Ecthyma gangrenosum caused by *Pseudomonas aeruginosa* in a patient with astrocytoma treated with chemotherapy

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Abstract  Ecthyma gangrenosum, presenting as embolic lesions caused by *Pseudomonas aeruginosa* infection, has distinct pathognomonic features and a high mortality rate in patients with bacteremia, but when recognized early is easily treated. In this case report we describe this disseminated infection in an adult patient treated with chemotherapy for an astrocytoma.

Keywords  Ecthyma gangrenosum · *Pseudomonas* · Hemorrhage · Bullous

Introduction

Lesions of the skin are common in cancer patients, usually occurring in the context of paraneoplastic skin events such as erythema nodosum and dermatological alterations due to allergic reactions or the side effects of systemic antitumor therapy, or in the context of erysipelas or cellulitis caused by either *Streptococcus* or *Staphylococcus*. However, in patients with chemotherapy-induced neutropenia (defined as <500 functional polymorphonuclear leukocytes/μl), innocent-looking macules or papules may be the first signs of sepsis with uncommon organisms.

Ecthyma gangrenosum is a rare cutaneous infection with painless, round, necrotic lesions consisting of a central black or gray-black eschar with surrounding erythema. The lesions are usually localized in nonpressure areas (in contrast to ischemic lesions), such as the extremities and gluteal and perineal regions. Solitary lesions have a better prognosis than multiple lesions, the latter being commonly related to life-threatening *Pseudomonas aeruginosa* bacteremia. Further, the prognosis is determined by an early diagnosis, the duration of neutropenia, and the start of appropriate antibiotic treatment. Predisposing factors, such as chemotherapy, extensive burns, hypogammaglobulinemia, hematological malignancies, and comorbidities (e.g., diabetes mellitus and malnutrition), contribute to the severity of the immunodeficient status, and thus negatively to the outcome [1]. Mortality rates of *Pseudomonas* sepsis in immunocompromised persons range from 38 to 96%, whereas the mortality rate in nonbacteremic patients is 15.4% [2]. The reported high mortality rate could be diminished by timely recognition of the early occurring pathognomonic characteristics of ecthyma gangrenosum, so that appropriate systemic antibiotic therapy can be initiated.

Case report

A 37-year-old Caucasian man with a history of low-grade oligo-astrocytoma in the right temporo-parietal lobe presented in October 2008 with epileptic insults, sensory neuropathy in the left hand, and coordination dysfunction of the left leg. Computed tomography (CT) of the brain showed progressive disease with an increase in size of the known lesion. As radiation therapy had previously been given, with a total dose of up to 50 Gy on the tumor site,
this was not an option anymore. Therefore, chemotherapy was opted for, and PVC chemotherapy, consisting of lomustine 110 mg/m² on day 1, procarbazine 60 mg/m² on days 8–21, and vincristine 1.4 mg/m² on days 8 and 29, to be repeated every 6 weeks, was started in November 2008. Further relevant comedication consisted of dexamethasone 2 mg, levetiracetam 1500 mg, valproate 1500 mg, and clobazam 10 mg, all twice daily. On the evening of day 40 of the first course, he was admitted with febrile neutropenia and cold chills. He had no worsening of his neurological symptoms. There were no localizing complaints. At physical examination blood pressure measured 139/69 mmHg with a pulse of 109 beats per min and a temperature of 38.5°C. On his arms, legs, abdomen, and pubic area several hemorrhagic bullous lesions with surrounding erythema were observed, some with central black necrosis (Figs. 1, 2). There were no other physical signs of possible infection found. Laboratory results showed 6.6 mmol/l hemoglobin (normal values 8.6–10.5 mmol/l), less than 0.05 × 10⁹/l neutrophils (normal values 1.4–8 × 10⁹/l), and 63 × 10⁹/l thrombocytes (normal values 150–370 × 10⁹/l), with 60 mg/dl C-reactive protein (maximum value in hospitalization period 170 mg/dl; normal values 0–9 mg/dl). Syndrome of inappropriate antidiuretic hormone secretion by central nervous malignancy was observed, with 125 mmol/l sodium (normal values 136–145 mmol/l) and 260 mOsm/kg serum osmolality (normal values 270–300 mOsm/kg). After several pictures of the skin lesions had been put on a secure intranet link, a dermatologist was consulted by phone, and suggested sampling bacterial cultures of skin and bullous fluid. Bacteriological sampling was taken from blood, urine, skin, and bullous fluid. Because of the patient’s neutropenic fever, we started imipenem/cilastatin 500/500 mg four times daily, intravenously, after the sampling. A day later, the consulted dermatologist examined the patient. At that time, the clinical picture, combined with positive culture of *Pseudomonas aeruginosa*, was typical for the diagnosis of ecthyma gangrenosum caused by disseminated *Pseudomonas* infection, and therefore histology of the skin lesions was not obtained. The antibiotic treatment was switched from imipenem/cilastatin to ceftazidime 1 g six times a day and tobramycin 7 mg/kg according to tobramycin blood levels, given intravenously [3, 4]. The clinical condition improved after 3 days of hospitalization, the fever subsided after 1 day of this antibiotic treatment, and the neutrophils were restored to a normal level in 9 days. After 1 week, the antibiotics were switched to ciprofloxacin 750 mg bid for 1 week, given orally, and the patient could be discharged. In the second chemotherapy course, lomustine and procarbazine were reduced to 75% of the original dose, and the patient remained without a recurrent neutropenic episode.

