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Case Report

Fixed drug eruption associated with fixed combination of fluoroquinolone-nitroimidazole

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ABSTRACT

Fixed drug eruptions (FDE) are cutaneous adverse drug reaction characterized by well demarcated erythematous plaques which on removal of the offending agent resolves with residual hyperpigmentation patches at the site. FDE to nitroimidazoles and fluoroquinolones have nevertheless been infrequently reported. Awareness about the adverse reaction to the fluoroquinolone-nitroimidazole combination drug and also the likelihood of recurrence with same or similar drugs and the possible cross reaction is eminent. Hereby we report one such case of FDE to ciprofloxacin/tinidazole combination.

Keywords: FDE, ciprofloxacin, Tinidazole

INTRODUCTION

Fixed drug eruptions (FDE) are cutaneous adverse drug reaction characterized by well demarcated erythematous plaques which on removal of the offending agent resolves with residual hyperpigmentation patches at the site.

The incidence is often between 20 and 40 years of age.¹ The lesions often occur at sites such as extremities, lips, genitalia and perianal regions. Antimicrobials are one of the most common drugs implicated to cause FDEs, especially reaction to nitroimidazoles and fluoroquinolones have nevertheless been infrequently reported.¹⁻³ Cross-reaction between fluoroquinolones, nitroimidazoles, fluoroquinolone-nitroimidazole combinations have also been previously reported.^{1,4} Awareness about the adverse reaction to the fluoroquinolone-nitroimidazole combination drug and also the likelihood of recurrence with same or similar drugs and the possible cross reaction is eminent.

Hereby we report one such case of FDE to ciprofloxacin/tinidazole combination.

CASE REPORT

A 39-year-old female patient presented with fever, nausea and vomiting. She was diagnosed as acute gastroenteritis and was prescribed with tablet ciprofloxacin 500 mg/tinidazole 600 mg fixed dose combination (FDC) twice daily along with tablet paracetamol.

After one day of consuming the medication, patient developed generalized multiple fluid filled eruptions over the upper arms which ruptured spontaneously leaving hyperpigmented patch at the sites on discontinuation of the FDC.

Upon further inquiry, she had no known allergy to paracetamol and also revealed that she had no history of exposure to the FDC of ciprofloxacin and tinidazole combination in the past. On the Naranjo's causality assessment scale, the adverse event was 5 indicating a 'probable' reaction to the FDC.



Figure 1: Hyperpigmented patches over the arm.

DISCUSSION

FDC containing ciprofloxacin a broad-spectrum quinolone antibiotic and tinidazole a nitroimidazole derivative antiprotozoal is widely used in the treatment of diarrhoea in India.² However, these FDC are not a part of WHO essential medicine list. Incidence of FDE, drug-induced anaphylaxis and erythema multiforme and vomiting have been reported with the use of Ciprofloxacin/tinidazole combination.^{2,3,5} Adverse drug reaction to this FDC makes it difficult to determine the possible offending drug due to cross sensitivity.¹ FDE involves activation of cell-mediated cytotoxic response with the offending drug act as a hapten that binds to basal keratinocytes, leading to an inflammatory response.⁵ Though rechallenge with the offending drug is the gold standard for confirmation of FDE, in this case it was not attempted as the patient was not willing for the fear of aggravation of the lesions.

CONCLUSION

Judicious use of the ciprofloxacin/tinidazole FDC is required in view of cosmetic disfigurement seen with

FDEs. Patients should be counselled on recurrence of FDE with the offending drug and possible cross-reactions of similar medications.

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