deficiency 5-15mg, riboflavin two times daily for two weeks is curative. Simple emollients make skin moist, reduce intensity of itching. Tricyclic antidepressant doxepin 25-50mg, per day helps some patients. Anxiolytic alprazolam or antipsychotic thioridazine is helpful in some. Systemic or topical therapy has to be given intermittently for a long time.

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# ANNULAR ERYTHEMATOUS LESION SECONDARY TO IMMUNOTHERAPY

## To the Editor

Immunotherapy is a relatively safe treatment. However systemic and cutaneous side effects can occur.<sup>1</sup> We report a case of annular erythematous lesion at the injection site in a patient receiving immunotherapy.

A 36- year- old male photographer was referred to the Dermatology department for evaluation of a skin rash following desensitization injections. He was suffering from allergic rhinitis since 8 years and had been prick tested with 22 allergens consisting of mite, pollen, fungi, insects, dusts, danders and foods obtained from Allergen Division Curewell (India) Ltd. He was tested positive for the following: Pollens - Chenopadium ablum, Ricinus communis, Cassia siamea; insects - Male and female cockroaches; dog epithelia; culvularia fungus and house dust mite. Immunotherapy was commenced with a mixture of allergens.

The patient initially received two injections per week from a vial containing 1:5000 dilution of the solution without developing any side effects. When injections with 1: 500 solution were commenced he developed pruritic erythematous papular lesions in an annular fashion around the injections site. The lesion used to appear within 24

hours and resolve completly within 10 days without any residual pigmentation or scarring. He developed this lesion following each injection taken weekly. No lesions appeared in other parts of the body and there was no history of angioedema. A skin biopsy revealed epidermis with foci of mild spongiosis and exocytosis of inflammatory cells. Dermis showed moderate perivascular and peri-follicular lymphocytic infiltrate.

Various dermatologic manifestations following desensitization treatment reported include local urticarial reactions which are by far the most common, <sup>2</sup> others being digital vasculitis, <sup>3</sup> persistent, itchy subcutaneous nodules <sup>4</sup> and cold urticaria. <sup>5</sup> In our case the skin lesion could have been a delayed hypersensitivity reaction as the patient developed it later in the course of therapy, sensitization having been induced with the initial injections.

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## CREEPING ERRUPTION

## To the Editor

Recently we have come across a case of creeping eruption in a 6-month-old infant on the back of trunk. A 6-month-old female child was brought to the Dermatology OPD with a tortuous, linear narrow lesion on the back of trunk of 3-4 weeks duration. It started as a red papule from which a linear cord-like lesion progressed. Mother of the child used to lay her child on ground during working in field. Examination revealed a linear and curved erythematous track of 1 to 1.5 cm. As the tunnel advanced on one end, the opposite end became scaly, crusted and finally cleared. There was no other skin lesion. General physical and systemic exmination did not reveal any abnormality.

Routine laboratory investigations on blood, urine and stool were normal. Two tablets of 500 mg thiabendazde, triturated in 10 gm petrolatum was applied twice a day over the lesion. The track cleared within two weeks.

Cutaneous larva migrans (creeping eruption) is caused by the larva of nematode parasites for which man is abnormal final host. *Ancylostoma braziliense* is the most frequent cause, though the larva of other hook worms may also produce it. The common area of skin involved

are the feet, buttocks and hands, though rarely it has been reported from other sites also.<sup>1,2</sup> In present case lesion was on back which is thought to be uncommon site in this part of the country. Demonstration of a persistent, progressive, erythematous, serpiginous eruption on common site of skin is usually sufficient for the diagnosis. Occurrence of larva migrans in the infant is quite rare though few cases have been reported.<sup>2,3</sup> A 68-day-old child has been reported with larva migrans of 65 days during a precipitated labour.<sup>3</sup> In our case age and site are unusual which prompted us to report this case.

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