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Microangiopathic hemolytic anemia in a female patient with systemic lupus erythematosus

A 34-YEAR-OLD FEMALE PATIENT with a history of systemic lupus erythematosus (SLE) presented to the emergency department after several days of worsening swelling of the lips and legs along with decreased oral intake. She also reported intermittent pleuritic chest pain and exertional dyspnea for the past 2 months.

The patient was previously diagnosed with SLE with cutaneous manifestations including malar rash and discoid lesions of the face and scalp, recurrent pericardial and pleural effusions, diffuse arthralgias, and positive serologic markers including positive antinuclear antibody, anti-dsDNA antibody, anti-Smith antibody, and low complement levels. A previous pericardial pleural effusion resulted in normalization of complement levels with corticosteroid therapy.

Medical history was significant for hypertension, controlled with amlodipine, as well as two unexplained consecutive first-trimester miscarriages and recurrent pulmonary embolism. However, prior investigations were not consistent with antiphospholipid antibody syndrome, with the absence of lupus anticoagulant, anticardiolipin immunoglobulin (Ig)G/IgM, and anti-B2 glycoprotein IgG/IgM on two separate occasions 6 months apart. Her home medications included the anticoagulant rivaroxaban, amlodipine, prednisone, hydroxychloroquine, methotrexate, and folic acid. However, the patient had been nonadherent for the previous 4 months owing to lack of insurance coverage and instead had been trying alternative remedies including herbal tea.

doi:10.3949/ccjm.89a.21066

The patient also had lymphocytopenia, which can be associated with viral infections, active disease in SLE, and immunosuppressive therapy including glucocorticoids.

EMERGENCY DEPARTMENT EVALUATION

In the emergency department, the patient's temperature was 36.7°C (98.1°F), heart rate was 92 beats per minute, blood pressure was 156/103 mm Hg, and peripheral oxygen saturation was 100% on room air.

Physical examination revealed lip-swelling, aphthous ulcers, pitting edema in both legs and feet, and a new nontender rash on palms and soles. There was no appreciable murmur, pericardial rub, or jugular venous distention. The patient was alert and oriented to time, place, and person, followed all commands, and had no focal neurologic deficits. Funduscopic examination revealed vessel attenuation, cotton wool spots, and dot-and-blot hemorrhages without any papilledema, consistent with grade III hypertensive retinopathy.

Laboratory testing revealed anemia, thrombocytopenia, and an elevated serum creatinine level of 1.82 mg/dL (increased from baseline of 0.8 mg/dL).

Urinalysis with microscopic examination yielded cloudy appearance with high hemoglobin, red blood cells, protein, granular casts, urine protein-to-creatinine ratio, and urine sediment with dysmorphic red blood cells and red blood cell casts. See **Table 1** for laboratory results.

A chest radiograph showed significant increase of the cardiac silhouette compared with prior radiographs, and subsequent echo-

Chest radiography showed significant increase of the cardiac silhouette compared with previous radiographs

TABLE 1

Irregular patient laboratory results

Test	Patient value	Reference range
Hemoglobin, g/dL	11.8	12.3–15.3
Hematocrit, %	36.0	36.0–46.0
White blood cell count, $\times 10^9/L$	2.62	3.70–11.0
Absolute lymphocyte count, $\times 10^9/L$	0.55	1.00–4.00
Platelet count, $\times 10^9/L$	30	150–400
Serum creatinine, mg/dL	1.82	0.58–0.96
Alanine aminotransferase, U/L	65	7–38
Aspartate aminotransferase, U/L	148	13–35
C-reactive protein, mg/dL	7.3	< 0.9
Lactate dehydrogenase, U/L	1317	135–214
Reticulocyte count, %	8.7	0.4–2.0
Absolute reticulocyte count, $\times 10^9/L$	348	18–100
Haptoglobin, mg/dL	< 10	31–238
Fibrinogen, mg/dL	411	200–400
D-dimer, ng/mL	2900	< 500
Protein-to-creatinine ratio	1.7	< 0.2
Urine		
Clarity	Cloudy	Clear
Hemoglobin, mg/dL	3+	Negative
Protein, mg/dL	≥ 300	Negative
White blood cell count, cells/high-power field	6–10	0–5
Red blood cell count, cells/high-power field	> 25	0–3
Protein-to-creatinine ratio	1.7	< .2
C3, g/dL	29	86–166
C4, g/dL	10	13–64
dsDNA, IU/mL	> 1000	< 30
ADAMTS13, %	54	> 60

