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A Case Report of Hemophagocytic Lymphohistiocytosis Secondary to Disseminated Tuberculosis

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INTRODUCTION

Hemophagocytic lymphohistiocytosis (HLH) is a life-threatening disorder characterized by excessive activation of the immune system causing tissue damage and organ dysfunction. Secondary HLH is more common in adults and is usually triggered by infection, malignancy, rheumatologic, and immunodeficiency syndromes. We present a case of HLH secondary to disseminated tuberculosis (TB).

CASE PRESENTATION

A 72-year-old woman with past medical history of hypertension and type 2 diabetes presented to the emergency department for one day history of fevers, poor oral intake, and fatigue. She was febrile to 102.9 F and noted to have a toxic appearance, but there were no focal abnormalities found on exam. Initial blood work was notable for a mild direct hyperbilirubinemia of 2.5mg/dL, AST/ALT of 162/112 IU/L and elevated INR of 1.4. Abdominal ultrasound showed gallbladder wall thickening. Due to concerns for ascending cholangitis, ERCP was performed and revealed a normal biliary system. Day 2 of hospitalization revealed new pancytopenia - her white blood cell count was 2.7 B/L with 44% bands, hemoglobin was 10.8 g/dL, and platelet count was 39 B/L. She also had rising hepatic function tests. Her ferritin on day 3 was significantly elevated at 4149 IU/L, raising concerns for HLH. On day 4, the patient became increasingly tachypneic and tachycardic with radiographic findings of pulmonary edema despite diuresis, as well as acute kidney injury with decreased urine output. She was intubated for increased work of breathing. Due to a high suspicion of HLH with the probability of HLH greater than 99% calculated by the HScore, methylprednisolone 120mg daily in split doses was initiated. Etoposide was not started due to her multiorgan dysfunction. Instead, anakinra, an IL-1 receptor antagonist was used in combination of dexamethasone. Anakinra as an immunomodulator has also been shown to be effective in the treatment of severe HLH in the critical care setting.

DISCUSSION

Although HLH secondary to TB (TB-HLH) has previously been reported in the medical literature, it remains a rare entity. Most patients with TB-HLH had short symptom duration and rapid progression leading to multi-organ dysfunction and ultimately, death.

The first HLH treatment protocol was published in 1994 (HLH-94) and has been widely used in clinical practice. HLH-94 includes an anti-inflammatory agent, dexamethasone, and a pro-apoptotic agent, etoposide. This regimen has been shown to improve survival but was primarily based on data in the pediatric setting, many with primary HLH. Furthermore, etoposide was not determined to be a safe option due to her multi-organ dysfunction. Instead, anakinra, an IL-1 receptor antagonist was used in combination of dexamethasone. Anakinra as an immunomodulator has also been shown to be effective in the treatment of severe HLH in the critical care setting.

Additionally, secondary HLH is often triggered by an acute infection or other condition (eg, rheumatologic condition) and treatment should be to address the underlying cause of immune activation. Although there is no consensus of standard treatment for TB-HLH, a systematic review of the international literature demonstrates that a delay or absence of antitubercular therapy (ATT) were associated with decreased survival. On the contrary, the combination of antitubercular therapy (ATT) with immunotherapy has been found to significantly reduce mortality. Our patient was initiated on a modified regimen of ATT as soon as the diagnosis of disseminated TB was made, but she still passed several days later. This case highlights the need for ongoing investigation for early detection and management of secondary HLH in the adult setting.
REFERENCES