### Discussion

Ecthyma gangrenosum is a cutaneous entity that is classically associated with septicemia and gram-negative bacteremia caused by *Pseudomonas aeruginosa* in immunocompromised patients. Usually, this disseminated cutaneous infection presents with embolic lesions with the formation of bullae. The exact mechanism of pathogenesis of this disease in a neutropenic patient is poorly understood. The *Pseudomonas* organism releases a variety of proteases in outer-membrane derived vesicles, regulated by a quorum sensing system. The virulence of these pathogenic factors leads to invasion of the medial and adventitial layers of the vascular wall of nearby blood vessels, resulting in septicemia, bacteremia, embolization, and dissemination [5, 6]. After invasion of the subcutaneous tissue, the bacilli are usually found in the collagen bundles of the dermis and panniculus [7]. Proliferation of the
organisms leads to the production and release of exotoxin A and proteases, resulting in the formation of bullae and ulceration. As the diagnosis in our patient was already clear from the positive bacterial culture of the bullous fluid, histology was not performed, but would typically show necrotizing hemorrhagic vasculitis with bacterial infiltration into the media and adventitia of vessels resulting in skin necrosis. The lesions are usually situated in the gluteal and perineal region, but can spread to the trunk and extremities in rare cases even to the face. Differential diagnosis includes pyoderma gangrenosum, other forms of vasculitis, and the presence of cryoglobulins or septic emboli from other microorganisms (Escherichia coli, Aeromonas hydrophilia, Pseudomonas cepacia, Candida, Fusarium species, and Aspergillus species).

Major risk factors for the development of ecthyma gangrenosum, such as chemotherapy, extensive burns, hypogammaglobulinemia, hematological malignancies, and comorbidities (e.g., diabetes mellitus and malnutrition), are all associated with an immunocompromised condition [2]. The combination of these different risk factors contributes to a life-threatening situation with a high mortality rate in the case of a systemic infection. Mortality rates vary from 38 to 96% in patients with bacteremia, and are approximately 15% in patients with a nonbacteremic condition. Multiple skin lesions predict a worse prognosis and can be diagnosed in both septicemic and nonsepticemic patients. However, ecthyma gangrenosum caused by Pseudomonas aeruginosa can develop in only a small number of cases in the absence of bacteremia [8, 9]. The recommended treatment for both bacteremic and nonbacteremic ecthyma gangrenosum combines an anti-

References