Eruptive Pseudoangiomatosis: Study of 7 Cases

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Abstract. Introduction. Eruptive pseudoangiomatosis consists in the acute development of small vascular lesions in the face and extremities that resolve in several weeks without scarring. Lesions are described as 3-4 mm asymptomatic macules and papules with peripheral whitish halo that blanch upon pressure. Initially it was considered a disease limited to children but it has also been described in adults. It overlaps with the entity known in Japan as «erythema punctatum Higuchi», possibly caused by an insect named Culex pipiens pallens.

Methods. We report a serie of 7 patients that consulted for lesions compatible with eruptive pseudoangiomatosis. We performed a detailed clinical history and histological, microbiological and serological studies. Follow-up time was up to 4 years.

Results. Eighty-five percent of patients were women and the mean age was 62 years. All cases appeared in spring/summer and 71 % relapsed. Lesions predominated in the face and extremities and the outbreak lasted 2-4 weeks. The anamnesis did not disclose any specific etiologic agent in any of the cases. Complete laboratory tests including serologies and cultures were negative or within normal limits. Histological study revealed vascular dilatation in all cases with endothelial cell protrusion and a peripheral lymphohistiocytic infiltrate.

Conclusions. Currently, the etiology of this entity is not well established although it probably represents a reactive disorder to different etiologic processes.

Key words: eruptive pseudoangiomatosis, erythema punctatum Higuchi, Culex pipiens pallens.

Introduction

Eruptive pseudoangiomatosis is characterized by outbreaks of erythematous, maculopapular lesions measuring 2 to 4 mm in diameter. They look like normal
angiomas and are generally located on the face and limbs. They blanch when pressed and many of them, particularly those on the arms or legs, are surrounded by a light-colored halo measuring 1 to 4 mm in diameter. They are asymptomatic or slightly pruritic, and resolve without residual scarring in a period of 2 to 18 days in children and 1 to 3 months in adults. The eruptions may recur.

Histological findings are unremarkable although there may be noticeable capillary dilation with plump endothelial cells protruding into the lumen. The affected vessel is normally surrounded by a predominantly lymphohistiocytic infiltrate of varying intensity. There is no vasculitis or vascular proliferation and the epidermis is generally unaffected.

Differential diagnosis should be based on the appearance of the lesions, clinical signs, and histological findings. It should include, among others, cherry hemangiomas, telangiectasias, and bacillary angiomatosis, and eruptive angiomatosis.

We describe our experience with eruptive angiomatosis and present a brief review of the few articles that discuss this disorder.

Materials and Methods

We present a series of 7 patients who visited our department in the last 10 years with a clinical picture compatible with eruptive pseudoangiomatosis. We obtained a detailed clinical history and performed a histological study in all cases. Complementary tests (routine laboratory and serological tests and cultures) and clinical follow-up were performed in most patients.

Results

The series included 6 women (86%) and 1 man (14 %), with a mean age of 62 years (Table 1). All the patients had

