Aquagenic urticaria: diagnostic and management challenges

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Abstract: Aquagenic urticaria (AU) is a rare inducible form of physical urticaria, which occurs in response to cutaneous exposure to water, including sweat and tears. Patients present with characteristic 1–3 mm folliculocentric wheals with surrounding 1–3 cm erythematous flares within 20–30 minutes following skin contact with water. In rare cases, there are concomitant systemic symptoms, such as wheezing or shortness of breath. The pathogenesis of AU is poorly understood at this time, and it appears to be mediated in both a histamine-dependent and independent manner. Diagnosis is based on eliciting a thorough clinical history combined with a water challenge test. Some patients may need to undergo further testing to exclude other physical urticarias. Rarely, multiple physical urticarias can be present in one patient, which can complicate diagnosis and treatment. Currently, the first-line therapy for AU is an oral administration of nonsedating, second-generation H1 antihistamines, but many patients may require further interventions to have adequate symptomatic control. In this review, we discuss the diagnostic and management challenges of AU. We review the key diagnostic features that differentiate AU from other physical urticarias. We additionally describe a therapeutic ladder for the treatment of AU and the rationale supporting these treatments.

Keywords: Aquagenic urticaria, physical urticaria, inducible urticaria, diagnosis, management

Introduction

Water is ubiquitous in our daily lives and generally regarded as harmless, yet for some patients, water is a source of great discomfort. Aquagenic urticaria (AU), a form of physical urticaria that occurs with cutaneous exposure to water, including sweat and tears, was first described in the literature in 1964 by Shelley and Rawnsley.1 The latest World Allergy Organization guidelines on urticaria now classify AU as an inducible type of chronic urticaria, whereas this condition and other physical urticarias were previously grouped together due to their inducible nature by specific physical stimuli.2 In this review, we will use both terms, inducible urticaria and physical urticaria, synonymously to describe AU. While AU is very rare, with ~50 cases reported in the literature at the time of writing this review, it causes significant morbidity for affected patients.

Among the limited cases reported in the literature, there seems to be a higher prevalence among females with the disease onset typically occurring during puberty or postpuberty. However, there have been reports of childhood-onset disease.3–6 One familial cohort presented with coexistent Bernard–Soulier syndrome in the affected patients, raising the possibility of an associated genetic locus for AU;
however, no specific gene or locus for AU has been identified thus far. AU has also rarely been reported in association with systemic conditions, including HIV infection, and occult papillary carcinoma of the thyroid gland.8,9

The mechanism of AU is poorly understood. In the 1960s, Shelley and Rawnsley were the first to propose a mechanism; they hypothesized that water reacts with sebum or sebaceous glands to form a toxic substance, which stimulates mast cell degranulation and subsequent histamine release, leading to the development of urticarial lesions.1 In 1981, Tkach hypothesized that the mechanism of AU has to do with sudden changes in osmotic pressure surrounding hair follicles, leading to increased passive diffusion of water.10 This sudden change in the pressure results in indirect provocation of urticaria. More recently, Gallo et al have described cases of localized AU following epilation, which seem to corroborate Tkach’s hypothesis.11,12 Another proposed mechanism involves existence of water-soluble antigens in the epidermis, which dissolves and diffuses across the dermis with resulting histamine release.13 A more recent study by Luong and Nguyen in 1998, however, suggests a mechanism that may be completely independent of histamine release; they reported several patients with AU who exhibited no increase in histamine levels upon exposure to water, which was sufficient to induce urticaria.3 Depending on the sensitivity of the assay used, a rise in the plasma histamine level may not have been sufficiently detected in these patients. Regardless, a histamine-independent mechanism is still conceivable based on the observation that pretreatment with scopolamine (acetylcholine antagonist) prior to contact with water can suppress wheal formation.14 The lack of a clear pathogenesis for AU has contributed to difficulty in proposing evidence-based treatments for affected patients.

