to T+3, there was no difference for the variable studied, comprising respiratory function tests. Conclusions: *A. xylosoxidans*-colonized CF patients have inherently more severe respiratory disease than uncolonized patients. However, *A. xylosoxidans* colonization did not affect the evolution of respiratory function.