

Unresponsive Thrombotic Thrombocytopenic Purpura (TTP): Challenges and Solutions

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Abstract: Thrombotic thrombocytopenic purpura (TTP) is a thrombotic microangiopathy secondary to a severely decreased A Disintegrin And Metalloprotease with ThromboSpondin type 1 repeats 13 (ADAMTS13) activity, resulting in the formation of widespread von Willebrand factor - and platelet-rich microthrombi. ADAMTS13 deficiency is mainly acquired through anti-ADAMTS13 autoantibodies in adults. With modern standards of care, unresponsive TTP has become rarer with a frequency of refractory/relapsing forms dropping from >40% to <10%. As patients with unresponsive TTP are at increased risk of mortality, prompt recognition and early therapeutic intensification are mandatory. Therapeutic options at the disposal of clinicians caring for patients with refractory TTP consist of increased ADAMTS13 supplementation, increased immunosuppression, and inhibition of von Willebrand factor adhesion to platelets. In this work, we focus on possible therapies for the management of patients with unresponsive TTP, and propose an algorithm for the management of these difficult cases.

Keywords: thrombotic thrombocytopenic purpura, refractory, relapsing, rituximab, caplacizumab

Introduction

Thrombotic thrombocytopenic purpura (TTP) is a thrombotic microangiopathy (TMA), characterized by the spontaneous formation of thrombi in the microcirculation.¹ The diagnosis of TMA relies on the association of hemolytic mechanical anemia, peripheral thrombocytopenia and various signs of visceral ischemia ascribable to microvessel thrombosis. In TTP, a severe ADAMTS13 (A Disintegrin And Metalloprotease with ThromboSpondin type 1 repeats 13) activity deficiency is responsible for an excess of unusually large von Willebrand factor (vWF) multimers. Under high shear stress conditions associated with the microcirculation, these vWF multimers spontaneously aggregate with platelets, leading to the different signs of the disease.^{2,3} Of note vWF and ADAMTS13 play a key role in various other thrombotic conditions and in cardiovascular diseases.^{4–7}

ADAMTS13 deficiency can be secondary to mutations or polymorphisms of the ADAMTS13 gene (9q34) in hereditary TTP (Upshaw Schulman syndrome, Online Mendelian Inheritance in Man number, 274150)⁸ or can be acquired through anti-ADAMTS13 antibodies in the majority of adult forms (immune TTP).⁹ TTP is a rare disease with an incidence rate estimated as 1.5–6/million/year, affecting mostly young women.^{10–12} It can be a devastating condition, leading to death in more than 80% of cases if not treated promptly with plasmatherapy.¹³

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Modern standards of care for suspected cases of TTP include therapeutic plasma exchanges (TPE), systemic corticosteroids, anti-CD20 monoclonal antibody and anti-vWF domain A1 nanobody.¹⁴ With this therapeutic schedule, a vast majority of patients will reach TTP remission and mortality rate is less than 5%.¹⁵ However, in some patients, first line therapy fails and additional treatments are required in emergency.¹⁶ The aim of this review is to explore current options at the disposal of clinicians faced with cases of unresponsive TTP in adults.

Challenge 1: Recognize Unresponsive TTP

Refractory TTP was first defined in the 2003 British Committee for Standards in Haematology guidelines as cases who fail to reach a platelet rate above 150 G/L after 7 TPE sessions.¹⁷ In the 2017 consensus on standardization of terminology on TTP,¹⁸ response to therapy was defined as a platelet rate above 150 G/L and Lactate De Hydrogenase (LDH) level <1.5 the upper norm of the laboratory after 5 TPE sessions. Remission consists of a durable response 30 days after the end of therapy (at that time, last TPE session). The 2021 consensus introduced several new notions.¹⁹ First, clinical response now also requires the absence of signs of new or progressive signs of ischemic organ injury, in addition to platelet and LDH rates improvement. Second, it introduces the notion of biological recovery, based on ADAMTS13 activity normalization, underpinned by the fact that TTP patients failing to reach ADAMTS13 activity recovery at the end of treatment are at very high risk of clinical relapse.

