A Case of Sweet’s Syndrome in a Patient with Liver Cirrhosis Caused by Chronic Hepatitis B

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Sweet’s syndrome (SS), also known as acute febrile neutrophilic dermatosis, is characterized by the sudden onset of painful erythematous skin lesions together with fever and neutrophilia. SS can be associated with several disorders, such as malignancy, autoimmune disease, and infections. However, SS associated with liver cirrhosis is uncommon. We report a case of SS in a patient who was diagnosed with liver cirrhosis caused by chronic hepatitis B. (Korean J Gastroenterol 2012;59:441-444)

Key Words: Sweet’s syndrome; Liver cirrhosis; Chronic hepatitis B

INTRODUCTION

Sweet’s syndrome (SS) is the prototype of neutrophilic dermatosis, which is an uncommon disorder characterized by abrupt onset of fever, leukocytosis, and painful red, popular skin rash. Additionally, SS is histologically characterized by diffuse infiltration of mature neutrophil in the dermis without vasculitis. Lesions usually develop on the face, neck, arms, and hands and are asymmetrically distributed. However, SS can be generalized, and patients often complain of discomfort with its associated signs and symptoms, including malaise, fever, elevated erythrocyte sedimentation rate (ESR), and elevated CRP levels, which mimic an infectious process.

SS accompanied with liver cirrhosis is uncommon, and there are few reports of SS associated with acute or chronic hepatitis. Here, we report a case of SS in a patient who was diagnosed with liver cirrhosis caused by chronic hepatitis B.

CASE REPORT

A 45-year-old man, who was diagnosed with liver cirrhosis related with chronic hepatitis B, visited our department with multiple, purplish-red infiltrated papules on the left thigh, right inguinal region, and both arms during the past 4 days. He had no specific respiratory or gastrointestinal symptoms except generalized edema and a distended abdomen.

One year ago he was admitted to the hospital with gastric ulcer bleeding caused by alcohol ingestion. At that time, he was diagnosed with liver cirrhosis caused by chronic hepatitis B. Since then, he has not consumed alcohol and has regularly visited the hospital.

At 2 months before admission, he experienced hepatic encephalopathy caused by high protein diet and his physical condition deteriorated. At 1 week before admission, he had a distended abdomen, and both lower legs were edematous. He did not take any other medications except ursodeoxycholic...
acids and diuretics as prescribed by doctors.

On admission, his vital signs were as follows: blood pressure, 100/60 mmHg; pulse rate, 80/minute; respiration rate, 20/minute; and body temperature, 38.5°C. He looked acutely ill but was alert. His heart and lung sounds were normal. He had a moderately distended abdomen, edematous lower legs, and swelling of the scrotum. A skin examination revealed multiple tender, erythematous and annular lesions on both arms and the trunk (Fig. 1A, B) as well as ulcerative lesions on the left inguinal region (Fig. 1C).

Peripheral blood tests showed white blood cells, 8,040/mm³ (neutrophils, 78.2%); hemoglobin, 9.0 g/dL; and platelets, 68,000/mm³. Routine chemistry showed total protein, 6.2 g/dL; albumin, 2.5 g/dL; total bilirubin, 2.1 mg/dL; ALP, 95 IU/L; AST, 47 IU/L; ALT, 27 IU/L; and prothrombin time, 20.2 seconds (normal range, 10-14 seconds; INR, 1.78). HBsAg and HBeAg were positive. The serum HBV DNA titer was 9,737,316 IU/mL. His ESR was 42 mm/hour (normal range, 0-10 mm/hour) and CRP was 0.6 mg/dL (normal range, 0-0.5 mg/dL).

Tumor markers (α-fetoprotein and carcinoembryonic antigen) were within normal ranges, and anti-nuclear antibody, anti-mitochondrial antibody, and human immunodeficiency virus screening were negative. An ascites study showed white blood cells, 243/mm³ (polymorphonuclear cells, 70%); protein, 0.5 g/dL; albumin, 0.3 g/dL; and microorganisms were negative in a culture.

Endoscopic findings revealed a well-defined active ulcer at the lesser curvature side of the distal body and duodenal bulb. F1 sized, red color sign (-) esophageal varices, which were classified by the Japanese Society for Portal Hypertension, were noted on the lower one-third of the esophagus. A biopsy specimen from the gastric ulcer site revealed a benign ulcer with Helicobacter pylori.

A chest X-ray finding was normal. Abdominal computed tomography showed a shrunken liver, splenomegaly, and a moderate amount of ascites.

Before histological confirmation, he received antibiotic treatment for 3 days to prevent wound infection. A histological examination of tissue obtained from the inguinal region showed dense infiltration of mature neutrophils in the superficial dermis without vascular infiltration (Fig. 3). After histological confirmation, he received 30 mg/day oral prednisolone, and the fever declined and normalized on the fifth day after medication. After 1 week of prednisolone, the cutaneous lesions had almost completely healed without scars. Oral prednisolone was tapered gradually during a 2-week period. He did not receive any antiviral agents initially because of economic problems. During regular follow-up of his

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**Fig. 1.** Multiple tender, raised, annular erythematous lesions on the arms (A) and trunk (B). Round pustular lesions with central necrosis and skin ulcers on the left thigh (C).

**Fig. 2.** Abdominal computed tomography scan showed a shrunken liver, splenomegaly, and a moderate amount of ascites.
liver enzymes, the ALT level increased after 2 weeks of prednisolone. So, he took 100 mg lamivudine daily, and his ALT level gradually normalized.

