PAPULONECROTIC TUBERCULID
(CASE REPORT)

MOHAMMED MOHY EL- DIN
SELIM, MD And
BAHRAM AZADEH, MD, FRC Path.

Abstract

We present in this article a 48 years old female who suffered from papulonecrotic tuberculid for 4 years. The diagnosis is based on the clinical picture, the highly positive tuberculin test, the histopathologic findings and the history of contact with an open pulmonary tuberculosis.

Introduction

Tuberculids are chronic, recurrent skin disorders which affect persons who have had personal or family history of active tuberculosis with a strongly positive Mantoux test (1).

Tuberculids result from hematogenous spread of mycobacteria in a patient with a moderate to a high degree of immunity. The mycobacterium tuberculosis cannot be isolated from the lesion but the mycobacterial DNA could be demonstrated by Polymerase chain reaction (PCR), [2,3,4].

The relation between the degree of the patient immunity and virulence of the mycobacteria determines the development and features of the tuberculids [1,5]. The tuberculids include lichen scrofulosorum, populonecrotic tuberculid (PNT), erythema induratum (nodular vasculitis), erythema nodosum and recently a nodular granulomatous phlebitis of the skin was described as a form of tuberculid.

Case Report

A female born 1949 from Iraq. She has been suffering from sore red hands and feet. The duration of her complaint was 4 years for the hands and 2.5 years for the feet. Her condition was aggravated in winter or by exposure to cold and gets better in summer. She gave history of being in contact with a case of active pulmonary tuberculosis. On examination she was a normotensive female with good general condition. Both hands were reddish blue with tiny papules (3-4 mm) on the dorsal surface of hands and flexor aspect of both wrists. [Fig. 1,2,3,4,]. Some papules had central necrosis. Both feet showed reddish blue papules dorsally and in tendoachilles region [Fig. 5,6,7,8]. Most of the lesions on clearing leave atrophic pigmented scars.

The condition was provisionally diagnosed as panniosis, erythromelalgia and PNT.

Fig. 1 and 2: left and right hands showing bluish red colour and papular lesions
Fig. 3 and 4: left and right palms showing bluish red colour with papules at both wrists.

Fig. 5 and 6: left and right feet showing papular lesions.

Fig. 7 and 8: left and right tendo-aehes region showing papular lesions.
The investigations done showed the following results:

1. Microcytic anemia with WBC count of 4.3, high monocytes (15.5) low neutrophils (32.6), high eosinophils (6.6) and raised E.S.R. (51).

2. Blood biochemistry showed normal blood sugar, kidney function, liver function, blood proteins and high cholesterol (6.17 m.mol/l).

3. Serology and immunology showed normal ANA and anti DNA antibodies, normal C3 and C4, negative rheumatoid factor, cryoglobulins, treponema IgG and negative serology for hepatitis A, B, C and normal G6PD.

4. Tuberculin test was highly positive (more than 30 mm with central bulla and necrosis [Fig.9]

5. X-ray chest showed prominent hilae with increased bronchovascular marking and reticular shadowing seen on both sides. Both CP angles were clear and cardiac size was within normal limits.

6. Biopsy showed a relatively large, elliptical zone of dermal necrosis with vasculitis, inflammatory cellular infiltrates and ulceration of overlying epidermis. Inflammatory and necrotic changes were centered along philosebacous follicles and blood vessels [Fig.10]. Vessels adjacent to or in the cen-
ter of the necrotic zone showed fibrinoid necrosis of the walls and thrombotic occlusion of lumina with perivascular infiltrates of histiocytic macrophages, lymphocytes and nuclear debris [Fig.11]. Polymorphs and plasma cells were inconspicuous. Histiocytic granulomatous aggregates were present in some foci [Fig.12]. Vasculitic changes in the periphery of the lesion appeared more chronic with a perivascular dense mantle of lymphocytes and monohistiocytic cells without any noticable filirloid necrosis [Fig.13]. Scattered epitheloid cells were noticed but not in sufficient numbers to constitute a definite palisade. Well defined epitheloid cell tubercles were not present in multiple level examined. There was no caseation and Ziehl Neelsen staining did not reveal any acid fast bacilli. The histopathologic examination was consistent with the diagnosis of PNT.

