## Metastatic tubercular abscess associated with bone and lymph node involvement

## Sir,

Tuberculosis is seldom considered an etiology of subcutaneous abscess in our daily practice. Here, we report a rare case of metastatic tubercular abscess in a malnourished girl, with bone involvement and lymphadenopathy.

A 13-year-old malnourished girl presented with multiple abscesses predominantly involving her upper and lower extremities for the last 2 years. She had multiple, variably sized, subcutaneous discharging abscesses on the upper extremity, lower extremity and neck [Figures 1 and 2]. The lesions were fluctuant and non-tender without any sign of inflammation. Matted, enlarged lymph nodes were noted in the cervical [Figure 3] and inguinal regions [Figure 4]. Scars were observed at the sites of previous drainage. She started developing these abscesses on her neck and extremities which progressively increased in size. They had been drained unsuccessfully, only to reappear again. She also received multiple courses of systemic antibiotics and antifungals with no appreciable improvement. There was no history of fever, cough and breathlessness at presentation. The past history revealed an episode of pulmonary tuberculosis, five years back, along with weight loss. Family history was unremarkable. The patient had not been immunized with bacillus Calmette-Guérin (BCG) vaccine. There was pallor, and malnutrition (body mass index 15.5 kg/m<sup>2</sup>); the remainder of the general examination was normal.

Routine blood parameters were within normal limits, except for anemia (hemoglobin 6.8 mg/dl). The patient was non-reactive for antibodies to human immunodeficiency virus 1 and 2. Chest skiagram revealed upper zone opacification and pleural effusion affecting the right lung, suggestive of a tubercular focus. Bone skiagram revealed an erosion of the medial aspect of the navicular and the medial cuneiform bone suggestive of tubercular osteomyelitis [Figure 5]. Examination of other systems did not reveal any abnormality.

Tuberculin skin test (Mantoux test) was positive ( $15 \text{ mm} \times 12 \text{ mm}$ ) with 5 TU purified protein derivative (PPD). Skin biopsy from the margin of abscess over the left foot showed a thickened epidermis with granulation tissue in the dermis along with a chronic inflammatory cell infiltrate and hemosiderin-laden macrophages. However, there was no granuloma, caseation necrosis or any features suggestive of a malignancy. Acid-fast bacilli were conspicuous in the discharge (Ehrlich–Ziehl–Neelsen stain) [Figure 6]. The patient

Access this article online	
Quick Response Code:	Website:
	www.ijavi.com DOI: 10.4103/0378.6323.188651
	10.4100/03/0-0020.100001

did not give consent for pus culture in Lowenstein–Jensen medium, despite repeated requests, because of economic constraints. Clinico-pathological correlation, positive tuberculin skin test and the past history of pulmonary tuberculosis led us to a final diagnosis of metastatic tubercular abscess. She was treated category 1 regimen of anti-tubercular drug therapy, after consultation with the department of pulmonary medicine, for 6 months. On follow-up after 5 months, ulcers had healed and lymphadenopathy resolved considerably [Figures 7-9], thus confirming our diagnosis retrospectively. Treatment was continued for 1 more month as per protocol.



Figure 1: Multiple, discharging subcutaneous abscesses on the right upper and lower limbs

This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 3.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as the author is credited and the new creations are licensed under the identical terms.

For reprints contact: reprints@medknow.com

How to cite this article: Sancheti K, Podder I, Saha M, Chowdhury SN, Bandyopadhyay D. Metastatic tubercular abscess associated with bone and lymph node involvement. Indian J Dermatol Venereol Leprol 2017;83:276.

Received: February, 2016. Accepted: April, 2016.



Figure 2: Necrotic ulcer on the right hand



Figure 4: Inguinal lymphadenopathy

Extrapulmonary tuberculosis is seen in about one-third of all tuberculosis cases with higher preponderance in children.<sup>1</sup> Cutaneous tuberculosis can have a myriad of clinical presentations.<sup>2</sup> Lesions occurring due to exogenous inoculation of *Mycobacterium tuberculosis* are tubercular chancre, tuberculosis verrucosa cutis and occasionally lupus vulgaris. Endogenous cutaneous infection may lead to scrofuloderma, acute military tuberculosis, tuberculous gumma, orificial tuberculosis and lupus vulgaris. In addition, there may be cutaneous immune reactions to *M. tuberculosis* called tuberculids. Lupus vulgaris has been reported to be the most common form of cutaneous tuberculosis, followed by scrofuloderma, tuberculous addition, 3

Metastatic tuberculous abscess occurs due to hematogenous dissemination from a primary focus during lowered immunity, resulting in single or multiple lesions. It is seen particularly in malnourished children or immunosuppressed patients, following local trauma, and in association with underlying lymphoma.<sup>4-6</sup> The extremities are more often affected than the trunk.<sup>2</sup>



Figure 3: Cervical lymphadenopathy



Figure 5: X-ray of the right foot showing erosion of the medial aspect of the navicular and the medial cuneiform bone, suggestive of tubercular osteomyelitis

Some notable differential diagnoses may be staphylococcal abscess, other mixed bacterial infections, sporotrichosis, leishmaniasis, atypical mycobacterial infections (predominantly *Mycobacterium ulcerans*), deep fungal infections, syphilitic gumma and leprosy. In our case, the past history of pulmonary tuberculosis, presence of tubercular focus on the chest X-ray and presence of acid-fast bacilli in Ehrlich–Ziehl–Neelsen stain helped to rule out the differentials.

