Thrombotic Thrombocytopenic Purpura

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Introduction

Thrombotic microangiopathy comprises a spectrum of clinical and laboratory findings including microangiopathic hemolytic anemia, thrombocytopenia, and thrombosis of capillaries and arterioles. Platelet and hyaline thrombi with complete or partial occlusion of these vessels are integral histopathological findings. These findings are seen irrespective of cause and organ involved. Pathogenesis and prognosis may differ for different forms of microangiopathy and clinical features often overlap.

Thrombotic angiopathy may be classified in a variety of ways based on various factors. A classical way of classification is dividing the spectrum of disease as Classical TTP, HUS or disease entity with severe ADAMTS 13 deficiency. Though the above classification favors dominant organ specific involvement, there is a large overlap in clinical features amongst these various groups making such a distinction inadequate. Though clinically-pathologically TTP and HUS overlap, both these entities appear to have distinct pathogenesis. Evidence shows a high incidence of ADAMTS13 deficiency in patients with clinically TTP but not HUS [1]. On the other hand familial and sporadic cases of HUS have been associated with deficiency of complement regulatory proteins, particularly factor H or membrane cofactor protein. Though TTP and HUS affect many of the same organ systems, the frequency with which they do so differs. Even histopathologic features of TTP and HUS are also distinct. Plasma exchange is highly effective for idiopathic TTP [2]. But is usually ineffective for sporadic HUS. Thus for a better understanding it seems reasonable to continue to distinguish these clinical entities while recognizing extensive overlap in clinical features.

Classical TTP is associated with dominant neurological abnormalities. Renal involvement is common but with minimal instances of oliguric acute renal failure. This entity is usually associated with acquired autoantibodies that inhibit disintegrin and metalloprotease thrombospondin type 1 repeats (ADAMTS 13). Inherited deficiency of ADAMTS13, also known as Upshaw-Schulman syndrome, is a congenital form of TTP. Features of microangiopathy with acute renal failure (oliguric/ anuric) is considered by some to represent HUS. HUS may be classified as Diarrhea associated HUS (also known as PRIMARY/ TYPICAL /EPIDEMIC HUS) due to identifiable enteric infection with Shigatoxin producing bacilli, most commonly Escherichia coli O157:H7 [3]. The Diarrhea NOT associated HUS (secondary/atypical/ sporadic HUS) comprises a group that is not associated with Shiga Toxin producing organism in patients without an obvious predisposing condition. In patients with severe ADAMTS 13 deficiency states both renal and neurological involvement is uncommon. On the other hand, some patients present with both neurological and renal involvement and are grouped into a broad TTP-HUS category.

There are other secondary causes of Micrangiopathy. These include cancer, infections, stem cell transplant, chemotherapy and variety of drugs. In most of these cases ADAMTS13 levels are normal and management is basically aimed at correcting the primary cause.

Epidemiology

Before the plasma exchange era, survival of patients with TTP was only 10%. When plasma exchange was reported to increase survival to 78%, compared with 51% survival for patients treated with plasma infusion. The diagnosis of TTP became an indication for plasma exchange treatment. Oklahoma TTP-HUS Registry is a population based cohort registry of patients maintained since 1989 based on request received for plasma exchange treatment [4]. As the registry is based on request for plasma exchange, many children with typical HUS are not included in the registry because they are not treated with plasma exchange. Because it is an all inclusive study with no patients excluded when plasma exchange request was received, incidence data was calculated in a recent paper by Terrel [5] ADAMTS 13 activity and inhibitors of ADAMTS 13 activity are measured on all samples whose samples activity was less than 20%. Severe ADAMTS13 deficiency is defined as activity less than 10% [5].

The whole registry is divided into three categories: all patients with clinically suspected TTP-HUS, patients with idiopathic TTP-HUS, and patients with severe ADAMTS-13 deficiency. ADAMTS 13 activity and inhibitors of ADAMTS 13 activity are measured on all samples whose samples activity was less than 20%. Severe ADAMTS13 deficiency is defined as activity less than 10% [6].