Unresponsive TTP in this work will, though, include cases presenting with:

1. Failure to reach clinical response after 5 TPE sessions (non-response or insufficient clinical response).
2. Occurrence of any new clinical or biological sign of TMA while in therapy, or within 30 days following the end of treatment (TPE or caplacizumab) (exacerbation).
3. Re-occurrence of TTP after a period of remission (relapse).

Figure 1 illustrates these definitions, according to the timeline of the disease.

Challenge 2: Confirm TTP Refractoriness Ascertain the Diagnosis of TTP

TTP is sometimes a difficult clinical diagnosis to make.²⁰

First, TTP may be mistaken with other diseases that do not belong to the TMA syndrome. Examples of such diseases that may present as TMA and usually do not respond to standard TTP therapy are provided in Table 1.

TMA syndrome consists of a number of different diseases and TTP is far from being the most frequent (estimated frequency being 3% of all TMA syndromes).²¹ ADAMTS13 activity study is of critical importance to differentiate TTP from other TMA syndromes with potentially similar clinical presentations (secondary TMA, complement-mediated hemolytic uremic syndrome (HUS), shigatoxin associated HUS, etc). Laboratory tests should be performed on blood samples retrieved before the onset of plasmatherapy. This way, a diagnosis of refractory TTP can readily be excluded if

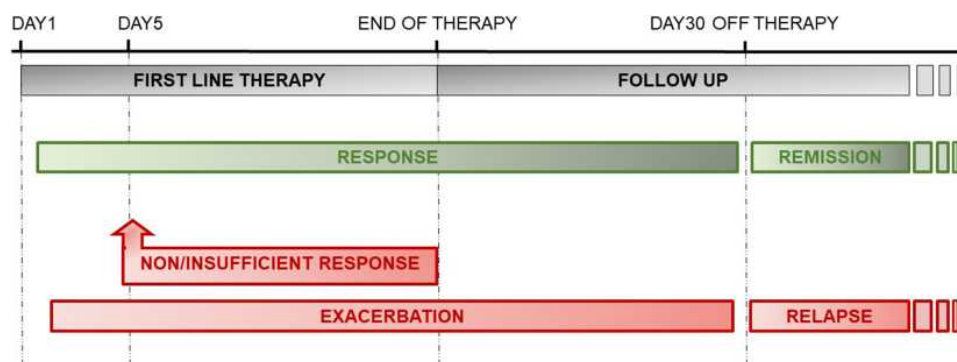


Figure 1 Timeline of acute TTP management with favorable (green) and unfavorable (red) outcomes. Non/insufficient response is defined as failure to reach platelet rate >150G/L and/or LDH rate <1.5N and/or occurrence of new/progressive signs of ischemic organ involvement after 5 TPE sessions; Exacerbation is defined as recurrence of any sign of TTP after a phase of clinical response, up to 30 days after the end of therapy; Remission is defined as clinical response maintained over 30 days after cessation of therapy; Relapse is defined as the recurrence of any sign of TTP after remission (ADAMTS13 activity decrease or clinical/biological signs of thrombotic microangiopathy).

Table 1 TMA Differential Diagnosis

	Causes of Cytopenia	Causes of Organ Dysfunction	Evocative Signs
Severe vitamin B9/B12 deficiency	Ineffective erythropoiesis and thrombopoiesis with intra-medullary destruction	Tissular ischemia secondary to profound anemia, bleedings secondary to thrombocytopenia, neurological symptoms secondary to vitamin deficiency	Macrocytosis, unregenerative anemia, megaloblastosis on bone marrow smear
Hematological malignancies/metastatic cancers	Bone marrow involvement by multiple myeloma/cancer metastasis Ineffective erythropoiesis and thrombopoiesis in myelodysplastic syndrome	Tissular ischemia secondary to profound anemia, bleedings secondary to thrombocytopenia, organ involvement of malignancies	Systemic symptoms of cancer, unregenerative anemia, neoplastic cells on bone marrow examination
Evans syndrome	Erythrocytes and platelets destruction secondary to autoantibodies	Tissular ischemia secondary to profound anemia, bleedings secondary to thrombocytopenia	Positive DAT
Hemophagocytic syndrome	Blood cells and precursors phagocytosis by activated macrophages	Cytokine storm, organ involvement of the underlying cause of hemophagocytic syndrome	Fever, HLH2004 criteria

Abbreviations: DAT, direct antiglobulin test; HLH2004, hemophagocytic lymphohistiocytosis.

patients displayed a detectable activity at onset of therapy and eventually fail first line therapy.^{22,23}

Search for Confounders

Dissociated clinical and biological evolution is unusual in TTP patients. In the case of persistent thrombocytopenia and/or anemia in patients with satisfying clinical evolution on first-line therapy, clinicians should look for alternate diagnoses others than refractory TTP. Alternative causes of persistent or new organ dysfunctions should also systematically be sought, especially if hematological involvement is improving. Examples of confounders are provided in Table 2.

