Acute Localized Exanthematous Pustulosis Caused by Flurbiprofen

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To the Editor:
Acute generalized exanthematous pustulosis (AGEP) is an acute skin reaction that is characterized by generalized, nonfollicular, pinhead-sized, sterile pustules on an erythematous and edematous background. The eruption can be accompanied by fever and neutrophilic leukocytosis. Skin symptoms arise quickly (within a few hours), most commonly following drug administration. The medications most frequently responsible are beta-lactam antibiotics, macrolides, calcium channel blockers, and antimalarials. Pustules spontaneously resolve in 15 days and generalized desquamation occurs approximately 2 weeks later. The estimated incidence rate of AGEP is approximately 1 to 5 cases per million per year. Acute localized exanthematous pustulosis (ALEP) is a less common form of AGEP. We report a case of ALEP localized on the face that was caused by flurbiprofen, a propionic acid derivative from the family of nonsteroidal anti-inflammatory drugs (NSAIDs).

A 40-year-old woman was referred to the dermatology department due to the sudden onset of multiple pustules on the face. One week earlier she started oral flurbiprofen (8.75 mg daily) for a sore throat. After 3 days of therapy, multiple pruritic, erythematous and edematous lesions appeared abruptly on the face with associated multiple small nonfollicular pustules. At presentation the patient was febrile (temperature, 38.2°C) and presented with bilateral ocular edema and superficial small nonfollicular pustules on an erythematous background over the face, scalp, and oral mucosa (Figure 1). The rest of the body was not involved. The patient denied prior adverse reactions to other drugs. The white blood cell count was 15,000/µL (reference range, 4500–11,000/µL), with an increased neutrophil count (12,000/µL [reference range, 1800–7800/µL]). The erythrocyte sedimentation rate and C-reactive protein level was elevated (erythrocyte sedimentation rate, 53 mm/h [reference range, 0–20 mm/h]; C-reactive protein, 98 mg/dL [reference range, 0–5 mg/dL]). Bacterial and fungal cultures of skin lesions were negative. The results of a viral polymerase chain reaction analysis proved the absence of varicella-zoster virus or herpes simplex virus. Histopathology of a skin biopsy specimen showed subcorneal pustules composed of neutrophils and...
eosinophils, epidermal spongiosis, some necrotic keratinocytes, vacuolization of the basal layer, papillary edema, and a perivascular neutrophil and lymphocyte infiltrate (Figure 2). A leukocytoclastic infiltrate within and around the walls of blood vessels at the superficial level of the dermis and red cell extravasation in the epidermis was present. She discontinued use of flurbiprofen and was treated with a systemic corticosteroid (methylprednisolone 0.5 mg/kg daily). The pustules rapidly resolved within 7 days after discontinuation of flurbiprofen and were followed by transient scaling and discrete residual hyperpigmentation.

Acute localized exanthematous pustulosis is a less common form of a pustular drug eruption in which lesions are consistent with AGEP but typically are localized to the face, neck, or chest. The definition of ALEP was introduced by Prange et al1 to describe a woman who was diagnosed with a localized pustular eruption on the face without a generalized distribution as in AGEP. In the past, this localized eruption was described under different names (eg, localized pustular eruption, localized toxin follicular pustuloderma, nongeneralized acute exanthematic pustulosis).2-5 According to a PubMed search of articles indexed for MEDLINE using the terms localized pustulosis, localized pustular eruption, and localized pustuloderma, only 16 separate cases of ALEP have been documented since the report by Prange et al.1 The medications most frequently responsible are antibiotics. Three cases developed following administration of amoxicillin2-5; 2 cases of amoxicillin–clavulanic acid2-5; 1 of penicillin1; 1 of azithromycin5; 1 of levofloxacin10; and 1 of combination of cephalosporin, sulfamethoxazole-trimethoprim, and vancomycin.11 Other nonantibiotic causative drugs include sulfamethoxazole-trimethoprim,12 infliximab,13 sorafenib,14 docetaxel,15 finasteride,16 ibuprofen,17 and paracetamol.18 In reported cases, the lesions are consistent with the characteristics of AGEP both clinically and histopathologically but are localized typically to the face, neck, or chest. In the majority of patients with ALEP, the absence of fever has been observed, but it does not appear distinctive for diagnosis. Our patient represents another case of ALEP with flurbiprofen as the causative drug. The close relationship between the administration of the drug and the development of the pustules, the rapid acute resolution as soon as treatment was interrupted, and the histologic findings all supported the diagnosis of ALEP following administration of flurbiprofen. This NSAID—2-fluoro-α-methyl-(1,1'-biphenyl)-4-acetic acid—is a prostaglandin synthetase inhibitor with anti-inflammatory activity. It is a propionic acid derivative that is similar to ibuprofen, which was once involved in the occurrence of ALEP.17 In 2009, Rastogi et al17 reported a case of a 64-year-old woman with an acute outbreak of multiple pustular lesions and underlying erythema affecting the cheeks and chin without fever who had been taking ibuprofen for a toothache. The case is similar to ours and confirms that NSAIDs can induce ALEP. Compared with other NSAIDs, propionic acid derivatives are usually well tolerated and serious adverse reactions rarely have been documented.19

The physiopathologic mechanisms of ALEP are unknown but likely are similar to AGEP. The demonstration of drug-specific positive patch test
responses and in vitro lymphocyte proliferative responses in patients with a history of AGEP strongly suggests that this adverse cutaneous reaction occurs via a drug-specific T cell-mediated process.\textsuperscript{20}

Further study is needed to understand the etiopathogenesis of the localized form of the disease and to facilitate a correct diagnosis of this rare disorder.

REFERENCES