Clinical presentation

Patients with AU will present with characteristic 1–3 mm folliculocentric wheals and surrounding 1–3 cm erythematous flares within 20–30 minutes following skin contact with water.1 Patients can also experience associated symptoms, including pruritus, burning, and uncomfortable prickling.1 Urticarial lesions will typically resolve within 30–60 minutes of cessation of water contact with the skin. Lesions most commonly appear on the trunk and upper arms, usually sparing the palms and soles. The affected areas are generally refractory to repeated stimulations for several hours. Rarely, patients can experience systemic symptoms such as wheezing or shortness of breath.3,8

There are also some uncommon clinical presentations of AU. While urticaria is thought to occur in response to any form of water in AU, there have been reports of patients who have had reactions depending on the salinity of the water. For example, a patient reacted to tap water, snow, and sweat, but could still swim in the ocean without urticaria.11,15 In patients who have decreased thickness of the stratum corneum following epilation or cutaneous exposure to organic solvents, there can be an exaggerated urticarial response to water.3,10 Additionally, in AU patients with associated systemic disease, the urticarial response seems to be often more dramatic, consisting of large edematous plaques rather than the classic punctate perifollicular wheals.8,9

Diagnostic challenges

Diagnosis of AU is largely based on a history of recurrent urticaria after exposure to water combined with a water challenge test. The test can be administered in a variety of ways; however, the standard method is to apply water at room temperature to a cloth and apply this damp cloth to the patient’s skin for 20 minutes, with an urticarial reaction indicating a positive test.2 The temperature of water applied to the patient’s skin is important because significant heat or cold exposure can potentially induce other physical urticarias, giving a false-positive result. A physical examination should also focus on testing for dermatographism and searching for any systemic signs such as wheezing. On laboratory evaluation, the level of serum immunoglobulin E should be normal, which can help to differentiate AU from immunoglobulin E-mediated allergic reaction. When patients present with angioedema along with urticaria, it might be helpful to investigate the etiology of angioedema by checking the level of C1-esterase inhibitor. Hereditary and acquired angioedema present with angioedema only and the level of functional C1-esterase inhibitor should be normal. It is important to note that the serum histamine levels may or may not be elevated following exposure to water.3 Finally, the histopathology of AU is consistent with nonspecific urticaria; therefore, it does not play a role in guiding the diagnosis.

The main challenge in diagnosing AU lies in differentiating this condition from other types of physical urticaria (eg, cholinergic urticaria, heat urticaria, cold urticaria, pressure urticaria, and exercise-induced urticaria). Patients should be subjected to provocative testing for these specific types. The lesions of cholinergic urticaria appear extremely similar to those of AU, but will arise in response to cholinergic stimuli (ie, the rise in the internal core body temperature), such as exercise, sweating, stressful emotions, or eating spicy food. Therefore, cholinergic urticaria will test negative in response to room temperature water challenge.3 However, it
is possible, albeit rare, for patients to have concomitant physical urticarias, which can further complicate the diagnostic process.\textsuperscript{16,17} For example, Bayle et al described a case of a female with AU, dermatographism, and cholinergic urticaria whose cholinergic urticaria responded to oral cetirizine but without resolution of AU.\textsuperscript{16} As AU, cholinergic, cold, and heat urticarias can all be induced by exposure to water, it is important to differentiate among these conditions with careful evaluation and appropriate testing (Table 1).

In generating differential diagnoses, one must also consider clinical subtypes of AU. Aquagenic pruritus is a condition in which the patient will develop pruritus without any skin lesions after exposure to water.\textsuperscript{19} Unlike AU, aquagenic pruritus is associated with polycythemia vera and is unresponsive to conventional AU therapies. Salt-dependent AU is a condition with specificity for the salinity of water. As such, the patients will develop urticaria in response to seawater (and 3.5% NaCl solution, which is isosmolar to seawater), but not tap water or hyperosmolar and nonionic solution such as 20% glucose.\textsuperscript{12} Several reports describe a localized version of salt-dependent AU in young females.\textsuperscript{11,12,15} Finally, it is important to ask patients about family history, as there is a reported cohort with familial AU.

