Strawberry gums: a clinicopathological manifestation diagnostic of Wegener’s granulomatosis?

S S Napier, J A Allen, C R Irwin, D R McCluskey

Abstract

Aims—To highlight an uncommon but characteristic gingival lesion associated with Wegener’s granulomatosis, emphasising the presence of pseudoepltheliomatous hyperplasia, microabscesses, and multinucleate giant cells; and the paucity of the currently accepted histopathological criteria of Wegener’s granulomatosis—namely necrosis, vasculitis, and granuloma.

Methods—The histopathological features of a gingival biopsy specimen from a case of Wegener’s granulomatosis limited to the upper aerodigestive tract in a 36 year old woman were compared with those of 14 similar reported cases.

Results—Pseudoepltheliomatous hyperplasia, microabscesses, and multinucleate giant cells were recorded as present in almost all cases of gingival Wegener’s granulomatosis. Necrosis, vasculitis, and granuloma formation were present in only a few cases.

Conclusions—The combination of pseudoepltheliomatous hyperplasia, microabscesses, and multinucleate giant cells in a gingival biopsy specimen from erythematous, swollen gums, clinically resembling over-ripe strawberries, in a patient with a severe systemic upset, is so typical of Wegener’s granulomatosis as to be virtually diagnostic. Recognition of this characteristic lesion may aid early diagnosis and treatment in cases where other diagnostic variables are absent.


Prompt definitive diagnosis is important in Wegener’s granulomatosis: the prognosis has been greatly improved by early instigation of appropriate treatment.1 Serological tests can be of help, especially the identification of high cytoplasmic staining anti-neutrophil cytoplasmic antibody (cANCA) titres. But these may be negative in 15–33% of cases, particularly those without renal disease.2 Current histopathological criteria for the diagnosis of Wegener’s granulomatosis are stringent, requiring identification of vasculitis, ill defined granuloma, multinucleate giant cells, and necrosis,3 4 although some authors maintain that the patterns of the necrosis are sufficiently characteristic to allow a diagnosis to be made from this feature alone.5 6 The lesions may be sparsely distributed through the tissues and are frequently not identified in small biopsy specimens, even in those from sites of active disease in the nose, lung, or kidney.

It is worth emphasising that Wegener’s granulomatosis cannot be diagnosed by the clinician or pathologist in isolation. It requires good interaction between clinical and laboratory disciplines, and careful correlation of the results of all available investigations with the clinical presentation.7

We draw attention to histopathological features, which have been consistently described, though perhaps not widely recognised, that occur in conjunction with a highly characteristic gingival lesion in Wegener’s granulomatosis. To the best of our knowledge, this particular combination of clinical and histopathological features, when seen against the background of a systemic upset, has not been described before in association with any condition other than Wegener’s granulomatosis.

Case report

A 36 year old woman presented to our unit complaining of a sore mouth and malaise of three weeks duration. Examination principally showed extremely swollen, erythematous gingivae, particularly in the upper and lower premolar regions, where there were natural teeth in good condition. The gums had a granular appearance and were flecked with yellow, resembling over-ripe strawberries (fig 1). The favoured clinical diagnosis was gingival Wegener’s granulomatosis. An incisional biopsy of affected gum was performed.

The gingival biopsy specimen showed extensive pseudoepltheliomatous hyperplasia, with the stratified squamous epithelium extending into the stroma as deeply penetrating tongues and crypts. Within the oedematous connective tissues, there was an intense inflammatory infiltrate composed predominantly of neutrophils and eosinophils, with lesser numbers of plasma cells, lymphocytes, and macrophages. Focal clustering of neutrophils and eosinophils was noted, with microabscess formation. One of these microabscesses was seen rupturing into an epithelial crypt, and draining to the surface as a bead of pus. Several isolated multinucleate giant cells were present, but no necrosis, vasculitis, or granuloma formation were identified (figs 2–4). Special stains failed to reveal fungi. The pattern and degree of epithelial hyperplasia, the intensity of the inflammatory...
infiltrate and its tendency to cluster into microabscesses, together with the presence of several multinucleate giant cells were felt to be strongly suggestive of gingival Wegener’s granulomatosis.

The patient was admitted to hospital to assess the extent of the disease. Examination of ear, nose, and throat, sinus and chest radiographs, and serological tests (including full blood picture, erythrocyte sedimentation rate (ESR), autoantibody screen, urinalysis and cANCA titres) were performed. Except for an ESR, which on two occasions was greater than 90 mm first hour, and a C-reactive protein concentration of 37.1 mg/ml (normal adult range under 6.0 mg/ml), all other investigations were normal or negative. A granular area was identified on the left side of the nasal septum, from which the patient had complained of an episode of epistaxis. Although not biopsied, this area was felt to be “consistent with Wegener’s granulomatosis” on clinical grounds.