The standard treatment regimen involves 2 months of 4-drugs therapy (isoniazid, rifampin, pyrazinamide and ethambutol) followed by 4 months of 2-drugs therapy (isoniazid and rifampin).<sup>7</sup> Various observations revealed the response of lupus vulgaris and tuberculosis verrucosa cutis to be alike with all the three regimens. The localized lesions subside completely after 4 months of therapy and the most extensive forms take 5 months to resolve. The patients with scrofuloderma responded similarly to both the triple drug regimens. The discharge, sinuses and ulcers cleared in 6 months, but the lymph nodes took longer to regress; up to 7 months in localized scrofuloderma and 9 months in more



**Figure 6:** Smear of discharge showing multiple acid-fast bacilli within a macrophage (marked by arrow) (Ehrlich–Ziehl–Neelsen, ×1000)



Figure 8: Healed cervical lymphadenopathy

widespread scrofuloderma.<sup>8</sup> In our case, the lesions improved considerably after five months.

Metastatic tuberculous abscesses are usually associated with involvement of the bones of hands and feet and lymphadenopathy.<sup>9,10</sup> Association has also been reported with polymyositis at different sites such as mediastinum, central nervous system and chest wall.<sup>11-13</sup> In our case also, lymphadenopathy (cervical and inguinal) was present, along with bony involvement of hands and feet.

Financial support and sponsorship Nil.

**Conflicts of interest** There are no conflicts of interest.

> Karan Sancheti, Indrashis Podder, Maitrayee Saha<sup>1</sup>, Satyendra Nath Chowdhury, Debabrata Bandyopadhyay



Figure 7: Healed lesion on the hand



Figure 9: Healed inguinal lymphadenopathy

Departments of Dermatology and <sup>1</sup>Pathology, Medical College and Hospital, Kolkata, West Bengal, India Correspondence: Dr. Indrashis Podder,

Block AD-71, Salt Lake City, Kolkata - 700 064, West Bengal, India. E-mail: ipodder88@gmail.com

## References

- Darmstadt GL, Sidbury R. The skin. In: Behrman E, Kliegman RM, Jenson HB, editors. Nelson Textbook of Paediatrics. 17<sup>th</sup> ed. Philadelphia: WB Saunders Company; 2004. p. 2153-250.
- Yates VM, Rook GA. Mycobacterial infections. In: Burns T, Breathnach S, Cox N, Griffiths C, editors. Rook's Textbook of Dermatology. 7<sup>th</sup> ed. London: Blackwell Science; 2004. p. 1427-67.
- Kumar B, Rai R, Kaur I, Sahoo B, Muralidhar S, Radotra BD. Childhood cutaneous tuberculosis: A study over 25 years from northern India. Int J Dermatol 2001;40:26-32.
- Maejima H, Arai S, Ebata T, Takeda H, Kusunoki M, Katsuoka K. Tuberculous gumma associated with idiopathic thrombocytopenic purpura: Report of a Japanese female patient. J Dermatol 2007;34:86-91.
- 5. Vidal D, Barnadas M, Pérez M, Coll P, Alomar A. Tuberculous gumma following venepuncture 2001;144:601-3.

- Kalaria VG, Kapila R, Schwartz RA. Tuberculous gumma (cutaneous metastatic tuberculous abscess) with underlying lymphoma. Cutis 2000;66:277-9.
- Bravo FG, Gotuzzo E. Cutaneous tuberculosis. Clin Dermatol 2007;25:173-80.
- Ramesh V, Misra RS, Saxena U, Mukherjee A. Comparative efficacy of drug regimens in skin tuberculosis. Clin Exp Dermatol 1991;16:106-9.
- Sezgin B, Atilganoglu U, Yigit O, Ergün SS, Cambaz N, Demirkesen C. Concomitant cutaneous metastatic tuberculous abscesses and multifocal skeletal tuberculosis. Indian J Dermatol 2008;53:149-53.
- Latsios D, Chloros D, Spyratos D, Dagdilelis L, Sichletidis L. Iliopsoas tuberculous abscess associated with cervical and axillary tuberculous lymphadenopathy. BMJ Case Rep 2011;2011. pii: Bcr0320114037.
- Silva GA, Motta RN, Carvalho Rde S, Lupi O, Azevedo MC, Ferry FR. Cutaneous tuberculous gummas in a patient with polymyositis. An Bras Dermatol 2013;88:98-101.
- 12. Pacheco C, Silva E, Miranda J, Duarte R. Cutaneous tuberculosis as metastatic tuberculous abscess. J Bras Pneumol 2015;41:200-2.
- Agrawal A, Shanthi V, Ramakrishna BA. Tubercular cerebellar abscess in a child. J Mahatma Gandhi Inst Med Sci 2015;20:77-8.