Discussion

PNT is characterized by papules eruption affecting extremities and occurring in more or less symmetric crops. The papules are dusky red, symptomless and usually not itchy but were reported in one case to be severely itchy (7). The sites involved in PNT are the dorsum of hands and feet, the elbows, the knees, the palms, the soles, the buttocks and perineum, the glans peris was reported to be affected (8,9), and pernio areas may be favoured (10). PNT affects patients who have strongly positive tuberculin test which may even show large bullae (7).

The diagnosis of PNT in the present case is based on the clinical presentation, the history of contact with a family case of active pulmonary tuberculosis and the highly positive tuberculin test. It was reported that mycobacterial tuberculosis DNA has been detected by PCR in 77.8% of PNT specimens (11). Wilson-Jones and Winkelmann believe that PNT presents a characteristic clinical and histopathological pattern and that a special granulomatous type of small-vessel vasculitis explains the unique pathological features of the disease. Immunohisto-chemical labeling of biopsy specimens (13), revealed a preponderance of T-lymphocytes, monocytes-macrophages and Largerhans cells indicating type IV hypersensitivity reaction. B lymphocytes were sparse. Conditions that may be confused with PNT on clinical or histopathological ground include perforating callagenosis, perforating granuloma annulare, necrotizing or septic vasculitis, Churg-Strauss granuloma, populeonecrotic syphilid, pityriasis lichenoides chronica, lymphomatoid populus, popular articularia, miliary tuberculosis, chordodermitatis nodularis, suppurative falkculities and infectious causes of palisading granulomas (12,13,14).

In the 12 cases described by Wilson-Jones and Winkelmann (12), the age of the patients varied between 7 years and 30 years. The females predominated (9 of 12) with a female to male ration 3:1 and tuberculin test was positive in 100% of cases. The number of locations affected in each patient varied from one to six with a mean of 3. The principal sites affected were the hands (8 of 12); the legs (5 of 12); the feet and buttocks (4 of 12); the knees, elbows, thighs and arms (3 of 12); fingers, trunk and ears (1 of 12). The duration of the lesion varied from 5 weeks to 15 years.

Our patient, a female 48 years old had the PNT on hands and feet and she winter and improves in summer. We provisionally thought of perniosis and the investigations done were negative to cryoglobulins, hepatitis B, C & A, rheumatoid factor, ANA, and anti DNA antibodies. She had raised ESR and highly positive Mantoux test. Wilson-Jones and Winkelmann (12) thought that one of their patients had chilblains and 6 of the 7 patients they tested for ESR had raised ESR readings which became normal after responding to anti tuberculous therapy in 3 to 12 weeks and they suggested that ESR could be useful to monitor the effect of treatment. The influence of cold and the predilection of PNT to perniotic areas was stressed (10).

PNT was reported to evolve in lupus vulgaris (15), or may be associated with erythema induration (12,15,16). Excutaneous tuberculosis was found in 30-40% of cases with high incidence of nodal affection (15,17). Therefore in patients with PNT it is important to search for any associated tuberculous in lymph nodes, lungs and bones (12).

Some consider PNT the most common tuberculid (18). Tuberculids in general are treated with antituberculous drugs (1), treatment is indicated even in absence of M. tuberculosis in histologic specimens or culture of skin lesion (19). Triple agent chemotherapy (rifampicin 900mg/d, isonatzed 400 mg/d and ethambutol 1200 mg/d), could lead to remission in two months with mainenance of treatment for 12 months (20). Long term triple regimen of treatment is recommended for PNT (21).
REFERENCE