CASE REPORT

Unusual fixed drug eruption due to rifampicin

A. K. JAISWAL, S. S. VAISHAMPAYAN, R. VERMA & K. VARADA
Department of Skin, STD and Leprosy, Base Hospital, Lucknow 226002, India

Accepted for publication 25 November 1999

Introduction

Cutaneous reactions due to rifampicin (RFM) are uncommon and amongst these fixed drug eruption (FDE) is especially rare. There have been only three reports of FDE to RFM in the literature so far. In 1985, Naik et al. first reported a peculiar urticarial type of FDE due to RFM. Here, we describe a second such case with a similar unusual cutaneous drug eruption.

Case report

A 24-year-old soldier suffering from borderline tuberculoid leprosy (BT) was admitted to the Leprosy Center, Base Hospital, Lucknow for supervised institutional therapy. He was put on cap RFM 600 mg once a month and dapsone 100 mg daily. The patient developed a solitary itchy erythematous urticarial lesion of 3.5 cm diameter with typical peau d’orange appearance (Figure 1), on the right side of the chest wall about 1 h after the second monthly dose of RFM. The lesion subsided on its own within 1 h without any residual hyperpigmentation. A similar lesion at the same site was observed following the third monthly dose of RFM which also subsided within 1 h without treatment. At this stage, a clinical diagnosis of FDE was made and all drugs were discontinued. Two days later, the patient was subjected to a provocation test with 600 mg of RFM, following which he developed an urticarial lesion at the same site. The provocation test to dapsone did not evoke any cutaneous reaction. Skin biopsy of the lesion revealed slight dermal oedema and sparse eosinophilic infiltration. Interestingly, this cutaneous response to RFM ceased after the fifth dose of RFM.

In our patient, recurrence of a wheal at exactly the same site following administration of RFM is highly suggestive of FDE. Though FDE can be urticarial, the residual hyperpigmentation that follows a classical FDE was not seen. Moreover, the histopathology was...
also consistent with urticaria without pigmentory incontinence. The possibility of a type I lepra reaction was not considered, in view of the onset, further evolution of cutaneous eruption and no change in the pre-existing BT lesions.

The outstanding feature of this case is the unusual non-pigmenting FDE to RPM. Moreover, in contrast to the case reported by Naik et al., the urticarial FDE in our case ceased after the fifth dose of RPM. This is probably because sometimes the inducing drug can be readministered without exacerbation and there may be a refractory period after the occurrence of FDE.

To conclude, this case report raises the question whether the urticarial type of FDE due to RPM is truly rare, or is simply under-reported because of its trivial and transient nature. Further studies on this subject may provide an answer.

References