Hypertensive Emergency as the Initial Presentation of Acquired Thrombotic Thrombocytopenic Purpura: A Case Report

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Abstract

Thrombotic thrombocytopenic purpura (TTP) is a type of thrombotic microangiopathy (TMA) related to a deficiency of ADAMTS13 protein, which could lead to fatal outcomes. TTP presents a vast array of symptoms, making its diagnosis a challenge to physicians. In this report, we discuss the case of an 80-year-old male who presented with abdominal pain and hypotension with underlying severe thrombocytopenia and hemolysis. Although his presentation could also be secondary to malignant hypertension, he was promptly diagnosed with TTP based on these findings. TTP is a hematologic emergency, and it should be kept in the differential diagnosis when a patient presents with thrombocytopenia and hemolysis with or without accompanying symptoms.

Categories: Cardiology, Internal Medicine, Hematology
Keywords: adamat13 protein, purpura, malignant hypertension, thrombotic thrombocytopenic purpura, thrombotic microangiopathies

Introduction

Thrombotic thrombocytopenic purpura (TTP) is a rare and highly fatal disease. It forms part of thrombotic microangiopathies (TMAs) and is characterized by hemolytic anemia, thrombocytopenia, and multiorgan dysfunction due to microthrombosis in arterioles and capillaries [1]. TTP can present with a vast array of symptoms such as petechiae, neurological deficits, confusion, seizures, and even myocardial infarction. It is not uncommon to find abdominal pain as the primary concern of patients looking for medical attention, such as in the case presented here. However, not all patients with TTP are critically ill. The majority of patients do not present with clear symptoms of thrombocytopenia when they seek medical attention; they are usually diagnosed incidentally following a routine complete blood count [2]. The PLASMIC score (Platelet count; combined hemolysis variable; absence of Active cancer; absence of Stem-cell or solid-organ transplant; Mean corpuscular volume (MCV); International normalized ratio (INR); Creatinine) is key to establishing a prompt and accurate diagnosis in a patient with thrombocytopenia and hemolytic anemia. The TTP pentad is only present in 10% of patients and it is unreliable to use this as the sole parameter for the diagnosis. This pentad consists of fever, neurological abnormalities, kidney injury, abdominal pain, and thrombocytopenia [3]. Due to the ample and varied presentation of TTP, these parameters will aid in differentiating it from other causes of microangiopathies such as hypertensive emergencies that can present initially in a similar manner. In this report, we aim to highlight the crucial need for accurate differentiation between TTP and hypertensive emergency, as well as emphasize the importance of rapid management to reduce fatality in this type of hematologic emergency.

Case Presentation

An 80-year-old Indian male presented to the emergency department (ED) with a one-day history of epigastric and left lower quadrant pain with mild relief after ibuprofen administration. The patient had a history of hypertension, diabetes mellitus type 2, hyperlipidemia, prostate cancer, and subclinical hypothyroidism. His surgical history included an endoscopic retrograde cholangiopancreatography (ERCP) and laparoscopic cholecystectomy. His medications at that time included metformin, omeprazole, gabapentin, carvedilol, atorvastatin, ferrous sulfate, cyanocobalamin, levothyroxine, amlodipine, and aspirin. The patient disclosed that he was inconsistent with taking his medications and stated that his conditions were poorly controlled. He denied any alcohol, tobacco, or recreational drug use. Vital signs on admission evidenced high blood pressure of 182/95 mmHg, a pulse of 89 bpm, a respiratory rate of 19 breaths per minute, and a temperature of 36.4 °C. Initial laboratory test results included an international normalized ratio (INR) of 1.2, a serum creatinine of 1.6 mg/dL, a white blood cell count of 8,900 cells/mL, an hemoglobin of 10.6 g/dL, and a platelet count of 14,000 cells/μL. The patient was started on a regimen of intravenous fluids, blood transfusions, and aspirin. The patient was discharged after 48 hours of admission and was referred to a surgical team for diagnostic workup. The patient was discharged on aspirin and levothyroxine and scheduled for follow-up appointments with primary care. The patient was advised to follow a healthy diet and to continue taking medications as prescribed. The patient’s condition improved during follow-up appointments, and he was discharged on a regular diet and medications. The patient was advised to follow up with his primary care physician for further management.
the remaining 50% of cases in infections, cancer, pregnancy, and drugs (e.g., cyclosporine, quinine, clopidogrel). Idiopathic TTP is seen in triggers, as follows: autoimmune disorders (e.g., systemic lupus erythematosus), viral and bacterial.

There are two types of acquired TTP: idiopathic and non-idiopathic. Non-idiopathic TTP is seen in about 50% of patients diagnosed with this disorder. The most frequent conditions associated with TTP, serving as 50% of cases.

manifestations in TTP are hematuria and proteinuria, whereas renal manifestations in malignant hypertension include a rise in creatinine, BUN, hematuria, and proteinuria.

Thrombocytopenia. This leads to the accumulation of microthrombi in the capillaries that in turn lead to hemolysis. This also affects the blood flow in the microcirculation that causes end-organ damage, mainly in the central nervous, renal, and cardiovascular systems. TMA in a hypertensive emergency is secondary to shear stress due to the narrowing of vessels that leads to excessive fragmentation of erythrocytes and the consumption of platelets.

Also, renal function in TTP is not as affected as in malignant hypertension. Renal manifestations in TTP are hematuria and proteinuria, whereas renal manifestations in malignant hypertension include a rise in creatinine, BUN, hematuria, and proteinuria.