The age-sex-race standardized annual incidence rates were per 10(6):

11.29 For all patients with clinically suspected TTP-HUS
4.46 For patients with idiopathic TTP-HUS
1.74 For patients with severe ADAMTS-13 deficiency (< 5% activity)

In all three categories, the incidence rates were greater for women and for patients with African ancestry. For patients with severe ADAMTS-13 deficiency (< 5 % activity), the age-sex standardized incidence rate ratio of blacks to non-blacks was 9.29. This greater incidence in black females is comparable with the increased risk of autoimmune disease within this subgroup of population.

Another recent review article estimated the incidence of acute ITP in children and adults as below: [7].

For children: 1.9 - 6.4 per 10(5) children/year.
For adults: 3.3 per 10(5) adults/year.

Regional variation may occur in incidence as these data were...
mostly collected in Europe. Other evidence shows a peak incidence in the fourth decade and an association with obesity as well.

Clinical Features

The classical presentation of TTP is thrombocytopenia and hemolytic anemia with fragmentation red cells.

The clinical picture may be varying with 10-40% of patients complaining of an upper respiratory tract infection in preceding weeks. Patients may also present with nonspecific symptoms (malaise, fatigue, fever) lasting weeks that are unresponsive to symptomatic treatment. In this situation, the diagnosis could be confused until these non specific symptoms become severe or other organ system get involved. Neurologic involvement could range from headache and confusion to somnolence, seizures, aphasia, or coma. Symptoms may even fluctuate in severity that may be attributed to the repetitive formation and dissolution of microthrombi in the cerebral microvasculature.

Platelet counts may go below 20,000/mL, with mucou-cutaneous bleeding. Prothrombin time, partial thromboplastin time, and fibrinogen levels are usually normal or only mildly altered. Fibrinogen degradation products occur in 50% of patients. Hematuria, proteinuria, granular or red cell casts, and mild azotemia may be the presenting features of TTP but anuria and renal failure are uncommon. Abdominal pain with or without pancreatitis, and electrocardiographic abnormalities have also been observed. Below is a recent chart from Oklahoma TTP-HUS Registry with clinical features of patients with severe ADAMTS13 deficiency. It is important to note that 31% of the patients had only minor symptoms (confusion, headache) whereas 34% had no neurologic abnormalities. Many developed new neurologic abnormalities during the course of treatment.

Mechanism of Microvascular Thrombosis in TTP/Pathogenesis

Pathogenesis of thrombus formation in TTP

The pathogenesis of thrombi formation is more clearly understood in models with severe ADAMTS13 deficiency whether due to genetic mutations or inhibitory autoantibodies.

Microvascular thrombi consist of von-willbrand factor and platelets. Understanding the interaction between vWF and platelet is crucial for knowing why thrombi occur in TTP.

What is the role of vWF?

It is a glycoprotein in the plasma made of several multimers which are connected by disulfide bonds. The molecular weight could reach millions of Daltons. The vWF serves as a carrier for factor VIII. It also has a major role in mediating platelet adhesion and aggregation at site of endothelial Injury. It binds to platelet glycoprotein receptors Ib/IX/V and αIIbβ3. It is synthesized in endothelial cells and stored in Weibel–Palade bodies as multimers. Upon activation the multimer will adhere to the unusual large multimer due to their activated glycoprotein Ib/IIIa complexes leading to occlusive thrombi.

The ADAMTS13 gene

ADAMTS13 cleaves vWF polypeptide whenever this normally cryptic bond is rendered accessible by circulatory shear stress. This rendered endothelial vWF polymer into smaller multimers that are conformationally less flexible and less adhesive. In the absence of ADAMTS13 vWF multimers are conformationally unfolded but are not cleaved. This results in accumulation of hyperactive forms of vWF leading to platelet aggregation and microvascular thrombosis.

