Forgotten but not gone - Scrofuloderma in a migrant student from India

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CASE REPORT

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Abstract

A 34-year-old Indian student who immigrated to Australia five years ago presented with a four-week history of neck pain. Physical examination revealed two firm fixed cervical lymph nodes in the anterior triangle and midline region which were tender on palpation and erythematous on inspection. Cording phenomenon was found on ZN staining of FNA sample and mycobacterium tuberculosis (*M.tb*) PCR confirmed the diagnosis with incomplete resistance to isoniazid. Patient was treated with other three first line antituberculosis medications for nine months with an excellent outcome. Prednisolone was also used as adjunctive therapy and tapered during the course of treatment.

Key Words

Scrofuloderma, Tuberculosis, Cording phenomenon

Implications for Practice

- Scrofuloderma is the terminology mostly forgotten due to efficient control of mycobacterium tuberculosis (MTB) in developed countries.
- 2. Immigrants from developing or third world countries may be infected with *M.tb* and inexperience of doctors in developed countries may lead to a delay in diagnosis, treatment and eventually spread of disease in the community.

Background

Scrofuloderma, also known as "Tuberculosis colliquativa cutis",¹ is a type of tuberculous skin involvement in children and young adults. There is breakdown of skin overlying a

tuberculosis focus in the lymph node, bone or joint. This condition begins as subcutaneous nodules which then suppurates and gradually causes sinuses, tracks and scars.²

Progression of the disease may lead to irregular fixed, dense and fibrotic masses; however it could be fluctuant and discharge.³

Other types of cutaneous tuberculosis such as tuberculosis verrucosa cutis, tuberculosis gumma and orificial tuberculosis are also defined in dermatology references.^{2, 4}

Diagnosis of this condition is usually with needle aspiration or excisional biopsy of the mass and the demonstration of acid-fast bacilli by microbiological staining. Mycobacterium tuberculosis (MTB) PCR has high specificity for diagnosis.^{5, 6} Cording phenomenon (Figure 2) is a typical finding in MTB infection but can also be seen in other mycobacterial infections.

Treatment of scrofuloderma is similar to pulmonary tuberculosis and the role of corticosteroids is still unclear. Corticosteroids are proven to be effective in tuberculous meningitis and tuberculous pericarditis with active constrictive pattern.⁷⁻⁹ Use of corticosteroids in patients with compromised airway secondary to enlarged lymph nodes may be helpful.

Case details

A 34-year-old male patient was referred to the infectious diseases outpatient clinic by his general practitioner with a four-week history of tender cervical lymphadenopathy.

He was a student from India who migrated to Australia five years ago. He denied previous contact with any patient diagnosed with tuberculosis (TB). He had no significant childhood history of lung diseases or TB and neither a family history of TB. He had his BCG vaccination as a child. Since his arrival in Australia, he had not travelled to other countries.

Physical examination revealed normal vital signs with no respiratory distress. Head and neck examination was consistent with two hard, tender and fixed masses in the



left anterior and mid cervical area (Figure 1). There was no evidence of generalised lymphadenopathy or abdominal organomegaly. His chest X-ray was also normal.

Figure 1: Cervical tender fixed masses



He underwent a fine needle aspiration (FNA) of his cervical lymph nodes sent for culture, microbiology and histopathology. Mycobacterial culture and PCR was also requested considering his background and recent travel from India.

Tuberculin Skin Test (TST) was performed which was positive at 48 hours. Ziehl-Neelsen stain performed on his FNA sample showed cording phenomenon (Figure 2) M.tb PCR confirmed the diagnosis of Tuberculosis. Mycobacterium culture results incomplete showed resistance to Isoniazid with minimum inhibitory concentration (MIC) of more than 1 but less than 4 microgr/ml. Full susceptibility was reported to Rifampicin, Ethambutol and Pyrazinamide. Considering the excellent outcome of treatment in Isoniazid resistant tuberculosis, he was treated with three other first line drugs Rifampicin, Ethambutol, and Pyrazinamide for nine months.

Figure 2: Microscopy of the culture.



Figure 3: Lymphadenopathy has resolved.





After one month of treatment, his lymph nodes became more erythematous, tender and inflamed. High dose prednisolone was started with a diagnosis of paradoxical immune response. Prednisolone was later tapered off in term of treatment. After three months of treatment his lymph nodes disappeared with minor hyperpigmentation (Figure 3). He finished the total treatment course or nine months with no major complications and made a complete recovery.

References

1. Ramos-e-Silva M, Ribeiro de Castro MC. Section twelve: Mycobacterial infections. In: Rapini RP, Bolognia, JL, Jorizzo JL. Dermatology: 2-Volume Set. Mosby;2007.p.1221-43.

2. Tappeiner G. Tuberculosis and Infections with Atypical Mycobacteria. In: Wolff K, Goldsmith LA, Katz SI, Gilchrest BA, Paller AS, Leffell DJ. Fitzpatrick's Dermatology in General Medicine, 7th edn. New York: McGraw-Hill; 2008.p.1768-78.

3. Iftikhar U, Nadeem M, Aman S, Kazmi AH. Scrofuloderma: a common type of cutaneous tuberculosis. Journal of Pakistan Association of Dermatologists 2011;21:61-65.

4. Yates VM, Rook GAW. Mycobacterial infections. In: Burns T, Breathnach S, Cox N, Griffiths C, editors. Rook's Textbook of Dermatology, 7th edn. London: Blackwell Science; 2004. p. 28.1-28.39.

5. Rajakumar D, Rosenberg AM. Mycobacterium tuberculosis monoarthritis in a child. Pediatr Rheumatol 2008; 6: 2-10.

6. Tan WP, Tang MBY, Tan HH. Scrofuloderma from the acromioclavicular joint presenting as a chronic ulcer in an immunocompetent host. Singapore Med J. 2007 Sep;48(9):e243-5.

7. Thwaites GE, Nguyen DB, Nguyen HD, Hoang TQ, Do TT, Nguyen TC, Nguyen QH, Nguyen TT, Nguyen NH, Nguyen TN, Nguyen NL, Nguyen HD, Vu NT, Cao HH, Tran TH, Pham PM, Nguyen TD, Stepniewska K, White NJ, Tran TH, Farrar JJ.. Dexamethasone for the treatment of tuberculous meningitis in adolescents and adults. N Engl J Med. 2004 Oct 21;351(17):1741-51.

8. Schoeman JF, Van Zyl LE, Laubscher JA, Donald PR. Effect of corticosteroids on intracranial pressure, computed tomographic findings, and clinical outcome in young children with tuberculous meningitis. Pediatrics. 1997 Feb;99(2):226-31.

9. Dooley DP, Carpenter JL, Rademacher S. Adjunctive corticosteroid therapy for tuberculosis: a critical reappraisal of the literature. Clin Infect Dis. 1997Oct;25(4):872-87.

CONFLICTS OF INTEREST

The authors declare that they have no competing interests. We also declare that all the authors have approved the final version of this manuscript.