ADAMTS13 = a disintegrin and metalloproteinase with a thrombospondin type 1 motif, member 13

cardiography demonstrated a large pericardial effusion.

Possible diagnosis

Patient presentation of new aphthous ulcers, deteriorating kidney function, pancytopenia

with lymphocytopenia, and a new large pericardial effusion was suggestive of an active lupus flare. The lip-swelling was also concerning for angioedema, which has been reported as an unusual manifestation of SLE, with the

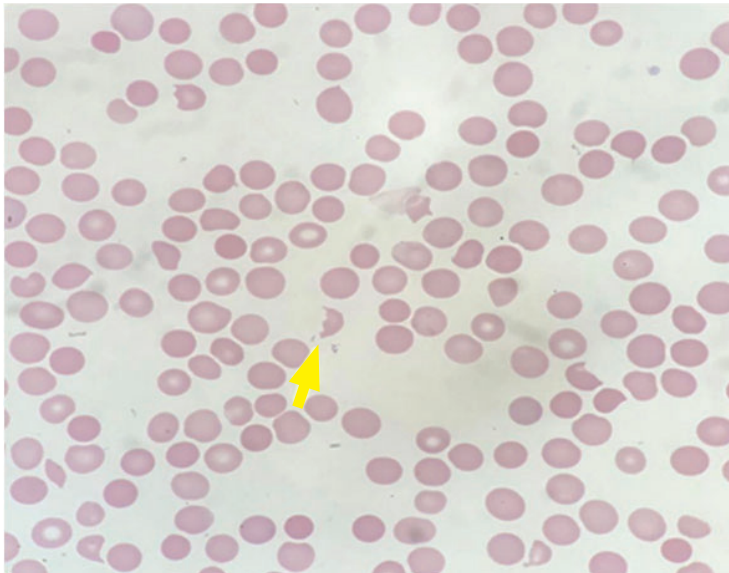


Figure 1. Peripheral smear. The yellow arrow indicates a schistocyte (hematoxylin and eosin stain, magnification $\times 400$).

The diagnosis of TMA can be inferred in patients with MAHA and thrombocytopenia in the appropriate clinical setting

proposed mechanism being acquired deficiency of C1 esterase inhibitor.^{1,2}

Peripheral smear revealed abundant schistocytes and thrombocytopenia (**Figure 1**). Other testing showed a reticulocyte index of 0.69% (indicative of hypoproliferation), a negative direct Coombs test, and prothrombin and partial thromboplastin times within normal limits.

At this point, microangiopathic hemolytic anemia (MAHA) was suspected, which is a descriptive term for nonimmune or Coombs-negative hemolysis resulting from intravascular red blood cell fragmentation that produces schistocytes on peripheral blood smear.

Further laboratory test results included low C3, low C4, high dsDNA, positive *Cri-thidia lucillae*, negative lupus anticoagulant, negative anticardiolipin IgG and IgM, negative anti-B2 glycoprotein IgG and IgM, low ADAMTS13 (a disintegrin and metalloproteinase with a thrombospondin type 1 motif, member 13) activity, and negative ADAMTS13 inhibitor.