<table>
<thead>
<tr>
<th>Time</th>
<th>Age, y</th>
<th>Sex</th>
<th>Lesion Site</th>
<th>Duration</th>
<th>Triggering Agent</th>
<th>Recurrence</th>
<th>Laboratory Tests</th>
<th>Biopsy</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1996-1999</td>
<td>52</td>
<td>F</td>
<td>Face, upper limbs, lower limbs</td>
<td>2-3 weeks</td>
<td>Not related</td>
<td>Yes (several) spring/summer</td>
<td>Normal (biochemistry, profiles, CBC and differential WBC count, ANA, anti-dsDNA, protein profile, C3, C4)</td>
</tr>
<tr>
<td>2</td>
<td>2000-2002</td>
<td>48</td>
<td>F</td>
<td>Face, sternal region, upper limbs</td>
<td>2 weeks</td>
<td>Not related</td>
<td>Improved with change in normal place of residence</td>
<td>Yes (several) spring/summer</td>
</tr>
<tr>
<td>3</td>
<td>2001-2005</td>
<td>38</td>
<td>M</td>
<td>Face, upper limbs</td>
<td>15-20 days</td>
<td>Not related</td>
<td>Yes, in spring</td>
<td>Normal (biochemistry, profiles, CBC and differential WBC count) ANA (–) ASO (–) CMV (–), EPV (–)</td>
</tr>
<tr>
<td>4</td>
<td>2002</td>
<td>77</td>
<td>F</td>
<td>Face, upper limbs</td>
<td>2 weeks</td>
<td>Mosquito bites Affected neighborhood</td>
<td>None</td>
<td>Not done</td>
</tr>
<tr>
<td>5</td>
<td>2005</td>
<td>73</td>
<td>F</td>
<td>Mostly lower limbs</td>
<td>2-4 weeks</td>
<td>Not related</td>
<td>Yes (3 episodes)</td>
<td>Normal (biochemistry, profiles, CBC and differential WBC count) Non-specific parvovirus</td>
</tr>
<tr>
<td>6</td>
<td>2005</td>
<td>68</td>
<td>F</td>
<td>Face, upper limbs</td>
<td>2 weeks</td>
<td>Not related</td>
<td>Yes, in summer (2 episodes)</td>
<td>None</td>
</tr>
<tr>
<td>7</td>
<td>2005</td>
<td>78</td>
<td>F</td>
<td>Face, upper limbs</td>
<td>15-20 days</td>
<td>Not related</td>
<td>None</td>
<td>None</td>
</tr>
</tbody>
</table>

Abbreviations: ANA, antinuclear antibodies; ASO, antistreptolysin O titer; CBC, complete blood count; CMV, cytomegalovirus; ds, double-stranded; EBV, Epstein–Barr virus; WBC, white blood cell
their first visit between the months of May and August. Five patients (71%) reported recurrences, particularly in the spring and summer. Follow-up lasted for up to 4 years. Several patients did not return for check-up visits or undergo the complementary tests indicated, probably because their lesions resolved spontaneously. Patients 5, 6, and 7 only came once but we were able to contact 2 of them (patients 6 and 7) by telephone after several months to assess progress.

The main areas of the body affected were the face and limbs (Figure 1) and the outbreaks lasted between 2 and 4 weeks (average of 18 days). None of the clinical histories provided any clues to a possible cause; only 1 patient suggested an association with mosquito bites and mentioned that other people in her neighborhood also had these lesions, although we were unable to confirm this. None of the patients had associated infections or systemic diseases. The results of the routine laboratory tests (laboratory workup, general profiles, liver function, lipid profile, antinuclear antibodies, antidouble-stranded DNA, protein profile, complement levels, and antistreptolysin-O titers), serological tests (cytomegalovirus, Epstein–Barr virus, enterovirus, parvovirus B19, entamoeba histolytica, fasciola hepatica, Echinococcus granulosus, Rickettsia, and Borrelia), and cultures (skin biopsy and viral, bacterial, and fungal cultures) were within normal ranges for all the

![Figure 1](A, B y C). Angiomatous lesions with a whitish peripheral halo on the face and limbs of 2 different patients.

![Figure 2](Image). The dermis shows dilated thin-walled vessels with a protruding endothelium surrounded by a lymphocytic infiltrate (hematoxylin–eosin, ×100)

![Figure 3](Image). Image of characteristic vessel at greater magnification (hematoxylin–eosin, ×200)
patients that underwent the tests. Histological examination revealed vascular dilation in all cases, and the endothelial cells protruded slightly towards the lumen of the vessel, which was surrounded by a predominantly lymphohistiocytic infiltrate (Figures 2 and 3).