Nevertheless, clinicians should always keep a high index of suspicion and if the cause of clinical or biological degradation is uncertain, patients must be considered as having refractory TTP and prompt therapeutic intensification must be performed.²³

ADAMTS13 activity control in patients on therapy may also prove useful, as organ dysfunctions appearing/persisting in patients with normalized ADAMTS13 activity are very unlikely to be related to refractory TTP.

Challenge 3: Choose Therapeutic Targets for Unresponsive TTP

First-Line Treatment

Figure 2 displays the different therapeutic targets available for the management of TTP.

ADAMTS13 Supplementation: Therapeutic Plasma Exchanges

TPE still are the cornerstone of therapy for patients with acquired TTP. As part of first line therapy they are performed at a volume of 1.5 estimated plasma volume (EPV), daily, with healthy donor plasma or derived products (ie cryosupernatant) until patients reach clinical remission.^{17,23,24} The main mechanisms of action of TPE in TTP are thought to be both the administration of massive amounts of functional ADAMTS13 and the removal of anti-ADAMTS13 autoantibodies and unusually large vWF multimers.²⁵ TPE should be preferred to plasma infusions in patients suspected of having TTP, as they are associated with a better outcome.²⁶ EPV may be calculated with the following formula:

$$\text{EPV} = \text{Total Blood Volume (TBV)} \times (1 - \text{Hematocrit})$$

TBV is estimated using either Nadler's formula (based on gender, height and weight) or Gilcher's rule of 5 (based on gender and morphotype).²⁷ With simplifications, EPV is often considered 40 mL/kg in critical care patients.

Immunomodulation

Steroids

Corticosteroids were the first immunomodulating drugs used in patients with TTP and they are part of the first line therapeutic schedule of patients with a suspicion of acquired TTP since 2003.^{17,28} In one study, the use of corticosteroids (prednisone 200 mg/d) as standalone

Table 2 Confounders for TTP Refractoriness

Manifestations	Potential Confounders	Proposed Reaction
Lack of response to first line therapy	Wrong diagnosis: other TMA syndrome (Complement mediated HUS, Shigatoxin-associated HUS, HELLP syndrome, secondary TMA)	- Verify ADAMTS13 study was performed - Verify appropriate TMA diagnostic workup was performed
	Incorrect/sub-optimal first-line therapy	- Verify quality of TPE procedure (substitution fluid = FFP, adequate venous access, no plasma intolerance, no catheter site thrombosis, etc) - Verify type, dosage and timing of immunomodulatory drugs administration are adequate (as soon as possible after last TPE session) - Verify quality of supportive care (blood pressure control, vitamin supplementation, adequate red cell transfusion, etc)
Dissociated clinical/biological response	Alternate diagnosis for clinical/biological anomaly: - Thrombocytopenia: sepsis, HIT, central, etc - Anemia: hemorrhage, iron/vitamin B9 deficiency, etc - Neurological symptoms: citrate overdose, sepsis, ICU delirium, etc - Renal function impairment: acute tubular necrosis, renal involvement of associated disease, etc	- Control ADAMTS13 activity - Perform differential diagnosis workup - Consider patient has refractory TTP until differential diagnosis is established

Abbreviations: TMA, thrombotic microangiopathy; HUS, hemolytic uremic syndrome; HELLP, hemolysis, elevated liver enzymes, low platelets; TPE, therapeutic plasma exchange; FFP, fresh frozen plasma; ICU, intensive care unit; HIT, heparin induced thrombocytopenia.

therapy in mild forms of TTP with no sign of neurological involvement (hematological signs only) induced durable remission in 52% of cases (28/54).²⁹

