European Respiratory Society Annual Congress 2013

Abstract Number: 611

Publication Number: 367

Abstract Group: 8.2. Transplantation

Keyword 1: Transplantation Keyword 2: Treatments Keyword 3: Bronchoalveolar lavage

Title: BAL neutrophilia increases mortality in lung transplant patients despite azithromycin treatment

Elly 4874 Vandermeulen elly.vandermeulen@med.kuleuven.be ¹, Stijn 4875 Verleden stijn.verleden@med.kuleuven.be ¹, David 4876 Ruttens david.ruttens@med.kuleuven.be MD ¹, Robin 4877 Vos robin.vos@uzleuven.be MD ¹, Dirk 4878 Van Raemdonck dirk.vanraemdonck@uzleuven.be MD ¹, Geert 4879 Verleden geert.verleden@uzleuven.be MD ¹ and Bart 4880 Vanaudenaerde bart.vanaudenaerede@med.kuleuven.be ¹. ¹ The Leuven Lung Transplant Group, Katholieke Universiteit Leuven, Leuven, Belgium .

Body: Introduction: Azithromycin can attenuate bronchoalveolar lavage (BAL) neutrophilia and can prevent or treat bronchiolitis obliterans syndrome (BOS). An increasing number of patients is found to have BAL neutrophilia despite azithromycin treatment, which might have an impact on BOS-free and overall survival. Methods: Medical records of all patients transplanted between 2002 and 2012 were retrospectively investigated. Three groups of patients were created. Group A: patients already treated with azithromycin before the onset of BOS having an increased BAL neutrophilia (>15%) (n=81); control group B: patients under azithromycin therapy with low BAL neutrophilia (<15%) (n=39) and control group C: patients without azithromycin therapy (n=170). Control patients were matched according to post-operative day of BAL sampling. Results: Group A had a significantly worse outcome compared to group C, in terms of BOS (p<0.0001) and survival (p<0.0001). Between both groups of azithromycin-treated patients, a significant difference towards survival (p=0.022) was demonstrated in favor of group B, although no significance was present towards BOS (p=0.14).

Discussion: Elevated BAL neutrophilia, despite treatment with azithromycin, implies a detrimental effect towards BOS and survival. This group of patients which constitutes 28% of our study population is steadily growing without an effective therapy being available up to now.