To the Editor:
Hydralazine-induced antineutrophil cytoplasmic antibody vasculitis (HIAV) is a rare side effect of hydralazine treatment and can have notable morbidity and mortality.

Incidence and prevalence of HIAV is unclear due to its rarity, but risk factors that have been identified are older age, a cumulative dose of 100 g of hydralazine at the time of presentation, female sex, thyroid disease, HLA-DR4 genotypes, slow hepatic acetylation, and the null gene for C4.

Symptoms of HIAV can include fever, malaise, arthralgia, weight loss, or even involvement of organs such as the kidneys and lungs.

If recognized early, cessation of hydralazine and supportive therapy generally are sufficient; however, severe cases may need management with high-dose corticosteroids, rituximab, and even plasmapheresis.

A 67-year-old woman presented with progressive, tense, hemorrhagic, and necrotic bullae on both sides of the face and neck as well as the extremities of 2 weeks’ duration. She had a history of hypertension and a thyroid nodule after unilateral thyroid lobectomy. A review of symptoms was positive for worsening dyspnea and progressive generalized weakness. Noteworthy medications included amiodipine, metoprolol, levothyroxine, and oral hydralazine 75 mg 3 times daily for 13 months.

Bullae first appeared on the patient’s scalp and quickly progressed with a cephalocaudal pattern with a propensity for the eyes, nostrils, and labial mucosa (Figure 1). The tongue was covered by an eschar, and she had diffuse periorbital edema. Additionally, concentric purpuric patches were noted on the thighs and lower legs (Figure 2).
Pertinent laboratory findings included a positive antinuclear antibody titer of 1:320 and perinuclear anti-neutrophil cytoplasmic antibody (ANCA) titer of 1:160, along with an elevated serum creatinine level (2.31 mg/dL [reference range, 0.6–1.2 mg/dL]). Bilateral perihilar infiltrates with bilateral pleural effusions were noted on a chest radiograph.

While hospitalized, she developed pulmonary hemorrhages and a progressive decline in respiratory status. She subsequently was admitted to the medical intensive care unit. Aggressive support was administered, and several skin biopsy specimens were obtained along with an endobronchial biopsy of the right middle lobe.

Skin histopathology revealed a necrotic vasculitis (Figure 3). Direct immunofluorescence was not performed. Lung histopathology showed fragments of bronchial tissue with acute and chronic inflammation, focal necrosis, granulation tissue formation, edema, and squamous metaplasia. Together with the clinical history, these findings were consistent with HIAV.

Hydralazine was immediately discontinued, and the patient was started on 65 mg daily of intravenous methylprednisolone; methylprednisolone was later changed to oral prednisone 30 mg daily. Due to multiple organ involvement—lung and kidney—intravenous rituximab 375 mg/m² every week for 4 weeks, per lymphoma protocol, was started. Within 2 weeks of beginning therapy, her renal function and respiratory status improved, and by week 4, the skin lesions had completely resolved. Although initially she did well on immunosuppressive therapy with resolution of all symptoms, the patient contracted *Clostridium difficile*–induced systemic inflammatory response syndrome after 5 weeks of therapy and died.

Hydralazine was first introduced in 1951 for adjunctive hypertension therapy due to its vasodilation effects.1-3 Since its introduction, it has been implicated in 2 important disease processes: HIAV and hydralazine-induced lupus.

Hydralazine-induced ANCA vasculitis was first documented in 1980; by 2011, multiple cases had been reported.1-7 Hydralazine-induced ANCA vasculitis has occurred in patients aged 11 to 79 years taking 50 to 300 mg daily. Symptom onset varies from 6 months to 14 years, with a mean exposure duration of 4.7 years and mean daily dose of 142 mg.1-7

Clinical manifestations range from less specific, such as fever, malaise, arthralgia, myalgia, and weight loss, to single tissue or organ involvement that may be fatal. The most frequent clinical features include kidney involvement (81%), cutaneous vasculitis (25%), arthralgia (24%), and pleuropulmonary involvement (19%). Cutaneous manifestations include but are not limited to palpable lower extremity purpura; morbilliform eruptions; and hemorrhagic blisters on the lower legs, arms, trunk, nasal septum, and uvula.1-4,8
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The most commonly affected organ is the kidney, which commonly presents as hematuria, proteinuria, and elevated serum creatinine level. Histopathologically, patients most likely will have necrotizing and crescentic glomerulonephritis that is pauci-immune by immunofluorescence.2,3 The lungs are the next most commonly affected organ, with a classic presentation of cough, dyspnea, and hemoptysis in the setting of intra-alveolar hemorrhage.4,5 When both the kidneys and lungs are involved, the patient is said to have pulmonary-renal syndrome that is characterized by lung infiltrates or nodules with or without hemorrhage, hemoptysis, and pleuritis in the setting of glomerulonephritis.1,6

Clear data on incidence and prevalence of HIAV does not exist due to the rarity of the disease and the lack of prospective studies. To identify a clear incidence and prevalence, prospective longitudinal studies with larger cohorts along with better recognition and diagnosis are needed.2,5,6 A few predisposing risk factors have been identified, including older age, a cumulative dose of 100 g at the time of presentation, female sex, a history of thyroid disease, HLA-DR4 genotypes, slow hepatic acetylation, and the null gene for C4.1,3,5,9-11 Our patient was an older woman with a history of thyroid disease who had been taking oral hydralazine 75 mg 3 times daily for 13 months. During this 13-month duration, she had no dose adjustments.

Currently, the pathomechanism for HIAV is unclear and may be multifactorial. There are 4 main theories:2,5,6,10,12:

1. Hydralazine and its metabolites accumulate inside neutrophils, then subsequently bind and alter the configuration of myeloperoxidase (MPO). This alteration leads to spreading of the autoimmune response to other autoantigens, making neutrophil proteins (eg, elastase, lactoferrin, nuclear antigens) immunogenic.

