Case Report

Anaphylactoid Purpura Associated with Streptococcal Cellulitis: A Case Report and Literature Review

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A 54-year-old Japanese man noticed painful swelling and redness of his left leg. He was admitted for treatment of cellulitis, which was accompanied with increased anti-streptolysin O and anti-streptokinase titers in his clinical course. After Piperacillin/Tazobactam administration, the skin lesion resolved. However, the patient then developed arthritis, palpable purpura, and intermittent abdominal pain, later found to be secondary to a severe duodenal ulcer. He was diagnosed with cellulitis-associated anaphylactoid purpura and was given prednisolone, which dramatically improved his symptoms. The anaphylactoid purpura was likely caused by Streptococcus-induced cellulitis, which was successfully treated with prednisolone. Association between these diseases is rare.

1. Introduction

Anaphylactoid purpura, also known as Henoch-Schönlein purpura or immunoglobulin A vasculitis, is a systemic vasculitis of the small vessels [1]. This disease is histologically characterized by deposition of immune complexes between antigens and immunoglobulin A, and perivascular leukocyte infiltration in the skin, joints, gastrointestinal system, and kidneys [2]. Although there is no identification of its etiology, anaphylactoid purpura has been reportedly linked to a wide array of pathogens, including infections (bacterial, viral, and parasitic), pharmacological agents, vaccinations, and malignancies [3]. With regard to infections, anaphylactoid purpura frequently develops after upper respiratory tract infections, including sinusitis and focal infections of the oral cavity [4–7]. Consequently, various infectious factors are associated with the development of anaphylactoid purpura; however, few reports have detailed the association between anaphylactoid purpura and cellulitis.

In this case report, we describe a patient diagnosed with Streptococcus-induced cellulitis and subsequent development of anaphylactoid purpura successfully treated with prednisolone. This case report is relevant for clinicians practicing in both inpatient and outpatient settings.

2. Case Report

A 54-year-old Japanese man noticed painful swelling and redness of his left lower leg, which prompted him to visit a general practitioner. His symptoms were suspected to be due to cellulitis, and he was subsequently admitted to our hospital for the treatment of the cellulitis. His daily consumption of alcohol was 180 ml of whisky and his tobacco use amounted to 30 cigarettes per day. His past medical history included chronic hypertension and hyperuricemia. His medications included benzbromarone 25 mg/day, amlodipine 5 mg/day, and telmisartan 40 mg/day. On physical examination, a swollen, painful erythematous region was observed on his left leg.
lower leg. The area of erythema and swelling rapidly increased in size within a short period (Figure 1). Examination of the bilateral femoral lymph nodes was unremarkable, with no evidence of localized lymphadenitis in both sides. His height and weight were 171 cm and 77.0 kg, respectively; therefore, his body mass index was calculated as 26.3 kg/m², which is classified as being overweight according to the World Health Organization criteria for adults [8]. His body temperature was 38.1 °C, his heart rate was tachycardic (101 beats/minute) with a regular rhythm, blood pressure was 124/78 mmHg, and oxygen saturation was 99% breathing room air. Laboratory data on admission showed an increased leukocyte count (13900/μL) and elevated C-reactive protein level (9.48 mg/dL). A blood culture examination was repeated twice with both anaerobic and aerobic tubes; but all results were negative. Although the route of bacterial invasion to the skin was not determined, and there were neither known risk factors for infection nor any systemic immunodeficiency concerns other than an overweight and alcohol polydipsia [9], his cellulitis was suspected to be bacterial in origin due to its rapid progression within a short period. Therefore, Piperacillin/Tazobactam (4.5 g IV every 8 hours) was administered aiming to prevent further skin deterioration, septic shock, and the risk of amputation of his left lower leg. After 9 days of treatment administering Piperacillin/Tazobactam, the skin lesion resolved. However, during the evening of Day 9, the patient subsequently developed palpable purpura on the left lower leg. Furthermore, between Days 14 and 19, the patient reported symptoms of intermittent abdominal pain and painful joints swelling to right ankle, suggestive of acute ankle arthritis. Results of an esophagogastroduodenoscopy, performed on Day 34, revealed a severe ulcer with irregular and widespread lesions, located in the region of the descending portion of the duodenum. A skin biopsy specimen from a purpuric lesion revealed leukocytoclastic vasculitis in the upper dermis (Figure 2(a)) and a duodenal biopsy specimen from the duodenal ulcer also showed eosinophil infiltration into the mucous membrane (Figure 2(b)). Based on these clinical events despite no previous history of upper respiratory tract infections or any deterioration in renal function including urinalysis, the patient was diagnosed with anaphylactoid purpura, which was associated with bacterial cellulitis. As such, a daily oral administration of 40 mg prednisolone was initiated on Day 35 of his hospital stay. Thereafter, his symptoms gradually resolved, and esophagogastroduodenoscopy on Day 48 showed a dramatic improvement of the duodenal ulcer (Figure 3). Despite negative blood culture results, the anti-streptolysin O (ASO) and anti-streptokinase (ASK) antibody titers were elevated at 336 IU/mL (normal range < 239 IU/mL) and 2560 (normal range < 2560) on Day 35, respectively. Therefore, it is likely that the anaphylactoid purpura in this case was associated with a streptococcal infection. This patient was discharged as fully recovered and has been followed up successfully on an outpatient basis since then.