■ IMPLICATIONS OF HEMOLYTIC ANEMIA WITH THROMBOCYTOPENIA

1 What is the implication of hemolytic anemia with thrombocytopenia in this patient?

- ☐ It may indicate possible development of thrombotic thrombocytopenic purpura (TTP)
- ☐ It is related to poorly controlled severe hypertension
- ☐ It is attributed to acquired complement-mediated thrombotic microangiopathy (TMA) associated with lupus nephritis
- ☐ It is attributed to Shiga toxin-mediated hemolytic uremic syndrome

The combination of characteristic laboratory data in our patient, including negative Coombs test, elevated lactate dehydrogenase (LDH), low haptoglobin, normal coagulation studies, and peripheral smear showing abundant schistocytes, were characteristic for MAHA. The combination with thrombocytopenia and worsening kidney function raised high suspicion for TMA.

TMA is a pathologic diagnosis determined by tissue biopsy in which abnormalities of arteriolar and capillary walls subsequently lead to microvascular thrombosis. As described previously, MAHA refers to Coombs-negative hemolysis resulting from intravascular red blood cell destruction, which may be caused by abnormalities in the microvasculature or even by intravascular devices such as prosthetic heart valves. Consequently, even though not all MAHA is caused by TMA, the diagnosis of TMA can be typically inferred in patients with MAHA and thrombocytopenia in the appropriate clinical setting.

Causes of TMA

TMA syndrome encompasses many disorders that include TTP, Shiga toxin-induced hemolytic uremic syndrome, drug-induced TMA, radiation-induced TMA, and complement-mediated TMA, which is also referred to as atypical hemolytic uremic syndrome.

Systemic disorders that can cause TMA include scleroderma, severe hypertension, preeclampsia/HELLP (hemolysis, elevated liver enzymes, and low platelets) syndrome, and antiphospholipid syndrome, especially the most severe form of antiphospholipid syndrome known as catastrophic antiphospholipid syndrome, which is characterized by disseminated intravascular thrombosis resulting in multiorgan failure within a short period of time (usually within 1 week).

Other causes. TMA is also well described in transplant recipients, both as a manifestation of transplant rejection and as possible toxicity from immunosuppressive medications such as cyclosporine and tacrolimus. Rarely, this syndrome can be caused by disorders involving regulators of the coagulation cascade (mutations in the gene encoding diacylglycerol kinase epsilon and thrombomodulin)³ and the molecules responsible for vitamin B₁₂ metabolism (mutations in the gene encoding methylmalonic aciduria and homocystinuria type C).⁴

Evaluation for the cause of TMA

Evaluation to determine the cause of TMA should initially focus on excluding the aforementioned systemic disorders associated with MAHA and thrombocytopenia. This should be followed by an investigation to distinguish among primary TMA syndromes, including the following:

- Assessment of exposure to drugs reported to cause TMA: chemotherapeutic agents (eg, gemcitabine, mitomycin), immunosuppressive agents (eg, cyclosporine, tacrolimus), vascular endothelial growth factor inhibitors (eg, bevacizumab), and illegal agents (eg, cocaine)
- Measurement of ADAMTS13 activity and ADAMTS13 inhibitor levels; elevations infer TTP
- Stool testing of Shiga toxin by immunoassay or polymerase chain reaction test to look for Shiga toxin-induced hemolytic uremic syndrome, especially in patients with diarrhea or known infectious diarrhea exposure
- Measurement of serum homocysteine and methylmalonic acid for cobalamin C deficiency-mediated TMA, which is caused by recessive (homozygous or compound heterozygous) mutations in the metabolism of the cobalamin-associated C (MMACHC) gene⁴
- Complement testing for complement-mediated TMA, specifically, for antibodies against complement factor (CF) H protein (anti-CFH antibody) and for complement gene mutations, including evaluation of complement regulatory genes (CFH, CFI, or membrane cofactor protein [CD46])

and complement effector genes (CFB or C3).

Kidney biopsy is not helpful for determining cause of primary TMA syndromes and may not distinguish primary TMA syndromes from other disorders such as systemic sclerosis or severe hypertension, which can also manifest the typical pathologic features of TMA.