Wegener’s granulomatosis limited to the upper aerodigestive tract was diagnosed, and in an effort to avoid the side effects of immunosuppressive treatment, a course of co-trimoxazole was started. There was clinical improvement within three days and complete resolution within one week, the ESR and C-reactive protein concentrations returning to normal values in line with the clinical condition.  

Discussion

After correlation of all positive findings Wegener’s granulomatosis was diagnosed in our case on the basis of three key features: (1) the multiple gingival lesions, which resembled over-ripe strawberries; (2) the patient’s complaint of malaise (supported by increased ESR and C-reactive protein concentration); and (3) the histopathological appearances in the gingival biopsy specimen. The serological tests only helped to reinforce the clinical impression of a systemic inflammatory process and did not confirm the diagnosis of Wegener’s granulomatosis. The cANCA titre, arguably the most useful investigation in cases of Wegener’s granulomatosis, was non-contributory. Even the presence of the nasal lesion could not be considered diagnostic.

This particular combination of clinical and histological features was first described in a peculiar form of progressive hyperplastic gingivitis by Milner, when the association with Wegener’s granulomatosis was made only at necropsy. Small numbers of similar cases have been reported since, most of which draw attention to the delay between initial presentation and diagnosis. These reports describe the histological features of pseudoeplitheliomatous hyperplasia, polymorph microabscesses, and giant cells in the gingival biopsy (table), the appearances being generally regarded as those of non-specific active chronic inflammation. The classic criteria of vasculitis, granulomata, and necrosis are

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PEH pseudoeplitheliomatous hyperplasia; + present; – absent; NS not specified. (In references 11 and 14 no specific details on the histopathological findings were reported).
notably absent in most gingival biopsy specimens, although vascular changes consistent with vasculitis,17 19 20 an inflammatory process described as “granulomatous”,10 15 16 and small areas of necrosis19 16 19 21 have been recorded. Often, only after disease progression with spread to “classical” sites or repeated biopsy is the diagnosis of Wegener’s granulomatosis established.

Although generally regarded as non-specific, the pseudoeosinophilic hyperplasia, the intense infiltrate of neutrophils and eosinophils with the formation of microabcesses, and the presence of multinucleate giant cells are seen in few other conditions affecting the gums. The degree of epithelial proliferation may suggest a squamous cell carcinoma, but the overall pattern is more that of a reactive rather than a neoplastic process. Fungal infections—for example, histoplasmosis and paracoccidioidomycosis (South American blastomycosis)—can usually be excluded by simple histochemical stains, in that the fungi may be identified either within the tissues or the giant cells with periodic acid Schiff or methylamine silver stains. Conditions such as orofacial granulomatosis, Crohn’s disease, and sarcoidosis, which may affect the gums and in which granulomata and multinucleate giant cells may occur, are quite distinct from Wegener’s granulomatosis in that they do not show polymorph infiltration, microabcess formation, or the same degree of pseudoeosinophilic hyperplasia.25 Eosinophilic granuloma (Langerhans’ cell histiocytosis) may produce red, swollen, painful gingivae. Eosinophil microabcesses, necrosis, and multinucleate giant cells are seen in this condition, but the striking histological feature is the dense infiltrate of the Langerhans’ cells—pseudoeosinophilic hyperplasia is not a feature.26

Most authors acknowledge that the clinico-pathological complex of “strawberry gums” plus the accompanying histopathological features of pseudoeosinophilic hyperplasia, microabcesses, and multinucleate giant cells is “highly suggestive” of Wegener’s granulomatosis.14-16 19 20 23 24 Indeed, we know of no association of the features of this complex with any other disease process. We feel that in an appropriate clinical setting they are so characteristic of gingival Wegener’s granulomatosis as to be virtually diagnostic, particularly since the classic criteria of vasculitis, granulomata, and necrosis occur only rarely in gingival biopsy specimens.

The diagnosis of a condition as rare, as variable in both its clinical and histopathological features, and as serious as Wegener’s granulomatosis will always require good clinico-pathological correlation, requiring more specific histological features to be identified to establish the diagnosis the less clinical support there is for Wegener’s granulomatosis.27 We believe that recognition of these lesions as part of the spectrum of Wegener’s granulomatosis may speed diagnosis, particularly those cases where other diagnostic variables are likely to be absent.

7 Colby TV, Tazelaar HD, Specks U, DeRemee RA. Nasal
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