3. Discussion

Anaphylactoid purpura is diagnosed according to the European League against Rheumatism/Pediatric Rheumatology International Trials Organization/Pediatric Rheumatology European Society (EULAR/PRINTO/PRES) criteria. These criteria include palpable purpura and at least one of the four following phenomena: abdominal pain, histopathologic evidence of leukocytoclastic vasculitis or proliferative glomerulonephritis with immunoglobulin A deposits, arthritis or arthralgia, and renal involvement [13]. The incidence of this disease in adults is reported as 0.8–1.8/100,000, which is relatively rare [3]. In our patient, the diagnosis of anaphylactoid purpura was determined clinically upon identification of palpable purpura, abdominal pain, and arthritis of the ankle, although histopathological findings of the skin and duodenal biopsy specimens could not confirm immunoglobulin A deposition because of technical problems.

Anaphylactoid purpura has been reportedly associated with bacterial and viral infections [3, 14]. In particular, upper respiratory tract infections precede the development of anaphylactoid purpura in approximately 30% to 50% of cases (4–6) and have been considered the most common cause in the pathogenesis of anaphylactoid purpura. These infections result in the formation of immunoglobulin A-containing immune complexes in the blood, which can then deposit in small vessels throughout the body, leading to systemic vasculitis [2]. We used the PubMed database for our literature search, with emphasis on publications prior to October 2016, in both the Japanese and English languages, using the Medical Subject Headings terms cellulitis and anaphylactoid purpura. Using these search criteria, only three reports were found, which discussed the association between these diseases. We then reviewed the literature for reports of cellulitis-associated anaphylactoid purpura [10–12] (Table 1). All reported cases, including our cases, as shown in Table 1, were from Japan, although why Japan is the only country to report these particular cases remains unclear. Beta-hemolytic streptococci and Staphylococcus aureus are most commonly implicated as the causative agents of cellulitis, although a number of other microorganisms can uncommonly result in cellulitis. Staphylococcus species were detected in the culture from the exudative fluid in only one patient with cellulitis.
### Table 1: Case reports of cellulitis-associated anaphylactoid purpura.