2. Hydralazine binds MPO in neutrophils, creating cytotoxic products that induce neutrophil apoptosis. Neutrophil apoptosis without priming then results in ANCA antigen presence on the neutrophil cell membrane and the formation of MPO-ANCA. Myeloperoxidase-ANCA then binds to these membrane-bound antigens that cause self-perpetuating, constitutive activation through cross-linking with proteinase 3 or MPO and Fcγ receptors.

3. Activated neutrophils in the presence of hydrogen peroxidase release MPO that converts hydralazine into a cytotoxic product that is immunogenic for T cells that activate ANCA-producing B cells.

4. Histone H3 trimethyl Lys27 (H3K27me3) levels are perturbed in HIAV, which leads to aberrant gene silencing of proteinase 3 and MPO. In contrast, the demethylase Jumonji domain-containing protein 3 for the H3K27me3 histone is increased in patients without HIAV. Based on this data and the data showing a role for hydralazine in reversing epigenetic silencing of tumor suppressor genes in cancer cells,13 it has been proposed that hydralazine may reverse epigenetic silencing of proteinase 3 and MPO.

Diagnosing HIAV is still difficult because physicians do not recognize the drug as the etiologic agent, there is extensive variability in duration between starting the drug and onset of symptoms, and there often is a failure to order the appropriate laboratory and invasive tests needed for evaluation and diagnosis.3,5,8,10,12 Despite these difficulties, a set of criteria and practices for diagnosis are delineated in Table 1, with the key diagnostic feature being resolution with hydralazine cessation.1,5,7,12

A comprehensive drug history from at least 6 months prior to presentation is essential. Biopsies also are strongly encouraged to confirm the presence of vasculitis and to determine its severity.8,12 If renal biopsies are performed, they typically show scant IgG, IgM, and C3 deposition that is characteristic of ANCA-positive pauci-immune glomerulonephritis. Compared to hydralazine-induced lupus, renal involvement in the setting of HIAV has a relative lack of immunoglobulin and complement deposition with histopathology and immunostaining.11

Laboratory test results including serum MPO-ANCA (perinuclear ANCA) with coexisting elastase and/or lactoferrin autoantibodies is characteristic of HIAV. Antinuclear antibody, antihistone, anti–double-stranded DNA, and antiphospholipid antibodies along with low complement levels also may be present.2,4,9,10,13,15 It is recommended that ANCA assays combine indirect immunofluorescence with antigen-specific enzyme-linked immunosorbent

TABLE 1. Diagnostic Criteria for Hydralazine-Induced ANCA Vasculitis

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<tr>
<td>1. The 1994 Chapel Hill Consensus Conference definition of AAV should be fulfilled by all patients</td>
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<td>2. Signs and symptoms should have a temporal relationship to hydralazine and should subside with hydralazine cessation</td>
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<td>3. Patients should have a positive serum ANCA, especially if multiantigenicity (eg, antilactoferrin, antielastase) is present</td>
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<td>4. Exclude other vasculitides and mimickers</td>
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Abbreviations: ANCA, antineutrophil cytoplasmic antibody; AAV, ANCA-associated vasculitis.
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TABLE 2. Recommended Therapeutic Approaches for Treatment of Hydralazine-Induced ANCA Vasculitis

1. Withdraw the causal agents, avoid rechallenges, and consider avoiding similar drug classes.

2. If organ involvement has occurred, corticosteroids, glucocorticoid-sparing medications, and/or rituximab should be initiated. Prednisone can be administered at 1 mg/kg/d for the first 4–8 wk followed by a gradual taper over the next 6–12 mo.

3. If severe organ involvement occurs (eg, necrotizing glomerulonephritis, focal segmental necrotizing glomerulonephritis, diffuse alveolar hemorrhage), then methylprednisolone pulse therapy (ie, 1000 mg/d IV for 3 d) should be started followed by a combined corticosteroid and immunosuppressive drug regimen.

4. Consider plasmapheresis if massive pulmonary hemorrhage is observed.

Abbreviations: ANCA, antineutrophil cytoplasmic antibody; IV, intravenous.

REFERENCES


With respect to its idiopathic counterpart, patients may only present with MPO-ANCA, while other aforementioned antibodies (eg, histone antibody, double-stranded DNA) are rarely found or are entirely absent. Patients with HIAV often have higher titers of MPO-ANCA. In hydralazine-induced lupus, patients rarely have MPO-ANCA.

When a diagnosis of HIAV is made, it cannot be confirmed until hydralazine is discontinued and the patient’s symptoms resolve. Therefore, it is both diagnostic and therapeutic to discontinue hydralazine when HIAV is suspected. If recognized when the patient is only presenting with nonspecific symptoms, simple hydralazine cessation may be all that is needed; however, because recognition and diagnosis of HIAV is difficult, most patients present when the disease is severe and has progressed to organ involvement.

Treatment recommendations are highlighted in Table 2. Glucocorticoid therapy is believed to work by preventing T-cell and B-cell maturation needed to produce MPO-ANCA. Rituximab, on the other hand, is suspected to act by clearing the peripheral blood of MPO-ANCA B cells. Of note, patients with HIAV are suspected. If recognized when the patient is only presenting with nonspecific symptoms, simple hydralazine cessation may be all that is needed; however, because recognition and diagnosis of HIAV is difficult, most patients present when the disease is severe and has progressed to organ involvement.

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Our patient highlights the importance of identifying individuals at risk for HIAV. We seek to increase recognition of this entity, as it is not commonly seen in a dermatologic setting and is associated with high morbidity and mortality, as seen in our patient.