Anti-CD20 Monoclonal Antibodies

Anti-CD20 monoclonal antibodies target B-lymphocytes implicated in the production of anti-ADAMTS13 autoantibodies in a vast majority of adult patients with acquired TTP.^{30,31} They were developed for the treatment of B-cell lymphomas and secondarily used in many autoimmune diseases. Most of the literature has described the use of rituximab, a chimeric mouse-human monoclonal anti-CD20 antibody, in these indications.^{32,33} Severe adverse side-effects include infusional reactions, serum sickness, hypogammaglobulinemia and infections including Hepatitis virus B reactivations and progressive multifocal leukoencephalopathy (due to JC-BK virus central nervous system infection). These manifestations were mostly described in patients with lymphomas receiving cytotoxic polychemotherapy along with rituximab.³⁴ In patients with TTP, rituximab seems to present a very favorable security profile.³⁰

Anti-CD20 monoclonal antibodies are recommended in case of refractory or relapsing TTP.³⁵ They are also increasingly used as part of first-line therapy, during the first 3 days after diagnosis.^{15,28,35} The dose varies from 375 mg/m² to 100–1000 mg fixed doses, administered

weekly³⁶ or on days 1-4-8-15,³⁷ considering TPE will remove a significant part of the monoclonal antibodies.³⁸

This recommendation is not based on randomized controlled studies but on case-control studies and case series.^{30,39} In TTP, authors have shown rituximab may improve time to clinical remission,³⁷ improve time to intensive care discharge and decrease the risk of TTP relapse during the year following administration.⁴⁰ There also seems to be a benefit with early use of rituximab (≤ 3 days after diagnosis) in comparison to late administration in terms of time to clinical remission.⁴¹

As the efficacy of rituximab is expected around 2 weeks after the drug's administration, early rituximab infusions are now recommended as a first line therapy in TTP, even before the results of ADAMTS13 studies are known.²⁸ If rituximab is used as second-line therapy, it is mandatory to continue first-line therapy (at least daily TPE) and adjunct fast-acting therapies may also be required. The number of rituximab injections may be adapted according to B-cell depletion monitored in blood.⁴²

Rituximab can also be used in patients that present with an ADAMTS13 activity decrease while in clinical remission of TTP. In this context, rituximab, as a single agent therapy in patients with no clinical or biological signs of TMA, may induce ADAMTS13 activity normalization and prevent progression to TTP clinical relapse (preemptive treatment).^{43–46} In this setting, sub-cutaneous administration is possible.⁴⁷

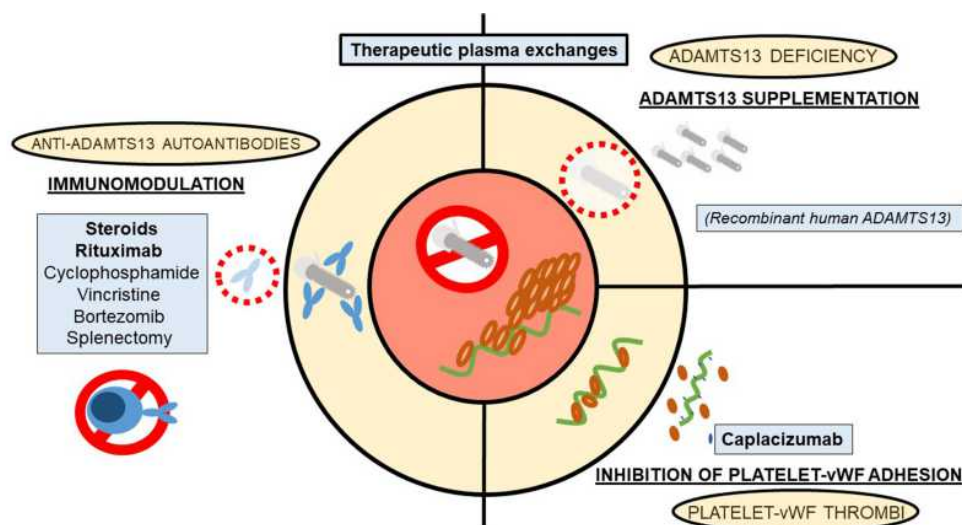


Figure 2 Pathophysiology, potential therapeutic targets and available therapy for the management of TTP.