Clinically, platelet thrombosis does not occur when ADAMTS13 is greater than 10% of normal values. But there is no lower threshold level for ADAMTS13 below which microvascular thrombosis invariably occurs. Various factors play a role in determining the level of ADAMTS13 protein including circulatory shear stress, platelet receptor levels, vWF and other unknown factors.

Laboratory Findings

The most important hematological parameters for the diagnosis of TTP/HUS are thrombocytopenia and microangiopathic hemolytic anemia. Thrombocytopenia is usually more severe in cases without predominant renal involvement; the platelet count are usually as low as < 20,000/mm³ at presentation [8]. Microangiopathic Hemolytic Anemia (MAHA) is a major diagnostic criterion for TTP and it is defined by the presence of schistocytes in the peripheral smear along with evidence of hemolysis. It was initially not clear as to how many schistocytes are required in peripheral smear for the diagnosis of TTP induced MAHA. But now it is concluded that more than 1% schistocytes, in the absence of other causes of thrombotic angiopathies, is highly suggestive of TTP-HUS [9] (Figure 1). Features of hemolytic anemia include increased reticulocyte count, increased lactate dehydrogenase levels (LDH), elevated indirect bilirubin; low serum haptoglobin levels with the intravascular hemolysis and increased levels of free plasma hemoglobin in severe cases [10-12]. Direct antiglobulin test is usually negative suggestive of the non-immune hemolytic process in TTP-HUS except in cases of neuraminidase-associated HUS [13,14]. Proteinuria, microscopic hematuria, granular or red cell casts may be seen with renal involvement. TTP rarely presents with severe renal failure and anuria, although up to 60% of TTP cases have renal dysfunction [8,15].

Measurement of ADAMTS 13 is necessary in the work up for TTP but it is not a diagnostic criterion because the levels may be normal in many patients that otherwise have all the features of TTP [16,17]. ADAMTS 13 levels less than 10% is considered specific for TTP, it can often be low in liver failure, sepsis, chronic kidney disease, connective tissue disorders, pregnancy and various other inflammatory conditions but the levels are not as low as less than 10% of the normal in these
If cultures or serology come back positive for exchange, plasma infusion, cryoprecipitate or solvent-detergent detected and confirmed. Plasma therapy can be in the form of plasma cases in children. But for all others with TTP plasma therapy should cases, treatment would be supportive similar to diarrhoea positive HUS producing diarrhoea and the patient does not have severe neurological and in the absence of any other causes of thrombotic angiopathies) recommended to start plasma therapy in all adult patients with suspected to make a diagnosis of TTP or HUS at presentation and thus it is survival in more than 80-90% of the patients [26]. It is often difficult initiation of plasma therapy can be fatal. Plasma therapy has improved conditions [18-20]. Patients with persistent anti-ADAMTS13 antibodies or undetectable levels of ADAMTS13 are at a higher risk for recurrence [16]. However, measurement of ADAMTS 13 during the remission is not routinely recommended [21]. The low levels are indicative of either congenital or acquired deficiency. This acquired deficiency can be a result of antibodies or inhibitors to ADAMTS 13. There are many assays used to detect these antibodies but there is no standardized and reliable assay widely available [15]. In cases of suspected diarrhea positive HUS, the diagnosis is based on detection of E.coli 0157:H7 or Shiga-toxin producing bacteria on sorbitol-MacConkey agar [22]. There are serological tests to detect the antibodies against the toxin and they are more sensitive compared to cultures but so far they are used only in research laboratories [23,24]. Patients with suspected atypical diarrhoea negative HUS can be tested for complement regulatory proteins like Complement Factor H (CFH), Complement Factor I (CFI) and Membrane Receptor Protein (MCP). Tests to identify genetic mutations for these complements are usually performed when the levels of the complements are low [25].

