

DRESS SYNDROME DURING TREATMENT OF INFECTION WITH NON-TUBERCULOUS MYCOBACTERIA

A. Ryba¹, S. Röseler², S. Blank¹, M. Gately¹, K-P. Ringel³, and U. Sommerwerck¹

¹Department of Pneumology, Allergology, Sleep and Respiratory Medicine, Augustinerinnen Hospital Cologne, Germany,

²Department of Otorhinolaryngology - Head and Neck Surgery, University Hospital RWTH Aachen, Germany

³Immunology Laboratories Aachen, Germany

Introduction

Drug Rash with Eosinophilia and Systemic Symptoms (DRESS) is a rare hypersensitivity syndrome with a mortality of up to ten percent.

Clinical case

A 75-year-old patient with CT morphological bronchiectasis and consolidating infiltrate with a cavity formation in the left lower lobe was initially treated empirically with ampicillin/sulbactam and subsequently with piperacillin/tazobactam and meropenem. When *Mycobacterium avium* was detected, therapy with rifampicin, ethambutol, and clarithromycin was used. Three weeks later the patient developed rapid respiratory deterioration requiring oxygen-high-flow therapy. Imaging showed extensive ground-glass infiltrates on both sides. There was also fever, initially urticarial and then a maculopapular rash, elevated CRP, elevated transaminases, and peripheral eosinophilia (1260/ μ l, corresponding to 13.4%). An acute viral, atypical or HIV infection as well as rheumatoid and collagenous genesis of the symptoms were excluded. The dermato-histological examination was inconclusive. The DRESS validation score was 3 points, corresponding to a possible DRESS syndrome. After short-term therapy with antihistamines and systemic cortisone, a significant regression of the symptoms was achieved. The lymphocyte transformation test (LTT) with clarithromycin was positive (3.5 SI; lymphoblasts 15.1%). With all other medications mentioned above, the LTT was negative.

Conclusion

We interpret the patient's symptoms as DRESS syndrome, most likely triggered by clarithromycin.