<table>
<thead>
<tr>
<th>Age</th>
<th>Sex</th>
<th>Location of cellulitis</th>
<th>AP occurrence from cellulitis (days)</th>
<th>Antibiotics for cellulitis treatment</th>
<th>Streptococcal detection by culture</th>
<th>ASO titer (IU/mL)</th>
<th>ASK (times)</th>
<th>Abdominal symptom</th>
<th>Treatment for AP</th>
<th>Reference number</th>
</tr>
</thead>
<tbody>
<tr>
<td>78</td>
<td>M</td>
<td>Right lower leg</td>
<td>6</td>
<td>FOM</td>
<td>Negative (C)</td>
<td>287</td>
<td>2560</td>
<td>(+)</td>
<td>No treatment</td>
<td>[10]</td>
</tr>
<tr>
<td>50</td>
<td>M</td>
<td>Left lower leg</td>
<td>7</td>
<td>CEZ</td>
<td>Positive (C)</td>
<td>1610</td>
<td>640</td>
<td>(−)</td>
<td>No treatment</td>
<td>[11]</td>
</tr>
<tr>
<td>41</td>
<td>M</td>
<td>Left lower leg</td>
<td>3</td>
<td>CEZ</td>
<td>Negative (C)</td>
<td>ND</td>
<td>ND</td>
<td>(−)</td>
<td>DDS</td>
<td>[11]</td>
</tr>
<tr>
<td>78</td>
<td>F</td>
<td>Both lower legs</td>
<td>14</td>
<td>CEZ→SBT/ABPC</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>(−)</td>
<td>PSL</td>
<td>[12]</td>
</tr>
<tr>
<td>54</td>
<td>M</td>
<td>Left lower leg</td>
<td>9</td>
<td>TAZ/PIPC</td>
<td>Negative (B)</td>
<td>336</td>
<td>2560</td>
<td>(+)</td>
<td>PSL</td>
<td>Current case</td>
</tr>
</tbody>
</table>

However, increased titers of ASO and ASK antibodies were confirmed in 3 cases. It was previously reported that the elevated titers of ASO and ASK antibodies were considered evidence of a recent streptococcal infection [15, 16]. Therefore, it is important to pay attention to the relationship between cellulitis, including streptococcal infection, and the development of anaphylactoid purpura. Regarding symptoms related to cellulitis-associated anaphylactoid purpura, only
two patients, including our patient, presented with abdominal pain. In particular, our patient demonstrated a severe duodenal ulcer confirmed by esophagogastroduodenoscopy and required transient total parenteral nutrition. Therefore, this can be considered a very rare case from the viewpoint of the severity of abdominal symptoms accompanied by an unusual manifestation even among rare cases of cellulitis-associated anaphylactoid purpura. As to the treatment of cellulitis-associated anaphylactoid purpura, two out of five patients showed clinical improvement with no specific treatment; however, these two patients presented with skin lesions only and no accompanying systemic symptoms. Prednisolone administration was needed in two patients: one showed recurrence, and the other, in our case, developed systemic manifestations including severe abdominal pain and arthritis. Thus, in the treatment of cellulitis-associated anaphylactoid purpura, patients who experience recurrence and/or systemic symptoms may require prednisolone during their clinical course. On the other hand, administration of Piperacillin/Tazobactam can be associated with the pathogenesis of anaphylactoid purpura and could have been a relevant factor in this case. This agent is well-known to be a broad-spectrum antibiotic widely used for skin and soft tissue infections, abdominal infections, respiratory infections, complicated urinary tract infections, and gynecological infections. It is effective mainly against methicillin-sensitive coagulase-negative Staphylococci, Streptococcus pyogenes, penicillin-sensitive Streptococcus pneumoniae, Haemophilus influenzae, Pseudomonas aeruginosa, and anaerobes [17]. Piperacillin/Tazobactam associated skin reaction, mainly petechial rash, was previously reported and disappeared completely within 3 days once this medication was discontinued [18]. In our case, palpable purpura progressively developed even after the discontinuation of Piperacillin/Tazobactam; therefore, it appears that there was limited association between anaphylactoid purpura pathogenesis and the administration of this agent in this case.

Pathological eosinophil infiltration was observed in the mucous membrane of the duodenal ulcer in this case. Recently, histopathologic results in 68 patients with Henoch-Schönlein purpura revealed eosinophil infiltration, in skin biopsy specimens in 51% of these patients, and also an inverse association between the presence of eosinophils and renal involvement [19]. Thus, the reason that renal involvement was absent in our patient’s clinical course might be associated with the presence of eosinophils in the duodenal lesion, although the mechanism of eosinophil infiltration remains unclear.

In conclusion, we described a case of streptococcal cellulitis-associated anaphylactoid purpura. Based on our experience, we suggest that not only do patients with cellulitis need to be examined and treated using antibiotic agents, but additionally attention needs to be paid to the development of anaphylactoid purpura during their clinical course.

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this article.

References


