A rare case of Hodgkin's lymphoma presenting with papulonecrotic tuberculid

Kundan Mishra,¹ Aditya Jandial,¹ Kaniyappan Nambiyar,² Pankaj Malhotra¹

¹Department of Internal Medicine, Post Graduate Institute of Medical Education and Research, Chandigarh, India
²Department of Pathology, Post Graduate Institute of Medical Education and Research, Chandigarh, India

Correspondence to
 Dr Kundan Mishra, mishrak20@rediffmail.com

Accepted 15 June 2017

DESCRIPTION

A 78-year-old man complained of pruritic skin lesions over the trunk and all limbs and occasional fever for 3 months. He had papulonodular skin lesions with erythematous base involving anterior as well as posterior aspects of the trunk and all limbs with predilection for extensor surface (figure 1). Mantoux test was negative. Skin biopsy showed features of papulonecrotic tuberculid (figure 2) with a negative *Mycobacterium tuberculosis* PCR and acid-fast bacilli stain. CT scan of the chest and abdomen showed multiple, enlarged mediastinal and retroperitoneal lymph nodes (largest 2.4 cm). Trucut biopsy and immunohistochemistry from retroperitoneal node revealed features of classical Hodgkin's lymphoma, nodular sclerosis

variety with CD15 and CD30 positivity. In the absence of conclusive evidence of tuberculosis, he was treated with chemotherapy alone (oral prednisolone, etoposide, procarbazine and cyclophosphamide). Two months later, he is asymptomatic with partial resolution of skin lesions.

Papulonecrotic tuberculid possibly results from hypersensitivity to *M. tuberculosis*. Isolation of *Mycobacterium* from cultures as well as by PCR has been described in the previously reported cases.¹ There has been evidence that antitubercular therapy is the treatment of this condition.²

Paraneoplastic skin lesions have been associated with Hodgkin's lymphoma like ichthyosis, acrokeratosis, urticaria, erythema multiforme and hyperpigmentation.³ However, there has not been any case report of Hodgkin's lymphoma presenting with papulonecrotic tuberculid when searched on PubMed. In our case, there was clinical and histopathological evidence of both the diseases in the same individual.

Figure 1 Papulonodular skin lesions with erythematous base involving lower limbs (A), upper limb (B), anterior as well as posterior aspect of trunk (C, D) with predilection for extensor surface. Inset (E) showing distinct papulonodular lesions.

Figure 2 (A) Skin biopsy showing wedge-shaped dermal necrosis with its broad base toward the epidermis and vasculitis, H&E x40. (B) Skin biopsy showing vasculitis and interstitial granuloma with histiocytes, lymphocytes and few epithelioid cells, H&E x 400.

Learning points

- ▶ Papulonecrotic tuberculid is a rare manifestation of hypersensitivity to *Mycobacterium tuberculosis*.
- ▶ In immunocompromised patient, the purified protein derivative test, acid-fast bacilli stain and culture all might be negative; however, typical histopathological features can clinch the diagnosis of papulonecrotic tuberculid.

Contributors KM, AJ, KN and PM were involved in patient care at various stages. KN prepared the histopathology slides. KM and AJ prepared the manuscript, while all authors read it and agreed to it.

Competing interests None declared.

Patient consent Obtained.

Provenance and peer review Not commissioned; externally peer reviewed.

© BMJ Publishing Group Ltd (unless otherwise stated in the text of the article) 2017. All rights reserved. No commercial use is permitted unless otherwise expressly granted.

REFERENCES

- 1 Tirumalae R, Yeliur IK, Antony M, et al. Papulonecrotic tuberculid-clinicopathologic and molecular features of 12 Indian patients. *Dermatol Pract Concept* 2014;4:17–22.
- 2 Sinha P, Sinha A, Baveja S, et al. A common but underreported entity of papulonecrotic tuberculid. *Med J Armed Forces India* 2015;71(Suppl 1):S237–9.

CrossMark

To cite: Mishra K, Jandial A, Nambiyar K, et al. *BMJ Case Rep* Published Online First: [please include Day Month Year]. doi:10.1136/bcr-2017-220484

3 Lucker GP, Steijlen PM. Acrokeratosis paraneoplastica (Bazex syndrome) occurring with

acquired ichthyosis in Hodgkin's disease. *Br J Dermatol* 1995;133:322–5.

Copyright 2017 BMJ Publishing Group. All rights reserved. For permission to reuse any of this content visit <http://group.bmj.com/group/rights-licensing/permissions>.

BMJ Case Report Fellows may re-use this article for personal use and teaching without any further permission.

Become a Fellow of BMJ Case Reports today and you can:

- ▶ Submit as many cases as you like
- ▶ Enjoy fast sympathetic peer review and rapid publication of accepted articles
- ▶ Access all the published articles
- ▶ Re-use any of the published material for personal use and teaching without further permission

For information on Institutional Fellowships contact consortiasales@bmjgroup.com

Visit casereports.bmj.com for more articles like this and to become a Fellow