What was the cause of this patient's TMA?

TTP is a primary TMA syndrome caused by severely reduced activity of the von Willebrand factor-cleaving protease ADAMTS13.⁵ When ADAMTS13 protease activity is reduced, large von Willebrand factor multimers can accumulate on the endothelial surface of small arterioles and capillaries, leading to potentially fatal platelet accumulation and subsequent mechanical red blood cell fragmentation.⁵ TTP can be acquired, from an autoantibody inhibitor to ADAMTS13, or it can be hereditary, owing to inherited mutations in ADAMTS13. Genetic testing of the ADAMTS13 gene is the definitive means of documenting the diagnosis of hereditary TTP in an individual with severe ADAMTS13 deficiency (activity < 10% of normal) without an inhibitor. On the other hand, acquired TTP can be confirmed by severe ADAMTS13 deficiency and the presence of an ADAMTS13 autoantibody inhibitor in the appropriate clinical setting (eg, a patient with MAHA and thrombocytopenia that responds to plasmapheresis).⁶

Hematologic recovery in TTP can be expected with plasmapheresis, as indicated by improvement in LDH (often within 1 day) and platelet count (within 2 to 3 days).^{7,8} TTP was ruled out in our patient on the basis of relatively normal ADAMTS13 activity, a negative ADAMTS 13 inhibitor, poor hematologic response to emergency plasmapheresis, and lack of hematologic recovery to suggest resolution of MAHA after 3 days of plasmapheresis.

Severe hypertension can also cause MAHA and thrombocytopenia with characteristic pathologic features of TMA in the kidneys.⁹ Grade III/IV retinopathy is more likely to be present, and recovery of kidney function and resolution of MAHA is typical with proper treatment of severe hypertension.^{9,10} Furthermore, severe hypertension may also occur in

TTP can be acquired, from an autoantibody inhibitor to ADAMTS13, or it can be hereditary, owing to inherited mutations in ADAMTS13

primary TMA with severe renal involvement, so the temporal relationship is important.⁹ Although the patient did have funduscopic findings consistent with grade III hypertensive retinopathy, kidney function and MAHA did not improve despite appropriate management of blood pressure, prompting us to continue our investigation into other causes of TMA.

Shiga toxin-producing pathogens (eg, *Escherichia coli* O157:H7) can induce hemolytic uremic syndrome and are often related to exposure to improperly prepared foods, inadequately cooked meats, or farm animals.¹¹ However, our patient did not report any severe abdominal pain or diarrhea with bloody stool, and DNA testing performed on the patient's stool sample for Shiga-toxin genes was negative. Additional workup investigating non-Shiga-toxin-producing infectious organisms associated with MAHA and thrombocytopenia was also unrevealing, including tests for bacterial (eg, *Clostridioides difficile*, *Legionella*), viral (eg, cytomegalovirus, hepatitis viruses, human immunodeficiency virus, varicella-zoster virus, Epstein-Barr virus), and fungal (eg, *Aspergillus*) infections.

Complement-mediated TMA or atypical hemolytic uremic syndrome is a chronic, systemic, life-threatening disease characterized by TMA and severe multiple end-organ damage caused by dysregulation and uncontrolled activation of the alternative complement pathway. Complement regulatory proteins, including CFH, CFI, and CD46, prevent inappropriate activation of the complement cascade.¹² The diagnostic workup for determining cause of complement-mediated TMA includes testing for inhibitory antibodies such as anti-CFH antibody, as well as genetic testing, including evaluation of the CFH, CFI, CD46, CFB, and C3 genes.¹³ On the other hand, testing for serum complement biomarkers such as CH50, C3, and C4 have no diagnostic role in complement-mediated TMA owing to low sensitivity and specificity. For instance, a low C3 level may only be appreciated in certain mutations such as variants in CFH, CFB, and C3.¹⁴