Clinicians should be cautious about the apparition of anti-rituximab antibodies in patients treated with multiple courses of rituximab. These autoantibodies have been associated with infusional reactions, serum sickness and loss of rituximab efficiency.⁴⁸ Patients who fail or become intolerant to rituximab may benefit from treatment with other anti-CD20 antibodies (eg obinutuzumab⁴⁹ or ofatumumab).⁵⁰

VWF Targeting Agents: Anti-vWF Domain A1 Nanobodies

Caplacizumab is an anti-vWF domain A1 nanobody, impairing the adhesion of vWF to platelet receptor gp Ib-IX-V. The therapeutic schedule includes a 10 mg intravenous loading dose administered before a TPE session followed by subcutaneous 10 mg injections after each TPE and up to 30 days after the end of plasmatherapy. In two randomized controlled studies, the addition of caplacizumab to standard of care shortened time to clinical remission in TTP patients.^{51,52} Real world data are also in favor of caplacizumab's early administration in patients suspected of having TTP.^{15,53} Caplacizumab is approved for the treatment of acquired TTP²⁸ but the cost-effectiveness of frontline caplacizumab administration in all TTP patients remains uncertain.^{54,55} Adverse side effects consist of hemorrhage, usually non severe.

Caplacizumab may be used in patients with refractory TTP who did not receive this drug frontline or who present with relapses after caplacizumab withdrawal.

Due to the mechanism of action of caplacizumab, that is the inhibition of the early stage of thrombi formation, this drug should be administered as soon as possible after the diagnosis of TTP.²⁸ It is a “time buying” or suspensive therapy that has no effect on anti-ADAMTS13 autoantibodies production, explaining a relatively high rate of relapses in patients that keep a severely decreased ADAMTS13 activity at the end of treatment. In these patients, intensification of the immunosuppressive regimen and caplacizumab continuation are recommended until ADAMTS13 activity is restored.

Unresponsive TTP Treatment

Therapeutic options for patients with unresponsive TTP are summarized in Table 3.

First-Line Treatment Adaptation ADAMTS13 Supplementation

In patients that do not respond to daily TPE or clinically deteriorate while on first line therapy, twice-daily TPE have been proposed, with a volume of 1 to 1.5 EPV per 12 hours.^{56,57} This costly and time-consuming procedure has not been evaluated in controlled studies but may prove helpful in patients with very severe forms of TTP, high titres of anti-ADAMTS13 autoantibodies and in whom immunosuppressive interventions have not yet reached their efficacy (during the first days of treatment). Clinicians must anticipate significant removal of therapeutic drugs with high molecular weights (ie monoclonal antibodies) and/or important albumin binding in patients receiving this therapeutic schedule.²⁵

Table 3 Therapeutic Options Currently Available for Unresponsive TTP Management

Molecule/ Procedure	Dosage	Time to Response	Considerations
First-line treatment adaptation			
High dose steroid pulse	1000 mg prednisone equivalent/24h, IV	<7 days	May be included in first-line therapeutic schedule, 1 to 3 pulses
Twice daily TPE	1–1.5 EPV/12h	<7 days	1 to 7 sessions (time-consuming)
Caplacizumab	10 mg, IV before first TPE then 10 mg/day, SC	<7 days	Usually included in first-line therapeutic schedule, duration of treatment 30 days after last TPE (and/or tailored by ADAMTS13 activity)
Rituximab	375 mg/m ² IV weekly for 4 weeks or D1-D4-D8-D15	>10 days	Fix doses (low or high, 100–1000 mg) may be used, number of injections may be adapted to CD19-B cells count monitoring at day 4 and after; administration of rituximab should take place immediately after a TPE session, SC route may be used for preemptive treatment
Second line therapy			
Vincristine	1.4 mg/m ² , maximum dose 2 mg, IV	<7 days	Usually 1 infusion (up to 1/week if necessary)
Cyclophosphamide	500–750 mg/m ² or 500 mg, IV	>7 days	Usually 1 infusion (up to 1/week if necessary)
Bortezomib	1–1.3 mg/m ² , IV or SC, D1-D4-D8-D11	<7 days	1 infusion may be sufficient
Splenectomy	N/A	< 7 days	Salvage therapy, laparoscopic technique may be preferred
N-acetylcysteine	300 mg/day continuous infusion, IV	<7 days	Limited clinical evidence in TTP, duration of therapy unknown
Ciclosporine A	300 mg/day PO or 2–3 mg/kg/day IV	< 7 days	Continue treatment after remission
Azathioprine	100 mg/day PO	Unknown	Very limited evidence in TTP
Intravenous Immunoglobulin	0.4 g/kg/day IV for 2–6 days	<7 days	Side effects may mimic TTP manifestations (acute kidney injury, headaches, thrombosis, etc)
Eculizumab	900 mg D1-D8-D15-D22 then 1200 mg/2–4 weeks	>7 days	Limited evidence in TTP; complement study should be performed to document potential associated alternate pathway anomalies, if TPE are maintained eculizumab has to be re-administered after each session

