Telaprevir-related DRESS syndrome complicating hepatitis C treatment

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ABSTRACT
In chronic hepatitis C patients telaprevir attracts attention with high sustained virologic response and short term treatment however it is associated a new spectrum of adverse events, especially several cutaneous manifestations. We report a 66-year-old female patient developed drug reaction with eosinophilia and systemic symptoms during telaprevir based hepatitis C treatment. All antivirals were discontinued and she was treated with supportive care. Systemic and cutaneous symptoms resolved in follow-up. Although rare, clinicians should be aware of potentially severe cutaneous skin reactions during telaprevir-based therapy. J Microbiol Infect Dis 2015;5(1): 36-37

Key words: Eosinophilia, hepatitis C, rash, telaprevir

Hepatitis C tedavisinde karşılaşılan telaprevir ilişkili DRESS sendromu

ÖZET

Anahtar kelimeler: Döküntü, eozinofili, hepatit C, telaprevir

INTRODUCTION
In chronic hepatitis C patients telaprevir (TVR) attracts attention with high sustained virological response and short term treatment however it is associated a new spectrum of adverse events.1 Several cutaneous manifestations related with TVR have been reported.2 We report a 66-year-old female patient developed drug reaction with eosinophilia and systemic symptoms (DRESS) during TVR based hepatitis C treatment. To our knowledge, our patient is the first reported case with telaprevir-related DRESS from Turkey.

CASE
A 66-year-old female with chronic hepatitis C in week seven of treatment with peginterferon alfa-2a, TVR and ribavirin, presented with generalized pruritic maculopapular eruption involving her trunk, abdomen, arms, face and legs bilaterally (Figure 1). The rash started during 6th week of treatment and subsequently generalized, involved >90% of her body surface area. The physical examination revealed fever, 38.5 °C, and the general especially facial oedema (Figure 2). She had malaise, nausea and vomiting. She was hospitalized. Complete blood count revealed: WBC: 3.5 µL (4.5-10), Hgb: 8.8 g/dL, plt: 93 µL, neutrophil 47%, eosinophil 19.6% and lymphocytes 27%. Serum biochemical tests were: AST: 85 IU/L (0-31), ALT: 158 IU/L (0-41), ALP 255 IU/L (30-120), BUN: 38 mg/dL (5-20), creatinine: 1.9 mg/dL (0.6-1), total IgE 1100 IU/ml (10-180). She was diagnosed DRESS. The antiviral medications were discontinued. She was treated with supportive care and hydration. Systemic antihistamines and topical steroids were started. In three days her fever was under control. Systemic symptoms resolved in ten days. Cutaneous symptoms resolved completely in four weeks subsequent to cessation of therapy. She had achieved rapid virologic response with undetectable virus at 4th week and also follow-up visit in third month HCV-PCR was still undetectable.
**DISCUSSION**

TVR has been shown to dramatically improve sustained virologic response however increased serious cutaneous adverse effects have been reported. Pruritus, eczematous or cutaneous eruption is observed in 56%, besides, severe cutaneous adverse reactions were reported in 3.7% of patients receiving TVR. In phase II and phase III trials 11 patients (0.4%) with DRESS and three patients with suspected Stevens-Johnson syndrome were reported. Rash is most often observed within the first 4 weeks, but can occur at any time. There is no identified predictive factor, and skin reaction mechanism is unknown. Our patient had complaints of pruritus since first weeks of treatment. She had advised to use emollient creams and lipid-rich lotions. However she presented with generalized rash in the seventh week of treatment diagnosed as DRESS. DRESS is a life-threatening condition that is characterized by the clinical triad of fever, rash, and internal organ involvement. The liver is the most frequently involved internal organ. Hypereosinophilia is the third common reported finding in DRESS syndrome. Our patient admitted with fever and the levels of aspartate aminotransferase and alanine aminotransferase increased by approximately 3-fold above the normal limits, she had 19.6% eosinophilia.

In the study of Roujeau’s et al, the incidence was significantly higher with age above 45 years, body mass index below 30, white race, and first HCV therapy. Our patient was elderly and her body index was 27, but she was receiving therapy for the second time because of relapse.

In telaprevir phase III trials, the rate of discontinuation of antiviral drugs due to skin manifestation was low. In the management of dermatological side effects, education of patient about good skin care practices is important. In case of severe cutaneous reaction, discontinuation is strongly recommended. Cutaneous and systemic symptoms usually improve after discontinuation and support care. Our patient’s fever was under control in a few days and skin lesions resolved in four weeks.

In conclusion, although majority of TVP-related cutaneous manifestations is mild or moderate, life-threatening severe conditions like DRESS should be kept in mind.

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**REFERENCES**