ACQUIRED HEMOPHILIA A: A CASE REPORT
Gunjan Shah, MD, Ajay Kandra, MD, Douglass A. Drelich, MD

Case Report
A 61-year-old African American male, with a past medical history of asthma and benign prostatic hypertrophy, presented from an outside hospital with complaints of hematuria and hematemesis. He had initially noted increased bruising two months prior to admission. He also had symptoms suggestive of a non-traumatic thigh hematoma the month prior. He developed hematuria one month prior to admission and underwent an outpatient cystoscopy one week prior to admission which was non-diagnostic. Cystoscopy was repeated on the day of admission localizing the hemorrhage to his right kidney with a biopsy specimen suggesting a low-grade papillary neoplasm. The patient subsequently developed hematemesis. Emergent esophagogastroduodenoscopy (EGD) at the outside hospital revealed a Mallory-Weiss tear. Epinephrine was injected with initial hemostasis. EGD was repeated after transfer due to recurrent hematemesis and clips were placed to control the hemorrhage.

The patient experienced continued hematuria requiring repeat cystoscopy with laser ablation, as well as biopsy of the right ureter and right mid-pole infundibulum. However, multiple repeated biopsies failed to confirm the presence of malignancy.

On admission, he was afebrile, with a heart rate of 94 bpm, blood pressure of 142/88 mm Hg, and oxygen saturation of 100% on room air. His physical exam was within normal limits. His initial hemoglobin was 7.6 g/dL, white blood cell count of 9,700 cells per cubic millimeter (cmm), and platelet count 138,000 cells per cubic millimeter. Coagulation studies revealed an elevated activated partial thromboplastin time (aPTT) of 76.5 seconds. An aPTT mixing study was performed with the aPTT correcting to 34.9 seconds (normal 23.6-35 seconds). Incubation was not performed. Additional laboratory evaluation demonstrated factor VIII <0.01 U/ml (normal 0.52-1.43 U/ml) and a factor VIII antibody was detected with a measured titer of 33.3 Bethesda Units (BU). Von Willebrand antigen and activity levels were in the normal ranges.

Given the undetectable Factor VIII level, lack of prior history of a bleeding diathesis, and the presence of an inhibitor with a high titer level, a diagnosis of Acquired Hemophilia A (AHA) was made.

The initial management strategy was to control the acute bleeding and eradicate the inhibitor. Hemostasis was initially obtained with recombinant Factor VIIa (rFVIIa) at 90 mcg/kg every 3-4 hours episodically for major bleeding for 6-12 hours and prior to planned invasive procedures. However due to a decrease in clinical response during the admission, it was switched to activated Prothrombin Complex Concentrate (aPCC), specifically factor eight inhibitor bypass activity (FEIBA). Immunosuppression was initiated with 60 mg of prednisone daily. He continued to experience severe hematuria requiring continuous bladder irrigation and multiple cystoscopies for clot evacuation and laser fulguration. Five days after initiation of prednisone, the patient failed to show significant clinical improvement, which warranted replacement with another type of immunosuppressive therapy. Pulsed high-dose dexamethasone 40 mg (Dex)45-49 and Rituximab35-40 were administered for a period of four days. However, the patient had a grade 4 reaction to the Rituximab consistent with anaphylaxis requiring cessation of the treatment. He then received intravenous immunoglobulin (IVIG) for two days52-57 without any clinical response.

Given his inadequate response to standard and high dose steroids and IVIG, and a factor VIII level of 2% with an inhibitor titer of 40 BU, oral cyclosporine was initiated on hospital day 22 and titrated to maintain a therapeutic goal of 200-400 ng/ml.41-44 His hematuria gradually improved, and he required no further transfusions.

Over the course of his 36-day hospitalization he received a total of 22 units of pRBC, 9 units of FFP, 34 doses of rFVIIa, and 6 doses of FEIBA. He was discharged on a therapeutic dose of oral cyclosporine with a Factor VIII level of 4% and an inhibitor titer of 57.1 BU at the time of discharge (Graph 1).

He was evaluated weekly following discharge during which he did not have any major hemorrhage. His cyclosporine dose was titrated based on trough levels. He continued pulsed, high-dose dexamethasone for 4 consecutive days every 28 days. His Factor VIII level demonstrated gradual improvement during this period and inhibitor levels gradually decreased. Following the fourth cycle of high-dose dexamethasone, his Factor VIII level rose to 29% and his inhibitor level became undetectable. Dexamethasone was discontinued following the fourth cycle after obtaining an undetectable inhibitor level. Cyclosporine weaning was initiated following two consecutive factor levels greater than 80%.

Discussion
Acquired Hemophilia A (AHA) is a rare condition in which autoantibodies, usually of the IgG class, are produced against Factor VIII. This results in low plasma Factor VIII levels. Its incidence ranges from 0.2-4 cases/million/year according to various reports.1,2,3,4,5 However, given the complexity of diagnosis, the condition may be under diagnosed.11,12,13 The incidence increases with age with a peak incidence in the 7th and 8th decade of life. There is a small peak in the 3rd decade likely corresponding to postpartum inhibitors.5 The mortality rate is high, ranging from 8%-22%. This is related to severe hemorrhage that can occur in 85%-90% of the patients.3,31 Approximately half of the cases are associated with underlying conditions including pregnancy, autoimmune disorders, malignancy, medications, dermatologic conditions, inflammatory bowel disease, and...
infections such as hepatitis B and C. The other half occur without evidence of a coexisting disorder (idiopathic). This condition commonly presents with mucosal bleeding (including epistaxis, gastrointestinal, and genitourinary), ecchymosis, and soft tissue hemorrhage including retroperitoneal bleeding. In contrast to congenital Hemophilia A, hemarthrosis is rare.