Complement activation is also common in lupus nephritis, and a subset of these patients (17.5%) can present with complement-mediated TMA,¹⁵ described as a separate clinical

entity that can be resistant to treatment with high-dose corticosteroids, immunotherapy, and plasmapheresis.¹⁶ The pathogenesis of renal TMA in lupus nephritis is yet to be determined and may be multifactorial, potentially attributable to antiphospholipid syndrome, malignant hypertension, or scleroderma.¹⁷

Our patient had normal plasma levels of complement factors CFH and CFB, along with negative anti-CFH antibody. Complement genetic studies were not obtained, and the patient did not report a family history of TMA disorders suggestive of a hereditary cause. However, one should be mindful that most complement mutations associated with TMA are heterozygous, and many family members with heterozygous mutations are asymptomatic.¹⁸ If performed in this patient, complement genetic studies might have provided a more specific diagnosis.

Although complement testing was unrevealing in this patient, a presumptive clinical diagnosis of complement-mediated TMA was made in light of the progressive kidney dysfunction, MAHA, thrombocytopenia, and absence of an alternate explanation for these findings (eg, no known drug exposure associated with drug-induced TMA, no antiphospholipid syndrome or scleroderma, no response after adequate control of hypertension, no ADAMTS13 deficiency, no Shiga toxin in stool).

■ CASE CONTINUED: CONDITION WORSENE

The differential diagnosis for the patient's acute kidney failure included lupus nephritis and TMA.

Plasmapheresis was performed for a total of 3 days, with no clinical improvement. Then, administration of eculizumab, a complement inhibitor, was commenced in light of the suspected diagnosis of complement-mediated TMA.

C5 inhibitors, including eculizumab and ravulizumab, prevent the cleavage of cell surface-bound complement protein C5 and inhibit generation of proinflammatory terminal complement proteins C5a and the membrane attack complex C5b-9, consequently reducing endothelial cell activation and thrombosis but also increasing the risk of developing mening-

Plasma exchange was started without positive results; eculizumab was administered for suspected complement-mediated thrombotic microangiopathy

gococcal infections.¹⁹ Therefore, in addition to meningococcal vaccination, the patient received antimicrobial prophylaxis with amoxicillin for the duration of eculizumab administration.

Complications

Pericardial effusion. During hospitalization, the patient required transfer to the cardiac intensive care unit because of pericardial effusion with tamponade. Pericardiocentesis was performed, and subsequent pericardial fluid studies were suggestive of an exudative effusion.²⁰ There was a predominance of mononuclear cells as well as negative Gram stain, bacterial/fungal cultures, and cytology, favoring active SLE flare as the cause of the exudative pericardial effusion.

Mental status changes. Moreover, the patient's clinical course was complicated by the development of altered mental status. She became less interactive, could not follow commands, and neurologic examination revealed slightly brisker reflexes in the right upper extremity. This was concerning for possible primary manifestation of neuropsychiatric SLE, as there was past evidence of systemic involvement of SLE.

Other potential etiologies that were considered included infectious encephalitis, given the patient's immunocompromised state, stroke secondary to central nervous system involvement by SLE, and posterior reversible encephalopathy syndrome, given her recent blood pressure fluctuations with systolic blood pressure higher than 200 mm Hg.

Magnetic resonance imaging of the brain was concerning for antibody-mediated striatal encephalitis. This finding is not specific for SLE, and viral encephalitis should also be considered in the differential diagnosis.

A lumbar puncture that was subsequently performed did not show any pleocytosis in the cerebrospinal fluid, rendering an intrathecal infection less likely as the etiology of the patient's acute encephalopathy and favoring the diagnosis of neuropsychiatric lupus.

Acute kidney injury. The patient's kidney function continued to deteriorate, requiring kidney replacement therapy within 5 days of initial presentation to the hospital. Kidney biopsy was considered at this time but was ul-

timately not performed due to severe thrombocytopenia.

■ WOULD KIDNEY BIOPSY HAVE HELPED?