Discussion

In 1969, Cherry et al. described the acute onset of hemangiomata-like lesions and high fever in 4 children, with symptoms resolving after just a few days. They performed cultures and serological tests, and isolated enteric cytopathic human orphan (ECHO) virus 25 in 2 of the patients and ECHO 32 virus in the other 2. Although they did not perform a skin biopsy, they suggested that given their appearance, the lesions might be due to capillary dilation caused either directly by a viral infection in the endothelium or indirectly via the binding of immunocomplexes to the endothelium. They believed that the disease was a variety of viral exanthem.

In 1993, Prose et al. saw a similar acute eruption in 3 children; they also suspected that it was of viral origin but could not confirm this. They performed a skin biopsy in 1 of the patients and found dilation of the dermal vessels, endothelial projections, and a discrete perivascular infiltrate without vascular proliferation. Electron microscopy did not reveal viral particles in the endothelium, as Cherry had suggested. In view of their findings, they proposed the term eruptive pseudoangiomatosis to describe the disorder. In 1994, Calza and Suarat described the onset of the disorder in 2 brothers and also suggested that it might have been caused by a transmissible vector such as a virus. In Spain, the first case, involving a male child, was reported in 1997. In 2000, Navarro et al. saw a similar clinical picture in a woman who had recovered from an infection of the upper airways and who had also tested positive for Epstein–Barr virus (immunoglobulin IgM and IgG).

Guillot et al. described 9 cases in adults and 4 in children. They noted that eruptions in adults were more common in women than in men, lasted longer than in children, and were not always preceded by fever. They added that they had not observed a specific age of onset and suggested a possible immunosuppressive role as several of the patients had a history of neoplastic disease, systemic corticosteroid treatment, or other situations that might compromise the immune system.

Five additional cases were described in 2000 by Neri et al., Laralde et al., and Angelo et al. In 2003, Joung and Kim described the case of 2 middle-aged Asian women who had developed lesions that were seasonal in nature (appearing and recurring in summer-autumn).

On the other hand, the entity erythema punctatum Higuchi was first described in Japan about 50 years ago. In 1965, Ohara et al. suggested that eruptive pseudoangiomatosis might be associated with mosquito bites, including bites by Culex pipiens pallens. Almost 40 years later, in 2004, Ban et al. described 26 cases of eruptive pseudoangiomatosis-like lesions in patients hospitalized between autumn 2001 and spring 2002. They attributed the lesions to mosquito bites and suggested that these might have been linked to construction work on nearby sewer systems. They also suggested that the process might be the same as that seen in eruptive pseudoangiomatosis and that the viral origin and transmission by insect bites could both be true.

Venturi et al. reported an outbreak of eruptive pseudoangiomatosis in 9 patients and 2 of the investigators in contact with them in an asylum in Parma, Italy. They believed that the origin was viral for several reasons, including the fact that they detected Epstein–Barr virus DNA by polymerase chain reaction in the leukocytes of 4 patients, suggesting a recent replication. One of the patients had a recurrent eruption, which until then had only been seen in children.

Like us, Neri et al. did not see viral particles in an ultrastructural biopsy study performed in a patient aged 7 years. On analyzing the lymphocytic perivascular infiltrate, however, they saw apoptosis and clusters of particles that were suggestive of viral origin. In their opinion, eruptive pseudodystrophy might be a specific response of the body to different viruses, as is the case with Gianotti–Crosti syndrome or the papular–purpuric gloves and socks syndrome.

In a recent letter, Restano et al. explained that several of the patients in their hospital had experienced similar eruptions in the summer months. Most of them were elderly women, immunodepressed patients, or bedridden patients. They attributed the disorder to mosquito bites and said that they had seen similar cases with flea bites.

Nonetheless, there are still doubts regarding the etiology of eruptive pseudoangiomatosis, and although the process seems to be relatively harmless, not being able to offer patients a logical explanation as to the cause of their lesions represents a problem. In our opinion, the disorder is a reactive clinical and pathological response to different etiological processes and is possibly underdiagnosed. Further studies and descriptions of cases are required to shed more light on this disease.

Conflicts of Interest

The authors declare no conflicts of interest.

References