Treatment

Plasma therapy

Plasma therapy remains the mainstay of treatment. A delay in the initiation of plasma therapy can be fatal. Plasma therapy has improved survival in more than 80-90% of the patients [26]. It is often difficult to make a diagnosis of TTP or HUS at presentation and thus it is recommended to start plasma therapy in all adult patients with suspected TTP or HUS (thrombocytopenia, microangiopathic hemolytic anemia with or without fever, neurological, renal manifestations and in the absence of any other causes of thrombotic angiopathies) [21,27]. If cultures or serology come back positive for E.coli or toxin producing diarrhoea and the patient does not have severe neurological manifestations then plasma therapy can be discontinued [27]. In such cases, treatment would be supportive similar to diarrhoea positive HUS cases in children. But for all others with TTP plasma therapy should be continued unless another etiology for the thrombotic angiopathy is detected and confirmed. Plasma therapy can be in the form of plasma exchange, plasma infusion, cryoprecipitate or solvent-detergent treated plasma. There are many studies to determine the efficacy of plasma exchange versus infusion but there is no proven benefit of one over the other [28]. Nonetheless, plasma exchange is preferred because larger amount of plasma can be given to the patient without fluid overload and it also allows more rapid removal of anti-ADAMTS 13 antibodies compared to plasma infusion but there are no studies to prove this benefit [27]. The goal of plasmapheresis is to exchange at least 1 to 2 plasma volumes per day (40-50ml/kg) [29,30]. Solvent-detergent treated plasma has lower rates of allergic reactions, however; ADAMTS13 concentration in solvent-treated plasma is 20% lower than that of fresh frozen plasma [31,32]. Efficacy of the plasma therapy is assessed by monitoring the LDH, platelet count and neurological signs. The therapy should be continued for at least two to three days after the platelet count and LDH are normal [33,34].

Plasma therapy is also beneficial in some secondary forms of TTP like ticlopidine or clopidogrel induced TTP or pregnancy associated TTP but its role in chemotherapy associated TTP and bone marrow transplantation associated TTP has not resulted in improved outcome [35,40].

Antiplatelet therapy

Antiplatelet agents are not recommended in the treatment of TTP. It does not change the outcome and survival and may even increase the chance of bleeding in some cases with severe thrombocytopenia [41,42].

Corticosteroids, immunosuppressive therapy and monoclonal antibodies

Corticosteroids as the sole agent to prevent recurrences of TTP have not proven to be of benefit. However, patients with ADAMTS 13 deficiency are started on steroids alone as recommended by many experts [43,44]. Combination of steroids with immunosuppressive drugs like vincristine, azathioprine and monoclonal antibody rituximab is used to prevent relapse in atypical HUS causes deficiency of complement proteins [45,46]. Rituximab by itself has been shown to significantly reduce the recurrence in cases of TTP caused by anti-ADAMTS13 antibodies. It has been used in patients that have failed the combination of plasmapheresis plus steroids. It can also be used as prophylaxis to prevent relapse in TTP patients with presence of anti-ADAMTS13 antibodies [47-50]. Anecdotal case reports with eculizumab, anti-C5 monoclonal antibody, in the use atypical HUS have been reported but clinical trials have yet to prove its role and benefit [51,52].

Splenectomy

Previously, it was considered in conjunctions with other modes of treatment and induced remission in 50% cases but now it is mainly reserved for cases that have not responded to plasmapheresis and immunosuppressive treatment [42,53].

Transplantation

Patients that progress to end stage renal disease are good candidates for renal transplantation but HUS can recur in the renal allograft [54]. This recurrence in the renal allograft may be avoided if liver transplantation is also done. Liver produces the complements implicated in the pathophysiology of recurrent HUS. But, this is not without its own downsfalls and patients need to be carefully selected for isolated renal versus renal and liver transplantation in case HUS cases caused by deficiency of complements and membrane cofactor protein [55,56].


Figure 1: Showing schistocytes in peripheral smear.


