Cutaneous tuberculosis and phlyctenular keratoconjunctivitis: A forgotten association

Archana Singal, Puneet Aggarwal, Deepika Pandhi, Jolly Rohatgi
Departments of Dermatology and STD and *Ophthalmology, University College of Medical Sciences and Guru Teg Bahadur Hospital, Delhi - 110 095, India.

Address for correspondence: Dr. Deepika Pandhi, B- 1/1101, Vasant Kunj, Delhi - 110 070, India.
E-mail: deepikapandhi@rediffmail.com

ABSTRACT

Cutaneous tuberculosis may be associated with concurrent systemic foci in the body such as lung, lymph node, bone or CNS. Phlyctenular keratoconjunctivitis (PKC) is a manifestation of immunological response to a variety of antigens in the eye, tubercular focus (evident or occult) being the commonest in India. Reports in the existing literature have shown lungs and lymph nodes to be the predominant underlying focus associated with PKC, whereas cutaneous tuberculosis has seldom been found in this situation. We report this forgotten association in two children with cutaneous tuberculosis, one each with lupus vulgaris and scrofuloderma, who also had PKC. Interestingly, one of the cases also had simultaneous lichen scrofulosorum, which is also an immunological response to tubercular antigen and manifests in the skin, thus showing immunological manifestation in two different organ systems along with cutaneous focus of tuberculosis.

Key Words: Lupus vulgaris, Scrofuloderma, Phlycten, Cell-mediated immunity

INTRODUCTION

Extra-pulmonary tuberculosis constitutes 10% of total tuberculosis and 1.5% of these have cutaneous disease.[1,2] Phlyctenular keratoconjunctivitis (PKC) represents an allergic cell mediated response in cornea and/or conjunctiva to some antigen, to which they have already become sensitized.[3] In a series from India, tuberculosis accounted for up to 77% cases of PKC.[3] However, only a few published reports documenting the association of cutaneous tuberculosis and PKC exist in the current world literature.[4-6] The present report highlights this association in two children.

CASE REPORTS

Case 1
A six year-old-boy presented with a red raised lesion on the right cheek for 6 months and photophobia with increased lacrimation in the left eye for 15 days. He did not have any constitutional symptoms. Cutaneous examination revealed a single, erythematous plaque of 2 x 6 cm size over the right preauricular region, with an ulcer showing unhealthy granulation tissue and scarring [Figure 1] and two discrete, non-tender, right submandibular lymph nodes. Ophthalmologic examination revealed marked circumcorneal congestion and multiple gray nodules...
along the limbus in the left eye [Figure 2]. The left cornea showed one healed ulcer and multiple fascicular ulcers, with a leash of blood vessels extending to the limbus. There was marked blepharospasm, photophobia and excessive lacrimation. Visual acuity was 6/6 and 6/18 in the right and left eye, respectively.

Hemogram, liver and kidney function tests were unremarkable. Erythrocyte sedimentation rate was 38 mm in the 1st hour and Mantoux test (1 PPD) was 16 mm after 48 hours. Stool examination revealed no ova or cyst. Bacterial culture from lid margin and conjunctiva was sterile. Fine needle aspiration cytology (FNAC) from the skin lesion and lymph node showed granulomatous infiltrate comprising epithelioid cells and occasional acid fast bacilli (AFB), on Ziehl-Neelsen staining. Skin biopsy revealed epithelioid cell granuloma, Langhans’ giant cells and minimal caseation necrosis without AFB in the upper dermis, consistent with lupus vulgaris.

Culture for *Mycobacterium tuberculosis* on Lowenstein Jensen (LJ) and radiometric liquid culture medium BACTEC® 460 TB System (Becton Dickinson Diagnostic Instrument Systems Sparks, Maryland, USA), did not show any growth.

