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Eosinophilic Pustular Folliculitis Involving Labial Mucosa, Which Improved with Naproxen

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Dear Editor:

Eosinophilic pustular folliculitis (EPF) is a sterile eosinophilic infiltration of hair follicles. There are three variants, including the classic, immunosuppression-associated, and infantile type¹. Although they are histologically indistinguishable from each other, some propose that the three types be regarded as different disease entities due to their different responses to treatment.

Histologically, the most diagnostic feature of EPF is the infiltration of eosinophils in hair follicles and perifollicular areas¹. However, the term 'folliculitis' is being challenged², on account of the fact that approximately 20% of patients with classic EPF have the disease on palms and soles¹.

Here, we report a case of EPF involving the labial mucosa, which improved with naproxen. An 11 year-old girl presented with itching erythematous plaques and clustering pustules on the lateral side of the nose and perioral area, with erosive lesions on the external lips (Fig. 1A). A biopsy was taken from the lateral side of the nose at a district hospital. The lesion at nose showed the same feature and connected with labial lesion. It demonstrated follicular and perifollicular infiltration by eosinophils and

other inflammatory cells (Fig. 2). The diagnosis of classic EPF was made at the hospital, and she was treated with systemic prednisolone, cyclosporine, dapsone, and/or topical corticosteroid, pimecrolimus. However, her symptoms waxed and waned over 6 months and showed improvement only with the systemic prednisolone. Routine blood test was within normal limits without eosinophilia. Because pruritus was the prominent symptom rather than pain and tenderness, infectious condition was less suspected. Suspecting EPF, we started her on naproxen, 250 mg twice a day; and after three months, her symptoms greatly improved (Fig. 1B).

The etiology and pathogenesis of EPF have still yet to be fully elucidated, and there are multiple treatment options. The utilization and efficacy of therapies seem to depend on the type of EPF². Topical corticosteroids tend to be the first choice in all EPF variants¹. In children, topical calcineurin inhibitors and oral antihistamines are also usually effective and are viewed as the first-line agents³. However, our case was recalcitrant to various treatments, including topical corticosteroids and topical calcineurin inhibitor. Only the naproxen had a remarkable effect.

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Fig. 1. (A) Erythematous plaques and clustering pustules on both lateral side of the nose with central clearing and peripheral extension, involving lips. (B) Greatly improved state after treatment with naproxen.

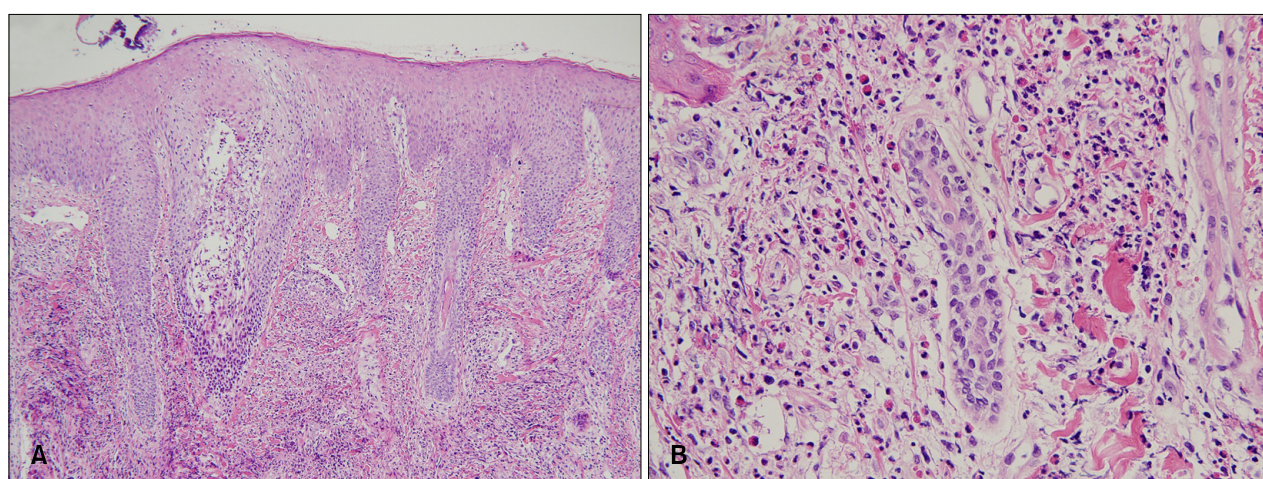


Fig. 2. (A) Inflammatory cell infiltration into hair follicle and around perifollicular area (H&E, ×100), (B) perifollicular eosinophilic and neutrophilic infiltration (H&E, ×400).

Nonsteroidal anti-inflammatory drugs (NSAID), particularly indomethacin, have been suggested as the treatment of choice in classic EPF². There are few reports dealing with naproxen, a type of NSAID widely being used, on EPF^{4,5}. We have reported the effects of naproxen as a first-line option on classic EPF in a Korean literature⁴. It showed 69% (11/16) of complete response rate in adult cases. This report implies that naproxen may also be safely used in children.

There has been only one case in a German literature, which involved oral mucosa in classic EPF. Our case had the labial lesions mainly on the external surface, which is

a keratinized stratified squamous epithelium rather than a true mucosa. But as there are no follicles in the external lip, as in the palms and soles, it is more likely that this disease may not be a real follicular disorder at all.

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Myopericytoma of the Facial Cheek

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Dear Editor:

A 44-year-old woman presented with a painful, solitary nodule on her right cheek. She noticed the flesh-colored nodule several years prior, and the nodule expanded slowly in size. There was no history of trauma. The patient's past medical history and family history were unremarkable. Upon physical examination, a 0.5×0.7 cm-sized, skin-colored, firm nodule on the right cheek was seen (Fig. 1). The biopsy specimen revealed a concentric perivascular proliferation of blank, spindle-shaped myoid-appearing cells (Fig. 2A, B). Immunohistochemical stain showed diffuse immunoreactivity on smooth muscle actin and was negative for desmin (Fig. 2C, D). The CD34 stain highlighted only the endothelium of the vessel, but the perivascular concentric myoid tumor cells were not immunoreactive (Fig. 2E). From the clinicopathological findings, the diagnosis of myopericytoma (MPC) was made. MPC is a rare, recently delineated tumor that originates from the perivascular myoid cells¹. It was described by Granter et al.² in 1998 and newly entered into a subgroup of perivascular tumors in the World Health Organization classification of soft tissue tumors³. MPCs are well-circum-

scribed and composed of a mixture of solid cellular areas intermixed with variable numbers of vascular channels. The latter are often elongated and display prominent branching, resulting in a stag-horn appearance. The cells in the solid areas are round, or short and spindle shaped with eosinophilic cytoplasm and vesicular nuclei. The presence of concentric layers of tumor cells around vascular channels resulting in a typical onion ring appearance is a hall mark of this tumor². However, MPC tumors exhibits a broad spectrum of growth patterns: solid classic, hemangiopericytoma-like, angioleiomyoma-like, hypocellular fibroma-like, solitary fibrous tumor-like, glomus tumor-like, cellular immature, intravascular and malignant subtypes. The present case was classified as angioleiomyoma-like pattern. By immunohistochemistry, the neoplastic cells



Fig. 1. A 0.5×0.7 cm sized, skin-colored, subcutaneous nodule of the right cheek.

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