Discussion

We discussed the case of a patient with acquired TTP who presented with and was initially treated as a case of hypertensive emergency. TTP is a rare and fatal hematological disease, with an annual prevalence of approximately 10 cases/million people and an annual incidence of one new case/million people.

Acquired TTP, as seen in this patient, is caused by the presence of autoantibodies against ADAMTS13. This metalloprotease is essential in the cleaving process of the von Willebrand factor (VWF). Without ADAMTS13, ultra-large VWF multimers with bounded platelets persist in the bloodstream, leading to thrombocytopenia. This leads to the accumulation of microthrombi in the capillaries that in turn lead to hemolysis. This also affects the blood flow in the microcirculation that causes end-organ damage, mainly in the central nervous, renal, and cardiovascular systems.

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Because of the difference in pathogenesis, the thrombocytopenia seen in TTP is much more severe (platelets of <15,000/mm³) than the one seen in malignant hypertension (platelets of around 60,000/mm³). Also, renal function in TTP is not as affected as in malignant hypertension. Renal manifestations in TTP are hematuria and proteinuria, whereas renal manifestations in malignant hypertension include a rise in creatinine, BUN, hematuria, and proteinuria.

There are two types of acquired TTP: idiopathic and non-idiopathic. Non-idiopathic TTP is seen in about 50% of patients diagnosed with this disorder. The most frequent conditions associated with TTP, serving as triggers, are as follows: autoimmune disorders (e.g., systemic lupus erythematosus), viral and bacterial infections, cancer, pregnancy, and drugs (e.g., cyclosporine, quinine, clopidogrel). Idiopathic TTP is seen in the remaining 50% of cases.
Diagnosing TTP presents a challenge to physicians since it has an ample array of symptoms and they may not be all evident in the initial consultation. The gold standard in diagnosing TTP is the ADAMTS13 activity assay. If its activity is less than 10%, it is diagnostic for TTP. Severe thrombocytopenia and hemolysis must also be present to confirm the diagnosis of TTP. A helpful tool in the prompt diagnosis of TTP when ADAMTS13 activity assay is not readily available is the PLASMIC score. It has been shown to be useful in the accurate diagnosis and in predicting the prognosis of TTP (Table 1). A PLASMIC score equal to or greater than 6 is highly predictive of ADAMTS13 deficiency and necessitates immediate treatment [4]. The key differences between TMA secondary to hypertension and acquired TTP are as follows: TMA secondary to malignant hypertension will be seen with extreme hypertension (diastolic blood pressure of >130 mmHg), a past medical history of uncontrolled hypertension, clear renal dysfunction, and moderate thrombocytopenia. A patient with TTP that presents with hypertension will not have retinal findings and will also not recover from thrombocytopenia after the management of hypertensive crisis [7].

| The PLASMIC score - modified | Points |
|-------------------------------|--------|
| Platelet count <30,000/10^9/L | 1      |
| Hemolysis variables present (increased reticulocytes, increased LDH, decreased haptoglobin, increased indirect bilirubin) | 1 |
| No active cancer              | 1      |
| No history of solid organ or stem cell transplant | 1 |
| MCV <90 fl                   | 1      |
| INR <1.5                     | 1      |
| Creatinine <2.0/mg           | 1      |

**TABLE 1: Modified PLASMIC score**

LDH: lactate dehydrogenase; MCV: mean corpuscular volume; INR: international normalized ratio

A presumptive diagnosis is made based on the presence of microangiopathic hemolytic anemia and thrombocytopenia. Even though the presence of severe deficiency of ADAMTS13 activity or an inhibitor to it confirms the diagnosis of acquired TTP, it is not essential for starting prompt treatment [8]. TTP is a medical emergency, and PEX remains the treatment of choice and should be started as soon as possible. Plasma exchange has been associated with a decrease in mortality from 80-90% to 10%. During this procedure, plasma is removed from the affected patient and replaced by a donor’s plasma. In the course of the therapy, autoantibodies and immune complexes are removed from the affected patient’s plasma, and ADAMTS13 and VWF-multimers analysis (MM) of normal composition are replaced [5]. PEX is performed daily until features of organ involvement are resolved, platelet count has recovered to levels >150,000/mm^3 for two consecutive days, and hemolysis has stopped (seen in decreased levels of LDH). Immunosuppression with corticosteroids (methylprednisolone 1 mg/kg/day) is frequently used along with PEX, in order to control the autoimmune response and to improve the efficacy of plasma exchange therapy. Prophylactic platelet transfusion can worsen the symptoms of TTP and therefore should be reserved for severe cases [9].

Rituximab, a monoclonal antibody, is usually added to the therapy regimen in patients with refractory TTP, given its benefit in suppressing the production of IgG antibodies against ADAMTS13, with improved platelet recovery and lower relapse rate (relapse defined as thrombocytopenia after more than 30 days of stopping PEX). The addition of rituximab has been shown to induce remission in patients for more than 19 months [10]. New therapeutic approaches are being developed, such as an anti-VWF agent called caplacizumab, to be used subcutaneously, which causes suppression of VWF for up to 48 hours. Several studies have demonstrated reduced time to platelet normalization, exacerbations, and relapse rates [9].

**Conclusions**

TTP is a life-threatening hematologic disorder that requires prompt diagnosis and immediate treatment with PEX and maintenance treatment with rituximab to decrease the chances of relapses. Although patients with TTP can present with hypertensive-like crises, it is imperative not to delay treatment with PEX. Using diagnostic parameters such as the PLASMIC score can aid in differentiating between these two conditions. Due to the uncommon nature of the illness and a lack of physician awareness, patients are often not promptly diagnosed, which can negatively affect their prognosis.

**Additional Information**
Disclosures

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