The diagnosis is confirmed by detection of a prolonged aPTT which does not correct by 1:1 mixing with normal plasma (aPTT mixing study). Occasionally, there is immediate correction but after incubation for 2 hours at 37°C there is reversal of the correction. Factor VIII levels are markedly reduced, and anti-FVIII antibodies are detectable.

The primary goals of treatment for AHA are to control acute bleeding and to suppress the autoantibody. Antibody eradication often requires treatment of the instigating condition when present. AHA can represent a transfusion emergency requiring intensive transfusion support. In patients with high titer inhibitors (>5 BU), bypassing agents, such as FEIBA and rVIIa, are the mainstay of therapy for acute hemorrhage. The recommended dose of FEIBA, ranges between 50–100 U/kg administered every 6–12 hours, not to exceed a single dose of 100 U/kg or a daily dose of 200 U/kg. In one study, FEIBA controlled hemorrhage in 86% of patients. The recommended dose of rVIIa ranges from 90 to 120 mcg/kg every 2–3 hours. A pooled analysis of data of 139 patients from different sources reported an efficacy rate of 88%. Venous thromboembolic disease (VTE) is an adverse event associated with both of these agents.

Factor VIII in large doses should be considered in patients known to have low-titer inhibitors (<5 BU). Porcine Factor VIII from pooled plasma, while not currently available in the US, has been used historically to increase plasma factor VIII levels, and it takes advantage of the minimal antigenic cross reactivity with human Factor VIII inhibitors. A recombinant porcine factor VIII, B-domain deleted (OBI-1) has recently been tested in a clinical trial in the US. A number of immunosuppressive therapies have been used to control inhibitors. However, current treatment is based on small, uncontrolled, single-center cohorts and meta-analysis. Corticosteroids and cytotoxic drugs, used either alone or in combination, have been regarded as the mainstay of therapy. An alternate modality of inhibitor suppression is IVIG; however, this has generally been less effective. Removal of the inhibitor via an immunoadsorption process has been

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**Figure 1. Factor VIII Level and Inhibitor Titer by Treatment Day**

- **Factor VIII Level (U/ml)**
- **Inhibitor Titer (BU)**
- Treatment Day
  - Dex-1
  - IVIG
  - Dex 2
  - Dex 3
  - Dex 4

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attempted.\textsuperscript{2,8,20} But this process requires central venous access, specialized equipment and training, and close FVIII monitoring.

Oral steroids used alone at 1 mg/kg prednisone equivalents per day for 3–6 weeks can cause remission in one third to one half of patients with acquired hemophilia.\textsuperscript{3,16,30,31} Cyclophosphamide has been used as initial therapy and in refractory patients with high-titer inhibitors (>5 BU) leading to higher remission rates of up to two-thirds versus steroids alone.\textsuperscript{1,3,11,16,32,33} However, a 2-year national surveillance report of acquired hemophilia A by the UK Haemophila Centre Doctors’ Organization concluded that there was no difference in inhibitor eradication or mortality between patients treated with steroids alone or with a combination of steroids and cytotoxic agents.\textsuperscript{11}

Several reports have demonstrated the effectiveness of Rituximab in acquired hemophilia A. The general high level of tolerance of this drug has led to initial use of this treatment. However, there is currently no data to establish its superiority over other treatments. The dose typically given is 375 mg/m\textsuperscript{2} infused weekly for 4 weeks and administered concomitantly with other immunosuppressive drugs.\textsuperscript{34–39}

Cyclosporine (CsA) has also been used for inhibitor suppression.\textsuperscript{40,43} Use of CsA requires monitoring drug levels to ensure efficacy and prevent toxicity. However, its use is limited due to availability of other strategies such as cytotoxic drugs and Rituximab.

We describe a case of AHA with high titer antibody that experienced clinical deterioration on initial monotherapy with prednisone. The patient was unable to tolerate Rituximab due to anaphylaxis and was also refractory to IVIG.

We report a novel strategy of combining Cyclosporine with pulsed, high dose dexamethasone at a dose of 40mg daily for four consecutive days every 28 days. Pulsed dose dexamethasone is an acceptable modality of immunosuppression and has been used for the treatment of Idiopathic Thrombocytopenic Purpura (ITP), another immune-mediated hematological disorder.\textsuperscript{44–46} In general it is tolerated well, without producing the long term effects of steroid toxicity. No randomized trials exist comparing the efficacy and tolerability of pulse dose dexamethasone with longer duration steroids. However, we have experienced good results in our institution for the treatment of ITP, and this has become our preferred first line therapeutic modality. The patient tolerated this regimen extremely well and went into complete remission with undetectable antibody titers after the fourth cycle of pulse dexamethasone. He is currently in remission and is being weaned from cyclosporine.

References
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