Letters to the Editor

Sir,

Symmetrical peripheral gangrene (SPG) is defined as symmetrical distal ischemic change at two or more sites in the absence of any vasculitis or large-vessel obstruction. SPG caused by malaria is very rare, and most of the cases are associated with Plasmodium falciparum. Herein, we present a case of SPG due to severe Plasmodium vivax malaria.

A 62-year-old female presented in the emergency department with complaints of fever for the last 4 days which was intermittent in nature and was associated with chills and rigor. On examination, she was febrile with an oral temperature of 104°F. Her pulse rate was 120/min, regular and all peripheral pulses were palpable, and blood pressure was 110/70 mmHg. There were multiple petechial spots over the lower limbs. Hemogram on the day of admission showed hemoglobin of 8.2 g/dl, total leukocyte count of 10,300/cumm (N73, L26, and E1), and platelet count of 22,000/cumm. Peripheral smear showed schizonts of P. vivax, and malaria kit test was also positive for P. vivax. Her renal (urea - 121 mg/dl, creatinine - 2.3 mg/dl) and liver function tests (bilirubin - 2.7 mg/dl, aspartate aminotransferase - 520 IU/L, alanine aminotransferase - 455 IU/L, and alkaline phosphatase - 258 U/L) were also deranged. Blood sugar was 40 mg/dl, and all electrolytes were normal except corrected serum calcium which was 7.5 mg/dl (normal: 8.5–10.5 mg/dl). Arterial blood gas analysis showed metabolic acidosis (pH - 7.29, pCO₂-30.2 mmHg, pO₂-62.5 mmHg, and bicarbonate - 14.5 mEq/L). Serum lactate dehydrogenase was 6690 U/L (normal: 200–400 U/L) and urinalysis was normal. A diagnosis of severe P. vivax malaria was made and she was started on injection artesunate at dose of 2.4 mg/kg. Dextrose containing intravenous fluid and oral paracetamol was given for hypoglycemia and fever, respectively. On the 3rd day of admission, she started complaining of burning sensation in her feet and right hand with blackish discoloration [Figures 1 and 2]. All peripheral pulses including dorsalis pedis were palpable. Prothrombin time was 10 s (normal: 11–15 s), activated partial thromboplastin time was 35 s (normal: 22–34 s), and D-dimer was 12 mcg/ml (normal: <0.50 mcg/ml fibrinogen equivalent units). Fibrin degradation product was 32 mcg/ml (normal: <10 mcg/ml). Other investigations such as blood culture, urine culture, HIV, hepatitis B surface antigen, anti-hepatitis C virus, anti-nuclear antibody, and antiphospholipid antibody were normal. Ultrasound abdomen, arterial Doppler of limbs, and echocardiography were normal. Injection artesunate was continued for 5 days and her platelet count improved to 1.6 lacs/cumm; she was then given 3 days course of artemisinin-based combination therapy (artemether and lumefantrine). However, there was gradual worsening of renal and liver function tests for which she received two cycles of hemodialysis. Blackish discoloration of hand and feet gradually progressed to develop dry gangrene by the end of 2nd week [Figure 3]. At time of discharge from the hospital, she was afebrile and all biochemical parameters were normal. She was discharged on tablet primaquine at a dose of 0.25 mg/kg/day for 2 weeks after testing glucose-6-phosphate dehydrogenase levels. She was further advised to follow-up in the Surgery Department for amputation of digits.

The mechanism of SPG in falciparum malaria is not known and is believed that heavy parasitemia causes activation of complement and coagulation pathway which eventually leads to thrombosis of microvasculature. Infected erythrocytes interact with various endothelial receptors such as intercellular cell adhesion molecule-1, vascular cell...
adhesion molecule-1, thrombospondin, and histidine-rich protein causing microcirculatory obstruction.[3] Disseminated intravascular coagulation is also a contributing factor in malaria-related SPG and is rarely associated with significant bleeding.[4] Management of SPG in malaria is supportive. There is no specific treatment for SPG per se and routine use of antiplatelets or anticoagulants is not indicated. The mortality rate of SPG is almost 35% and 70%–90% patients might need amputation after line of demarcation develops.[5]

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Conflicts of interest
There are no conflicts of interest.

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References

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