Abbreviations: EPV, estimated plasma volume; TPE, therapeutic plasma exchange; D, day.

Immunomodulation

As high dose pulses of steroids may be associated with improved outcomes in comparison with lower doses,⁵⁸ their use as second-line therapy in unresponsive cases of TTP may be proposed in patients not already undergoing treatment with corticosteroids or who received lower doses of steroids.

Second Line Therapy

Vincristine

Vincristine is a vinca-alcaloid used as anti-cancer chemotherapy or immunomodulating agents in various

indications. In TTP, vincristine may also play an inhibitory role on platelets glycoprotein receptors to vWF.⁵⁹

Patients with refractory TTP may respond to second-line therapy with 1.4 mg/m² (maximum dose 2 mg) vincristine intravenously. A single injection is usually administered, with an efficacy expected in less than a week and toxicities including cytopenias due to bone marrow depletion and peripheral neuropathy.^{22,59,60}

Cyclophosphamide

Cyclophosphamide is an alkylating agent used as anti-cancer chemotherapy and as an immunosuppressive

agent. It can be used in patients with refractory TTP with a single parenteral injection of 500 mg/m² or 750 mg/m², with an efficacy expected from one to two weeks after treatment and toxicities including bone marrow depletion.^{61,62} Cyclophosphamide may be particularly indicated in TTP cases associated with other autoimmune diseases, like systemic lupus erythematosus and Sicca syndrome.

Bortezomib

Bortezomib is a proteasome inhibitor approved for the treatment of multiple myeloma and other plasma cells disorders.⁶³ It may be a part of refractory TTP therapy at a dose of 1–1.3 mg/m², administered intravenously or subcutaneously, from 1 to 4 times (days 1-4-8-11) depending on response which can be expected in days following administration.^{64,65} Side-effects described in patients with refractory TTP are peripheral neuropathy, pulmonary and cardiac toxicities.

Other Immunomodulating Agents

Many other immunomodulating drugs have been tried in patients with refractory/relapsing TTP, including cyclosporine A,⁶⁶ azathioprine,⁶⁷ intravenous polyvalent human immunoglobulin⁶⁸ and eculizumab.^{69,70} Data concerning these therapies is scant, sometimes contradictory and limited to case reports or small case series, often pre-dating the advent of rituximab use.

N-Acetyl Cysteine

In clinical practice, N-acetyl cysteine is principally used as an antidote for acetaminophen poisoning.⁷¹ It has the capacity to disrupt the di-sulfite bonds linking vWF monomers, to disrupt vWF domain A1 and display an inhibitory effect on ADP- and collagen-induced platelet aggregation *in vitro* and in animal models.^{72–74} By these mechanisms, N-acetylcysteine could reduce the amount of unusually large vWF multimers found in TTP patients and reduce vWF-platelets adhesion.⁷⁵ It can be administered using the same protocol as in acetaminophen poisoning (continuous infusion of 300 mg/day), for an indeterminate duration (usually less than 1 week). Efficacy is expected in the days following N-acetylcysteine infusion. There is no significant side-effect of N-acetylcysteine described in TTP patients.