2 What would have been the value of kidney biopsy in this patient?

- ☐ It would have been helpful in differentiating a specific type of TMA
- ☐ It would have been unlikely to affect patient management, as the therapeutic window was missed early in the disease course because of bleeding concerns related to thrombocytopenia
- ☐ It would have ascertained the extent of ischemic injury sustained in relationship to renal TMA to facilitate decision-making regarding next steps in therapy
- ☐ It was contraindicated in this patient owing to increased risk of bleeding complications from azotemia and thrombocytopenia

Risk vs potential benefit of biopsy

Biopsy of the kidney may be performed for suspected TMA syndrome.²¹ However, kidney biopsy may not inform treatment decisions for a specific TMA as there is no consensus regarding features on kidney biopsy that definitively diagnose a specific TMA syndrome. Kidney biopsy is not usually done to evaluate patients with acute kidney injury, MAHA, and thrombocytopenia unless there is a specific management decision that would be influenced by the results.

Our patient's clinical condition was an example of a situation in which a kidney biopsy may have been helpful: it may have identified the extent of ischemic injury sustained in relationship to potential TMA, as well as the degree of irreversible kidney injury to support decision-making for next therapeutic treatment steps. Kidney biopsy may have justified the maintenance or discontinuation of immunosuppressive therapy for lupus nephritis, especially as the patient had not yet developed other systemic manifestations of SLE (eg, pericardial effusion, neuropsychiatric SLE) at the time kidney biopsy was being considered. The presence of extensive interstitial fibrosis or glomerulosclerosis (class VI advanced sclerosing lupus nephritis), as opposed to focal or

The patient was transferred to intensive care to manage pleural effusion and tamponade

diffuse proliferative glomerulonephritis (class III and IV lupus nephritis, respectively), might have prompted discontinuation of immunosuppression.

Although the association between kidney dysfunction and bleeding is well recognized, major complications have been noted to be less than 1%.²² Our patient did not undergo kidney biopsy owing to persistent thrombocytopenia, as thrombocytopenia increases the risk of major complications with kidney biopsy in patients with SLE.²³

SUBSEQUENT CLINICAL COURSE

Intravenous methylprednisolone and mycophenolate mofetil were initially employed as induction therapy for lupus nephritis. Mycophenolate mofetil was preferred over cyclophosphamide because mycophenolate mofetil has been shown to be more effective than intravenous cyclophosphamide in inducing remission of lupus nephritis.²⁴ The patient's race was a consideration as it has been demonstrated that African American patients with lupus nephritis respond better to mycophenolate mofetil than to intravenous cyclophosphamide (53.9 vs 40%; $P = 0.39$).²⁵

However, concern for neuropsychiatric SLE with antibody-mediated striatal encephalitis later prompted the discontinuation of mycophenolate mofetil in favor of cyclophosphamide, as cyclophosphamide is the only therapy that has demonstrated efficacy for neuropsychiatric SLE compared with methylprednisolone in a controlled clinical trial.²⁶

Although high-dose glucocorticoids and intravenous cyclophosphamide remain the cornerstone for patients with neuropsychiatric SLE, rituximab or intravenous immunoglobulin may be used if response is not achieved.²⁷ Furthermore, even without studies regarding efficacy of plasmapheresis in patients with lupus antibody-mediated striatal encephalitis, there is some evidence regarding its use in severe neuropsychiatric SLE.^{28,29} Consequently, plasmapheresis was briefly performed together with administration of cyclophosphamide, with subsequent improvement in the patient's mental status.

Although the hemolytic anemia and thrombocytopenia resolved after 6 weeks of

therapy with eculizumab, the patient's kidney function did not recover for more than 2 months during hospitalization. Eventually the decision was made to stop eculizumab and continue immunosuppressive therapy with cyclophosphamide upon discharge to a rehabilitation facility. Although the patient did not undergo kidney biopsy to guide therapy, the rationale for continued treatment with immunosuppressive therapy at discharge was owing to evidence of other systemic manifestations of SLE during the course of hospitalization, including pericardial effusion and lupus antibody-mediated striatal encephalitis.

