Norwegian scabies following topical steroid therapy

ROGER CLAYTON*  
M.B., B.S., M.R.C.P.

SIMON FARROW  
M.B., B.S.

Summary  
Following an outbreak of scabies in a hospital ward, a correct diagnosis of Norwegian scabies was made in an eczematous patient whose skin had unexpectedly deteriorated with topical potent fluorinated corticosteroid therapy.

Introduction  
Reports of Norwegian or crusted scabies occurring in England are rare (Calnan, 1950). In this condition itching is not a constant symptom, burrows may not be present and the acarus may be found on the head and neck. The acarus is seen within scales and the gross hyperkeratosis may give the skin the appearance of the bark of a tree. Norwegian scabies has been reported in mental defectives (Zoon and Mali, 1949; Calnan, 1950; Maguire and Kligman, 1960), in cases of lymphoma and leukaemia (Logan, Grant and Keczkes, 1967; Walshe, 1967) and during immunosuppressive therapy (Paterson, Allen and Beveridge, 1973). It is frequently diagnosed after the onset of an epidemic (Ingram, 1951; Wells, 1952). Recently an atypical case of scabies was reported in a 3-month-old child previously treated with topical fluorinated steroids (Macmillan, 1972). The following appears to be the first reported case of classical Norwegian scabies occurring in a patient treated with such steroids.

Case report  
A 62-year-old male patient with a long past history of endogenous eczema was admitted to hospital for treatment. He had extensive involvement being virtually erythrodermic, with much scaling from the waist down. Itching was minimal. He had been treated as an outpatient with about 50 g daily of 0.025% betamethasone 17-valerate. He was treated with approximately 50 g clobetasol propionate 0.05% (Dermovate) ointment daily. The condition of his skin became worse with gross hyperkeratosis, generalized dusky erythema, fissures in the flexures and generalized lymph node enlargement. He was started on a course of oral prednisolone, but without improvement and he became languid and irritable. At this time two other patients in the ward presented with an irritating rash diagnosed clinically as scabies. Shortly afterwards two nurses, six further patients and the ward houseman were similarly diagnosed. The skin patient was re-examined and many acari...
Case reports

Fig. 2.

Fig. 3.
were seen in his scales under the microscope. He was treated with four applications of gamma benzene hexachloride and his skin and mood quickly returned towards normal. All the patients, resident medical and nursing staff of the ward were similarly treated. Subsequently scabies was clinically diagnosed and successfully treated in the two flatmates of one of the nurses, six other nurses, a visitor to the patient, several of his family who visited, three patients attending out-patients who had been discharged before the mass anti-scabietic treatment, one of the ward consultants and his wife and the authors—a total of about thirty cases.

Discussion

The onset of itching in scabies is thought to be due to hypersensitivity to acarine antigen. The immune response is cell-mediated, probably with some skin IgE (Burgess, 1973). Indeed, with reinfection, itching starts at once (Mellanby, 1972). Failure of removal of acari by scratching is thought to be a major factor in the development of Norwegian scabies (Mellanby, 1972) hence its occurrence in mental defectives with or without sensory dysfunction, and debilitated patients with malnutrition and systemic disease. It has been postulated that immunosuppressant therapy may inhibit the primary immune response (Paterson et al., 1973). The occurrence of Norwegian scabies in patients with lymphoma and leukaemia may be the result of defective immune mechanisms. Topical steroids suppress cell mediated immunity in the skin (Weston et al., 1972), hence the lack of papular reaction and itching.

Acknowledgments

We thank Dr M. Feiwel, Dermatologist, St Charles' Hospital, London W.10, for permission to publish this case.

References


Norwegian scabies following topical steroid therapy

Roger Clayton and Simon Farrow

Postgrad Med J 1975 51: 657-659
doi: 10.1136/pgmj.51.599.657

Updated information and services can be found at:
http://pmj.bmj.com/content/51/599/657

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/