Splenectomy

Splenectomy has been proposed to patients with refractory or relapsing TTP, mainly before the rituximab era.^{61,76,77}

The presumed mechanisms of action of splenectomy in TTP are unclear, probably related to the removal of a major site of anti-ADAMTS13 autoantibodies production as well as a major site of platelet consumption. This procedure usually leads to prompt remission of the disease but can be difficult to perform in severely thrombocytopenic patients. Adverse side effects include surgical complications and thromboembolic disease.⁷⁶ Due to the risk of hemorrhagic complications, laparoscopic procedures should probably be preferred. Prophylactic platelet transfusion should be avoided whenever possible in the context of unresponsive TTP.⁷⁷

Unresponsive TTP: Management Algorithm Proposal

In our institution, first line therapy for suspected TTP includes daily TPE, corticosteroids, caplacizumab (despite doubts about its cost-effectiveness) and early rituximab administration. With this schedule, we observe a response rate over 90% and a mortality rate less than 5%.¹⁵ Considering the highly variable clinical presentations and first-line therapeutic management protocols of acquired TTP patients, it is difficult to recommend a universal management algorithm for refractory forms of the disease. In most of the literature on refractory TTP, patients receive multiple second-line therapies, sequentially at best, making it difficult to draw any conclusions about which drug the patients ultimately responded to.

Patients with a suspicion of TTP undergo a diagnostic workup, including ADAMTS13 study, before the onset of first-line emergency treatment. Frontline therapy includes TPE (60 mL/kg/day with fresh frozen plasma), corticosteroids (1 mg/kg/day prednisone equivalent, administered after TPE sessions), early rituximab (375 mg/m² administered after TPE sessions at day 1 and 4) and caplacizumab (10 mg/kg IV before first TPE then 10 mg/day SC after TPE sessions). In cases where patients respond to first-line treatment, TPE are stopped when platelet rate is measured above 150 G/L for 2 consecutive days. Corticosteroids are maintained at the same dose for 3 weeks and then tapered and stopped on 1 to 2 weeks. If remission of TTP occurs before day 8, B-cells depletion is monitored in the blood and patients may skip scheduled rituximab infusions on days 8 and 15 if CD-20 cells are not detectable anymore. Caplacizumab is administered for 4 weeks after the last TPE. Shorter or longer durations of caplacizumab are discussed on a case by case basis, depending on the results

of ADAMTS13 activity dosages performed weekly during that period.

In case of a refractory course of TTP, patients are thoroughly evaluated for confounding factors. If refractory TTP cannot be ruled out, second-line therapy is promptly started. According to our local protocols of care, the choice of therapeutic intensification depends on the kind of symptoms revealing refractory TTP.

In patients with persisting biological anomalies only (thrombocytopenia, hemolysis markers, etc), our local practice is to add vincristine infusions (1.4 mg/m², one single dose) and/or high dose steroid pulses (methylprednisolone 1000 mg/day for 3 days, administered after TPE sessions) to first-line therapy. In our experience, this schedule is sufficient to lead most patients to remission.

Patients with minor clinical events ascribable to TTP (headaches, brisk reflexes, angina, etc) while on first-line therapy receive vincristine and/or high dose steroid pulses and twice-daily TPE for 1 to 3 days depending on symptoms resolution.

Patients with major clinical events (seizures, coma, unstable angina, etc) receive vincristine, high dose steroid pulses, twice-daily TPE and additional immunosuppressive drugs (cyclophosphamide, bortezomib, etc).

In patients not responding to all previous therapies, splenectomy is used as salvage therapy.

Figure 3 illustrates the management of refractory TTP at our institution.

As TTP is a rare disease and a refractory course even rarer, management of these cases still relies mostly on expert clinicians' experience. Patients with refractory TTP should at least be evaluated by a physician used to TTP management, and in some cases be transferred to an expert team.

Future Perspective: Human Recombinant ADAMTS13

Human recombinant ADAMTS13 (rhADAMTS13) is not yet available in routine practice.

A Phase 1 study on rhADAMTS13 (BAX930) in patients with congenital TTP demonstrated safety and a pharmacokinetic profile comparable to fresh frozen plasma.⁷⁸ A Phase 3 study on rhADAMTS13 use as prophylaxis or on demand treatment in patients with congenital TTP is ongoing (NCT03393975).

