

Commentary: Antituberculosis Drug Induced Fixed Drug Eruption– A Case Report

Jitendra H. Vaghela*

Senior resident, Department of Pharmacology, Government Medical College, Bhavnagar, Gujarat, India

Article Info

Article Notes

Received: August 3, 2018

Accepted: October 20, 2018

*Correspondence:

Dr. Jitendra H. Vaghela, Senior resident, Department of Pharmacology, Government Medical College, Bhavnagar, Gujarat, India; Email: drjitendravaghela@gmail.com

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Fixed Drug Eruption (FDE) is characterized by a single or multiple oval, erythematous patches due to systemic exposure to a drug that mostly resolves with a residual hyper pigmentation¹. The overall incidence of FDE from FDC ranges from 3.77% to 15.34%^{2,3}. Fixed drug eruption is one of the serious conditions affecting individual's quality of life. Presently, it is known to occur due to many medications. Among the known medications, some antituberculosis drugs are also one of the causative drugs for fixed drug eruption⁴. Skin reactions due to some antituberculosis drugs are commonly found in patients with history of drug allergy⁵.

According to recent guidelines, use of fixed dose drug combinations (FDC) are promoted for its better compliance and advantages. World Health organization also advocates the use of FDCs. In India, under Revised National Tuberculosis Control Programme (RNTCP), from 1st April 2018 fixed dose drug combinations have been started for treatment of tuberculosis with daily given drug regimens. Previously thrice a week individual drug regimen was provided by the programme⁶. Fixed dose drug combinations have its own advantages. It decreases the number of tablets, frequency of tablets and also improves patient adherence to the treatment.

In present case, patient was prescribed FDC of antituberculosis drugs and developed FDE⁷. Fixed drug eruption (FDE) was caused by fixed dose combination (FDC) of antituberculosis drugs- tablet forecox (rifampicin 225 mg + isoniazid 150 mg + pyrazinamide 750 mg + ethambutol 400 mg) in 40-year-old male patient with previous history of drug allergy. Patient developed FDE after taking the third dose of tablet forecox for pulmonary tuberculosis. After taking the third dose, patient noted multiple, discrete, hyper pigmented patches on nape of neck; around mouth; both eyes; lower abdomen; back and upper abdomen. Patient was advised not to take antituberculosis drugs further and he was treated with injection dexamethasone 4 mg i.m. stat, then oral prednisolone 5 mg 6 hourly; injection ceftriaxone 1 gm i.v. 12 hourly; framycetin cream for local application; mucaine viscous gel per orally 12 hourly; betadine gargles and tablet multivitamins for 15 days. Tablet forecox was withdrawn and the reaction was recovered after 15 days of treatment for FDE. As per WHO-UMC and Naranjo's causality assessment criteria, the association between reaction and tablet forecox was possible and probable, respectively^{8,9}. The reaction was moderately (Level 4b) severe as per Modified Hartwig and Siegel's scale¹⁰.

Among patients from low socioeconomical class and having comorbid conditions, adverse drug reactions due to drugs may



Figure 1: Fixed drug eruption lesion

occur more frequently¹¹. In present case, patient was from lower socioeconomical class, labourer, alcoholic and suffering from comorbidity. Patient has been prescribed antituberculosis drug therapy by private practitioner. Patient history revealed that he had some unknown drug allergy. But, it is one of the limitations of case report that no confirmation was possible for the drug to which patient was allergic. We were unable to perform the patch test for identifying the culprit drug. Patient was diagnosed to suffer from comorbid condition of HIV infection, so patient was started on antiretroviral therapy during the time of hospitalisation. This can be also considered as one of the limitations of the study for identifying the culprit drug.

To conclude, fixed drug dose combinations are widely accepted but before using such combinations for each and every patient one has to keep in mind the possibility of history of drug allergy.

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