Tracheo-bronchial involvement in Sweet syndrome

A 70-year-old man with myeloﬁbrosis and Sweet syndrome (acute febrile neutrophilic dermatosis) presented with a 2 month history of cough with bloody sputum (ﬁgure 1). No inﬁltration was visible on a chest radiograph.

On bronchoscopic examination, a centrally depressed macular lesion with an eroded surface was revealed in the anterior tracheal wall, as well as several scattered erosions in the trachea and bronchi (ﬁgure 2A). A biopsy specimen of the tracheal lesion showed dense inﬂammatory inﬁltrates with intercellular oedema and some leucocytoclasis. Despite dense inﬂammatory inﬁltrates mainly consisting of neutrophils, there was no evidence of vasculitic changes (ﬁgure 3A). These histological characteristics of the tracheal lesion were similar to those of skin lesions in Sweet syndrome (ﬁgure 3B). Aetiological investigations yielded negative results, but the anti-Chlamydia pneumoniae immunoglobulin (IgM) titre was elevated. Symptoms remained aggravated despite azithromycin treatment. Oral prednisolone at 50 mg daily was started and then tapered. The patient’s symptoms decreased and the lesions showed scarring on bronchoscopic examination following this treatment (ﬁgure 2B).

Sweet syndrome is often associated with haematopoietic malignancy and myeloproliferative disorders such as myeloﬁbrosis.1

Figure 1  A well-deﬁned erythematous plaque on the dorsal surface of the hand.

Figure 2  Images from bronchoscopic examination before (A) and after (B) corticosteroid treatment.

Figure 3  (A) A bronchoscopic biopsy specimen of the tracheal lesion showing dense inﬂammatory inﬁltrates without vasculitic changes. (B) A biopsy specimen of the skin lesion showing the same characteristics.
Its pathogenesis is not definitively determined but it is postulated to be a multifactorial process involving a hypersensitivity reaction and cytokine dysregulation. Sweet syndrome can include internal organ involvement. In the literature, there is only one report of bronchial involvement, but that was associated with pulmonary infiltration. In the present case, tracheo-bronchial involvement without pulmonary infiltration may have been induced locally by chlamydophilal tracheobronchitis.

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Competing interests None.

Patient consent Obtained.

Provenance and peer review Not commissioned; externally peer reviewed.

Accepted 25 May 2010
Published Online First 6 September 2010


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