February 2017 Critical Care Case of the Month

Morgan Wong, DO
Nicholas Villalobos, MD

Department of Internal Medicine
University of New Mexico
Albuquerque, NM USA

History of Present Illness
A 68-year-old man presented to the emergency department with a one-day history of lower back pain, arthralgias, and malaise. The patient had a previous splenectomy and was concerned about influenza.

Past Medical History, Social History, and Family History
He has a history of osteoarthritis, seasonal allergies, and splenectomy. He is a nonsmoker. Family history is noncontributory.

Physical Examination
Upon admission, the patient’s vital signs were notable for a temperature of 35.3 degrees Celsius, blood pressure of 74/44 mmHg, oxygen saturation of 85% on room air with a respiratory rate of 24 breaths per minute. Physical exam was prominent for non-pitting edema of the distal upper and lower extremities, as well as diffuse macular rash of the palms and soles.

Laboratory
CBC
- White blood cell count of 6.77 X10^3 cells/uL
- Hemoglobin of 13.8 gm/dL
- Hematocrit of 43.7%
- Platelet count of 19 x 10^3 /uL
Chemistry
- Creatinine of 3.0 mg/dL
- CO2 < 10 mmol/L
- Anion gap >18 mmol/L
Liver function tests
- Alanine aminotransferase (ALT) of 511 U/L
- Aspartate aminotransferase (AST) of 529 U/L
- Total bilirubin of 1.0 mg/dL
Coagulation
- INR of 2.07
- Prothrombin time of 22.5 seconds
- Partial thromboplasrin time of 82.3 seconds
- Fibrinogen level was 71 mg/dL
Arterial blood gases
- pH of 6.91
- pCO₂ 54 mmHg
- pO₂ 263
- HCO₃ of 7.7 mmol/L
- Procalcitonin >200 ng/ml.

His blood peripheral smear was examined.

![Peripheral blood smear on admission](image)

**Figure 1:** Peripheral blood smear on admission.

Given the results of the preliminary laboratory results and peripheral smear what **hematologic abnormality** are you most concerned with at this time?

1. Autoimmune hemolytic anemia (AIHA)
2. Disseminated intravascular coagulopathy (DIC)
3. Microangiopathic hemolytic anemia (MAHA)
4. Thrombotic thrombocytopenic purpura (TTP)
DIC is a pathological imbalance in the procoagulant and anticoagulant systems, which leads to unregulated thrombin generation (1). DIC is secondary to many states, including sepsis/infection, malignancy, and trauma. DIC classically presents with a low platelet count, elevated INR, elevated PT and PTT times, as well as a decreased fibrinogen level. Peripheral smear demonstrates shistocytes, and low platelet levels that are also reflective of DIC. Microangiopathic hemolytic anemia is incorrect given it typically presents with normalized PT, and PTT times, as well as normalized fibrin levels. Thrombotic thrombocytopenia purpura and autoimmune hemolytic anemia would be unlikely given the patient’s normal bilirubin levels.

The patient was subsequently intubated and placed on norepinephrine for hypotension and piperacillin and tazobactam for presumed sepsis. Blood cultures were positive for gram-positive cocci in two out of two cultures. A lumbar puncture was performed which noted a total nucleate cell count of 1/uL, glucose of 57 mg/dL, protein 69 mg/dL, with an HSV IgG of 0.12. Lactic acid was 5.5mmol/L. *Streptococcus pneumoniae* antigen and urinary *Legionella* antigen were negative. Computed tomography (CT) scan of the brain, thorax, abdomen, and pelvis was notable for diffuse body wall edema as well as bilateral lower lobe atelectasis.

Given the patient’s medical history and clinical presentation what organism would have likely precipitated the current presentation?

1. *Haemophilus influenzae*
2. *Neisseria meningitidis*
3. *Streptococcus agalactiae*
4. *Streptococcus pneumoniae*
3. *Streptococcus agalactiae*

The patient presents with a history of splenectomy, making encapsulated organisms the most likely causative organism. *Streptococcus agalactiae* or Group B Streptococcus (GBS) is a gram positive encapsulated organism that has been documented to present with hypotension, coagulopathy, renal dysfunction, and erythematous rash (2). *Streptococcus pneumoniae* is unlikely given negative clinical history without notable findings on CT imaging, and negative streptococcal antigen. *Neisseria meningitidis* is unlikely given no sign of meningismus and a lumbar puncture with low cell count and normal glucose levels. *Haemophilus influenzae* is a gram-negative bacteria.

The resident astutely notes that a finger is necrotic distally (Figure 2).

![Figure 2. Photograph of patient’s necrotic finger.](image)

On further examination, the patient’s toes are also necrotic.

What **therapy** would you initiate?

1. Dobutamine
2. Epoprostenol
3. Norepinephrine
4. Phentolamine
Correct!

3. Norepinephrine

The patient has dry gangrene from a myriad of causes to include possible hypothermia with potential frostbite injury, toxic shock secondary to group B *Streptococcus*, disseminated intravascular coagulopathy, and now with the addition of vasopressor support; potential vasospasm. Group B *Streptococcus* has been shown to present with a toxic shock like syndrome (TSLs) from pyrogenic toxin(s) (2-4). The goal in therapy is to treat the underlying disease but preservation of the extremity is pivotal. In this case, continuation of norepinephrine with a plan to discontinue as soon as possible is the most appropriate plan (5). Phentolamine is incorrect as there is no evidence of pressor extravasation. There is limited evidence for the use of epoprostenol used solely as reversal for vasospasm. Dobutamine at this time is not a preferred vasopressor.

Once you receive speciation, what antibiotics should be added or augmented?

1. Addition of ampicillin and sulbactam and discontinuation of piperacillin and tazobactam
2. Addition of cefepime and discontinuation of piperacillin and tazobactam
3. Addition of cefepime only
4. Addition of clindamycin
Correct!

4. Addition of clindamycin

The addition of clindamycin for its inhibition of bacterial protein synthesis is crucial to reduce the severity of the disease (2-4). Linezolid otherwise, would be preferred in areas of high clindamycin resistance. The other choices do not act specifically to inhibit protein synthesis.

The mortality rate of toxic shock like syndrome secondary to Group B *Streptococcus agalactiae* approaches 50% in review of current literature with rates of infection anywhere from 22 to 30 per 10,000 patients per population (2-4). There are no current vaccinations available to patients for prevention of this encapsulated organism like there is for *Haemophilus influenza*, *Neisseria meningitidis*, and *Streptococcus pneumoniae*.

References