DISCUSSION

SS, also known as acute febrile neutrophilic dermatosis, was first described by Robert Douglas Sweet in 1964. SS is characterized by tender, erythematous papules and plaques, fever, and leukocytosis, which respond well to corticosteroid therapy. The importance of SS is a marked clinical expressiveness and frequent association with other systemic diseases.

SS occurs more frequently in females and can appear at any age, but peaks in patients in their 50s-70s. SS can be triggered by infections, hematological malignancies, other neoplasms, autoimmune diseases, pregnancy, trauma, and medications. Immune complex vasculitis, T-cell activation, and altered neutrophil function have been presumed to occur, but there is lacking experimental evidence.

Diagnostic criteria, which are comprised of two major criteria and four minor criteria, were recently modified by von den Driesch in 1994. Both major criteria and two of the four minor criteria should be satisfied for a definitive diagnosis of SS. The two major criteria are 1) abrupt onset of painful erythematous plaques or nodules and 2) histopathological evidence of a dense neutrophilic infiltrate without evidence of primary leukocytoclastic vasculitis. The minor criteria are 1) pyrexia (> 38°C); 2) an association with an underlying hematological or visceral malignancy, inflammatory disease, or pregnancy, or preceded by an upper respiratory or gastrointestinal infection or vaccination; 3) excellent response to treatment with systemic corticosteroids, potassium iodide, or colchicine; 4) abnormal laboratory values at presentation (three of the following four): ESR > 20 mm/hour, positive CRP, > 8,000 leukocytes, and > 70% neutrophils. The present case was sufficient to meet these criteria.

SS related skin lesions can be preceded or accompanied by fever and general discomfort and should be diagnosed differently from erythema multiforme, erythema elevatum diutinum, erythema nodosum, and pyoderma gangrenosum.

The histological differential diagnosis of SS includes conditions microscopically characterized by either neutrophilic dermatoses or neutrophilic panniculitis. Neutrophilic dermatoses include abscesses or cellulitis, intestinal bypass syndrome, leukocytoclastic vasculitis, pyoderma gangrenosum, and rheumatoid neutrophilic dermatitis. Leukemia cutis cannot occur concurrently with SS, but mimics the dermal changes of SS. In contrast to SS, the dermal infiltrate in patients with leukemia cutis consists of malignant immature leukocytes rather than mature neutrophils. The adipose tissue changes that occur with subcutaneous SS are similar to those of other conditions characterized by neutrophilic lobular panniculitis. Similar to pyoderma gangrenosum, SS is known to cause pathergy, in which lesions occur in areas of minor trauma, such as sites of scratches, bites, and venipuncture. The lesions may also be photodistributed or localized to the site of a previous phototoxic reaction.

In some patients with SS, skin lesions eventually resolve without any therapeutic management. However, the skin lesions usually persist for weeks to months without treatment. Systemic steroids are the gold standard therapy for SS. Dermatosis-associated symptoms improve rapidly after starting steroid therapy, and the cutaneous lesions resolve subsequently. Systemic corticosteroid therapy usually begins with 1 mg/kg/day prednisolone as a single oral morning dose. However, several studies have suggested starting prednisolone at 30 mg per day, so our patient was also started on 30 mg prednisolone and gradually tapered, and his symptoms and skin lesions improved promptly.
HBV infection is also an SS trigger factor. HBV is well known to cause several types of skin manifestations such as vasculitis, urticaria, and polyarteritis nodosa. But, the pathogenesis of SS-related hepatitis B is unclear.

SS pathogenesis is thought to be an abnormal immune reaction triggered by variety of bacterial, viral, or tumorous antigens. Physical condition including generalized edema or deteriorated immunity by hepatic decompensation is thought to provide additional chances for infection or altered T-cell activation and neutrophil function. Our case was a patient with liver cirrhosis. During the 2 months before he developed SS, he experienced hepatic encephalopathy and consecutive generalized edema. Deteriorated liver status is thought to be a more vulnerable condition than to be infected by a virus or bacteria.

SS skin lesions usually appear similar to other infectious diseases. But, SS skin lesions improve after using steroids not antibiotics. Thus, a differential diagnosis of SS should be considered if patients with skin lesions are not effectively controlled by antibiotics. If steroids are not indicated, dapsone, indomethacin, or potassium iodide can be used to improve symptoms.

Reactivation of HBV replication, which increases HBV DNA and ALT level, has been reported in 20-50% of hepatitis B carriers undergoing immunosuppressive or cancer chemotherapy and is more common when chemotherapeutic regimens include corticosteroids. Therefore, regular follow-up of serum HBV DNA and ALT level is very important.

According to the practice guidelines of the American Association for the Study of Liver Disease, prophylactic antiviral therapy should be administered to hepatitis B carriers at the onset of cancer chemotherapy or a finite course of immunotherapy and maintained for 6 months. Our patient also experienced an ALT flare up after prednisolone treatment. Thus, regular monitoring of HBV DNA, ALT level, and prophylactic antiviral therapy is very important for hepatitis B carriers who receive immunosuppressive agents including corticosteroids.

We report a case of SS in a patient with liver cirrhosis caused by chronic hepatitis B that was suspected to be triggered by his physical condition including generalized edema and deteriorated immunity by hepatic decompensation.