A diagnosis of lupus vulgaris with PKC was made and the patient was started on short course antitubercular therapy (ATT) with rifampicin (10 mg/kg/day), isoniazid (5 mg/kg/day), ethambutol (15 mg/kg/day) and pyrazinamide (25 mg/kg/day) for the initial 2 months, to be followed by rifampicin and isoniazid for 4 months. For eye lesions, topical drops (dexamethasone 0.01% and chloramphenicol 0.5%) twice daily, were prescribed. Follow up after 3 weeks showed complete resolution of eye lesions with residual nebulous corneal opacities. The skin lesion resolved after 2 months.

**Case 2**
A 5 year-old-girl complained of a nonhealing ulcer in the right groin, of 3 months duration. Her parents also noticed asymptomatic, grayish white patches in both eyes, since one month. Cutaneous examination revealed an ulcerated lesion with bluish undermined margins, adherent to underlying matted right inguinal lymph nodes. The patient also had grouped follicular and perifollicular, skin colored, pinhead sized papules on her trunk, suggestive of lichen scrofulosorum (LS). Ophthalmologic examination revealed severe photophobia, multiple phlyctens on cornea and limbus bilaterally, on right more than left and multiple nebular corneal opacities, suggestive of previous phlyctens. Visual acuity in right and left eye was 6/18 and 6/6, respectively.

Hematological investigations and stool examination were unremarkable. Bacterial culture from lid margin and conjunctiva was sterile. X-ray chest revealed multiple opacities in the left upper zone, suggestive of pulmonary tuberculosis. Mantoux test was 22 x 25 mm, with marked induration and vesiculation.
Phlyctenular keratoconjunctivitis is a nonspecific allergic response in the cornea and/or conjunctiva, to a variety of antigens. Conjunctival phlyctens are usually transient and asymptomatic, but occasionally in larger phlyctens, frank pustular conjunctivitis may develop with subsequent penetration into sclera, leading to permanent scar formation. Corneal phlyctens present with lacrimation, photophobia and blepharospasm and tend to leave opacities, leading to permanent vision impairment or occasionally blindness.[7]

DISCUSSION

Phlyctenular keratoconjunctivitis is a nonspecific allergic response in the cornea and/or conjunctiva, to a variety of antigens. Conjunctival phlyctens are usually transient and asymptomatic, but occasionally in larger phlyctens, frank pustular conjunctivitis may develop with subsequent penetration into sclera, leading to permanent scar formation. Corneal phlyctens present with lacrimation, photophobia and blepharospasm and tend to leave opacities, leading to permanent vision impairment or occasionally blindness.[7]

Tuberculosis as an etiological association, is being supplanted by staphylococcal infection and worm infestation in the developed world. Tuberculosis has been implicated in 77% (86/112) of all cases of PKC from India.[3] PKC lesions were observed to be more severe and recurrent in patients with tuberculosis. The most common underlying tubercular focus was found to be the lungs (56/86), followed by tubercular lymphadenopathy, but none had cutaneous disease.[3]

A tuberculid is a cutaneous cell mediated immunological response to a tubercular focus, often occult, elsewhere in the body. Lichen scrofulosorum is the commonest presentation.[2] Thus, both PKC and LS represent delayed hypersensitive immunological response to Mycobacterium tuberculosis in eyes and skin, respectively. This was observed in case 2, in the present report. PKC was considered to be a usual allergic phenomenon, attributed to primary tuberculosis in children over 50 years ago.[4] However, since then, only isolated cases along with papulonecrotic tuberculides have been reported.[5,6] Of our two cases, one had LS with scrofuloderma, while second had lupus vulgaris alone. Though PKC is most frequently reported in children, most series on pediatric tuberculosis fail to document this entity.[8,9]

The incidence of underlying systemic involvement with cutaneous tuberculosis has varied from 1.7 to 26.1%, with frequent involvement of lungs and lymph nodes.[8-10] Although most dermatologists diligently look for systemic disease, a detailed ophthalmologic evaluation is often overlooked. We suggest that a simple practice of ophthalmologic evaluation in all patients with cutaneous tuberculosis, will result in timely institution of local treatment of PKC and will subsequently ameliorate the unwanted complications of visual disability.

REFERENCES