WHICH MANAGEMENT STRATEGY?

3 Which management strategy has been shown to improve kidney outcomes in patients with complement-mediated TMA associated with SLE?

- ☐ Therapeutic plasma exchange
- ☐ Immunosuppressive therapy specific for lupus nephritis
- ☐ Complement-inhibiting therapy with eculizumab
- ☐ Supportive management with red blood cell and platelet transfusions

Therapeutic plasma exchange

The immediate management decision for a patient with suspected TMA is whether to perform plasma exchange for a presumptive diagnosis of TTP or anticomplement therapy for a presumptive diagnosis of complement-mediated TMA.

Plasma exchange should be initiated in adults as soon as diagnosis of TMA is suspected. This helps remove ADAMTS13 autoantibodies, factor H autoantibodies, and hyperfunctioning complement components as well as replaces ADAMTS13 and faulty complement regulators.²¹

Immunosuppressive therapy specific for lupus nephritis

Unfortunately, TMA associated with lupus nephritis is usually refractory to treatment with corticosteroids, cyclophosphamide, immunomodulation, and plasma exchange.^{30,31} However, a 2018 series involving patients with complement-mediated TMA related to

The patient was discharged to a rehabilitation facility on immunosuppressive therapy with cyclophosphamide

SLE reported complete recovery of hematologic parameters in all patients and recovery of kidney function in 80% of patients treated with eculizumab.¹⁶

Complement inhibition with eculizumab or ravulizumab

Initial evidence for the efficacy of eculizumab for treating TMA came from a prospective, open-label case-series³² of patients with complement-mediated TMA in which eculizumab improved kidney and hematologic outcomes and reduced thrombotic microangiopathic events including requirement for plasma exchange or infusion, reduction in platelet counts, and initiation of hemodialysis.³²

Although C5 inhibitors including eculizumab and ravulizumab are currently the only effective therapy designed to prevent the underlying mechanism of microvascular damage, which is dysregulation of the complement activation system, there are limited data on long-term outcomes in patients with TMA treated with terminal complement inhibition. There are also several unanswered questions regarding C5 inhibitor therapy, including optimal duration of therapy, appropriate strategies for monitoring therapeutic efficacy in patients with suboptimal response or during periods of infection or inflammation, and determining tailored therapeutic regimens in various groups of patients with TMA, including those with complement-mediated TMA.

At Cleveland Clinic, eculizumab therapy has been used broadly, not only for clinically suspected complement-mediated TMA, but also for other TMA syndromes. Owing to the lack of available information regarding man-

agement decisions for patients with TMA, investigation into management strategies for these patients is warranted as is research regarding kidney and hematologic outcomes after therapy.

TAKE-HOME POINTS

- Suspect complement-mediated TMA in patients with MAHA in whom other causes including TTP, antiphospholipid antibody syndrome, Shiga toxin-induced hemolytic-uremic syndrome, and uncontrolled hypertension have been excluded.
- Kidney biopsy does not help identify a specific type of TMA and therefore does not inform treatment decisions for specific TMAs. However, kidney biopsy may assist in evaluating the extent of ischemic injury sustained in relation to TMA and thus may characterize the degree of irreversible kidney injury and facilitate decision-making regarding next therapeutic steps, especially in cases that are refractory to C5 inhibitor therapy.
- C5 inhibitors, including eculizumab and ravulizumab, currently represent the only effective therapy designed to prevent the underlying mechanism of microvascular damage in complement-mediated TMA, which is dysregulation of the complement activation system. However, data are limited on long-term outcomes in patients with TMA treated with terminal complement inhibition.

DISCLOSURES

The authors report no relevant financial relationships which, in the context of their contributions, could be perceived as a potential conflict of interest.

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