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DRESS Syndrome During Treatment of Infection with Non-Tuberculous Mycobacteria

Ryba A^{1*}, Röseler S², Blank S¹, Gately M¹, Ringel KP³, Kunz L¹ and Sommerwerck U¹

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

*Corresponding author: Agnieszka Ryba MD, Department of Pneumology, Allergology, Sleep and Respiratory Medicine, Cellitinnen Augustinerinnen Hospital Cologne, Germany

Abstract

This case report discusses a diagnosis of a Drug Rush with Eosinophilia and Systemic Symptoms (DRESS) as a rare severe hypersensitivity reaction in a 75-year-old female patient, who developed rush, fever, elevated CRP, transaminases and peripheral eosinophilia and severe respiratory insufficiency caused by extensive bipulmonary ground-glass infiltrates during treatment of infection with non-tuberculous mycobacteria.

Keywords: Drug Rush with Eosinophilia and Systemic Symptoms; DRESS; Non-tuberculous mycobacteria; Mycobacterium avium; Lymphocyte transformation test; LTT

Introduction

Drug Rush with Eosinophilia and Systemic Symptoms (DRESS) is a rare (incidence of 1:1,000

to 1:10,000 drug administrations), multifaceted and complex hypersensitivity syndrome, the effects of which can be fatal with a mortality of up to ten percent [1].

Clinical Case

75-year-old female patient with CT А morphological bronchiectasis and consolidating infiltrate with a cavity formation in the left lower lobe (Figure 1) and with a neutrophilia of 74% in the Bronchoalveolar Lavage (BAL) was initially treated empirically with ampicillin/sulbactam and subsequently with piperacillin/tazobactam and meropenem if the CRP value was stagnant. When Mycobacterium avium was detected, therapy with rifampicin, ethambutol and clarithromycin was used. Three weeks after starting therapy, the patient developed rapid respiratory deterioration requiring O2 high-flow therapy. Imaging showed extensive ground-glass infiltrates on both sides (Figure 2). There was also fever, initially urticarial and then a maculopapular rash (Figure 3 [1]), elevated CRP, elevated transaminases and peripheral eosinophilia (1260/µl, corresponding to 13.4%). An acute viral, atypical or HIV infection as well as rheumatoid and collagenous genesis of the symptoms were excluded. The dermatohistological examination was

inconclusive. The DRESS validation score was 3 points (Table 1, [1-3]), 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% [4]). With all other medications mentioned above, the LTT was negative.



Figure 1: Bronchiectasis and consolidating infiltrate with a cavity formation in the left lower lobe.





Figure 3: Maculopapular rash [1].

Table 1: DRESS validation score [1].

Items	Score			Commonto
	-1	0	1	– comments
Fever \geq 38.5 °C	N/U	Y		
Enlarged lymph nodes		N/U	Y	>1 cm and \geq 2 different areas
Eosinophilia $\geqq 0.7 \times 10^9/L \text{ or } \geqq$		N/U	Y	Score 2, when $\ge 1.5 \times 10^9/L$ or $\ge 20\%$ if WBC
10% if WBC < 4.0 \times $10^9/L$				$4.0 \times 10^{9}/L$
Atypical lymphocytosis		N/U	Y	
Skin rash				Rash suggesting DRESS: \geq 2 symptoms: purpur
Extent > 50% of BSA		N/U	Y	lesions (other than legs), infiltration, facial
Rash suggesting DRESS	Ν	U	Y	edema, psoriasiform desquamation
Skin biopsy suggesting DRESS	Ν	Y/U		
Organ involvement		N	Y	Score 1 for each organ involvement, maximal
				score: 2
Rash resolution \ge 15 days	N/U	Y		
Excluding other causes		N/U	Y	Score 1 if 3 tests of the following tests were
				performed and all were negative: HAV, HBV, HC
				Mycoplasma, Chlamydia, ANA, blood culture

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ANA: anti-nuclear antibody; BSA: body surface area; HAV: hepatitis A virus; HBV: hepatitis B virus; HCV: hepatitis C virus; N: no; U: unknown; WBC: white blood cell; Y: yes.

Summary

We interpret the patient's symptoms as DRESS syndrome, most likely triggered by clarithromycin. Alternatively, if the prick test for azithromycin was negative, this was carefully increased in dosage, well tolerated and combined with clofazimine. Despite the negative LTT with rifampicin, it was not used because the risk of this medication for DRESS was known. After a negative prick test and patch test, ethambutol was added.

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