There is limited data suggesting the efficacy of rhADAMTS13 in acquired TTP, the mechanism of action being an overriding of anti-ADAMTS13 autoantibodies (ex-vivo and animal model).⁷⁹

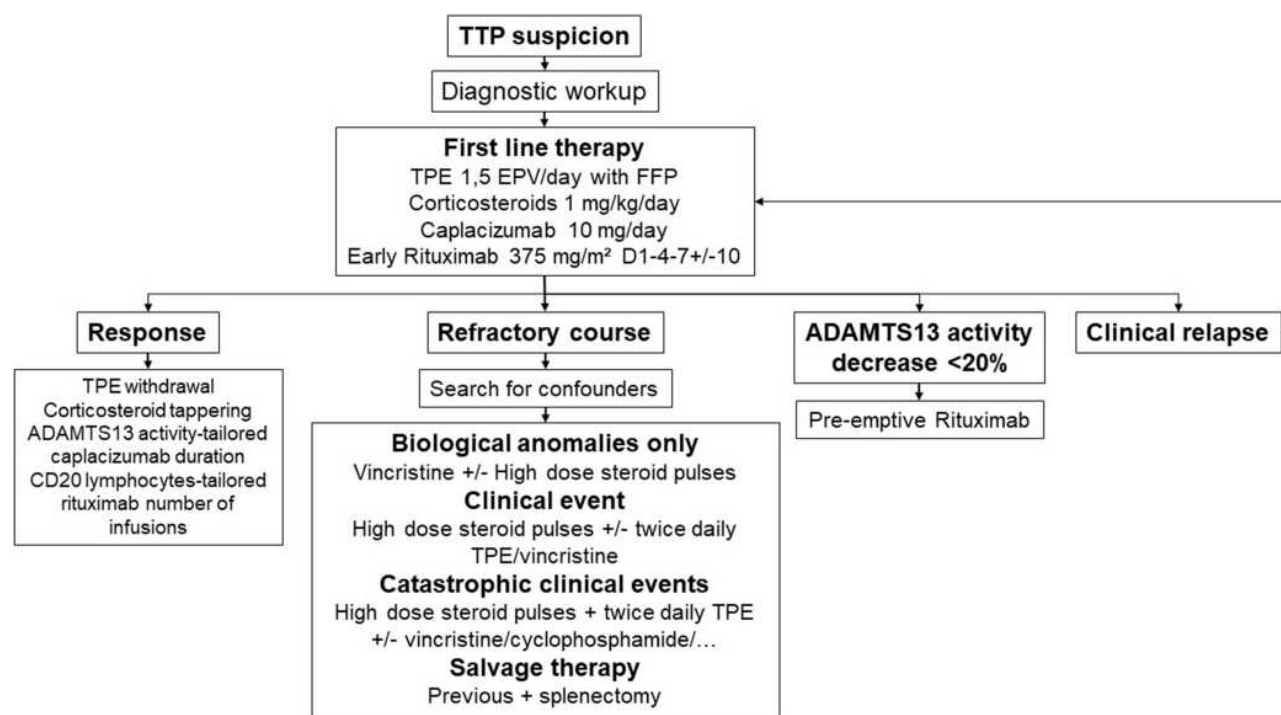


Figure 3 Unresponsive TTP management algorithm.

A Phase 2 study aiming at determining the potential role of rhADAMTS13 (SHP655) in addition to the standard of care in patients with acquired TTP is also ongoing (NCT03922308).

Conclusion

With modern standards of care, including TPE, corticosteroids, rituximab and caplacizumab, the mortality of acquired TTP has dropped from 10–20% to less than 5% and refractory/relapsing course from over 40% to less than 5%. As TTP is a rare disease, there will probably never be high quality data concerning the optimal management of these difficult cases. The algorithm for therapeutic escalation in unresponsive forms will ultimately depend on expert opinions and each center experience/preference. New therapeutic options, such as recombinant human ADAMTS13 (NCT03922308) and anfibatide (NCT04021173), are under evaluation and may hopefully lead to the disappearance of unresponsive TTP.

Disclosure

Dr Virginie Lemiale reports they belong to a research group which received fees from gilead, MSD, Alexion, celgene, and baxter, and that biomerieux pay travel for congress, outside the submitted work. Dr Sandrine Valade reports personal fees from Sanofi and Gilead/Kite, and non-financial support from Pfizer, outside the submitted work. Dr Eric Mariotte reports personal fees from Sanofi, outside the submitted work. The authors reported no other potential conflicts of interest for this work.

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