### Table 1 Differential diagnosis of aquagenic urticaria

<table>
<thead>
<tr>
<th>Differential diagnosis</th>
<th>Distinguishing features</th>
<th>Diagnostic testing</th>
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<tbody>
<tr>
<td>Aquagenic urticaria</td>
<td>Distinguish from evaporative cooling and cold urticaria</td>
<td>Wet cloth at room temperature applied for 20 minutes</td>
</tr>
<tr>
<td>Salt-dependent aquagenic urticaria</td>
<td>Sea water provokes lesions, with or without symptoms from tap or distilled water</td>
<td>3.5% NaCl at room temperature soaked in wet cloth applied for 20 minutes</td>
</tr>
<tr>
<td>Aquagenic pruritus</td>
<td>Pruritus without the appearance of skin lesions</td>
<td>Wet cloth at room temperature applied for 20 minutes</td>
</tr>
<tr>
<td>Dermatographism</td>
<td>Linear, pruritic hives from shear force, the most common physical urticaria</td>
<td>Linear stroking at various pressures (20–144 g/m\textsuperscript{2}) using dermatographometer, Fric test, or ballpoint pen</td>
</tr>
<tr>
<td>Cholinergic urticaria</td>
<td>Pinpoint diffuse papular lesions from increase in internal core body temperature</td>
<td>Exercise challenge to induce sweat for &gt;10 minutes or passive warming using hot water bath to raise body temperature &gt;1°C</td>
</tr>
<tr>
<td>Cold urticaria</td>
<td>Pruritic wheal and flare from cold contact, up to one-third of cases of physical urticaria</td>
<td>Placement of ice water in 50 mL beaker for 1–10 minutes, cold hand immersion for 5 minutes, total body cold exposure, evaporative cooling</td>
</tr>
<tr>
<td>Delayed pressure urticaria</td>
<td>Pruritus, swelling, and pain 4–8 hours after exposure may be associated with systemic symptoms of fatigue and arthralgia</td>
<td>100 g/m\textsuperscript{2} of pressure for 5–180 seconds on the forearm using a dermatographometer or 15 lb weight bearing on the shoulder or the lower leg for 15–20 minutes</td>
</tr>
<tr>
<td>Exercise-induced urticaria</td>
<td>Not induced by passive warming, larger lesions often associated with systemic symptoms</td>
<td>Exercise challenge as above</td>
</tr>
<tr>
<td>Local heat urticaria</td>
<td>Reaction limited to area of exposure</td>
<td>Placement of hot water (45°C–50°C) in beaker for 1–10 minutes</td>
</tr>
<tr>
<td>Solar urticaria</td>
<td>Immediate reaction to UV and visible light, resolves within 24 hours, distinguish from polymorphous light eruption</td>
<td>UVA, UVB, and visible light stimulation of variable intensity to establish minimal urticarial dose</td>
</tr>
<tr>
<td>Vibratory angioedema</td>
<td>Erythema and swelling beyond provocation site</td>
<td>Vortex vibratory stimulation for 4 minutes at 2,500 rpm</td>
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</tbody>
</table>


Abbreviation: UV, ultraviolet.
Table 2 Therapeutic options for aquagenic urticaria

<table>
<thead>
<tr>
<th>Therapeutic options</th>
<th>Line of therapy/notes</th>
<th>Proposed mechanism</th>
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<tbody>
<tr>
<td>Nondesatining, second-generation H1 antihistamines (eg, cetirizine)</td>
<td>First line at standard dose, second line consider dose increase up to fourfold</td>
<td>Antagonism or inverse agonism of H1 receptor, preventing histamine effects</td>
</tr>
<tr>
<td>First-generation H1 antihistamines (eg, hydroxyzine)</td>
<td>Third line</td>
<td></td>
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<tr>
<td>H2 antihistamines (eg, cimetidine)</td>
<td>Third line</td>
<td></td>
</tr>
<tr>
<td>Acetylcholine antagonists (eg, scopolamine)</td>
<td>Adjuvant with H1 antihistamines</td>
<td></td>
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<tr>
<td>Phototherapy (eg, psoralen plus UVA, UVB)</td>
<td>Adjuvant with oral therapy or second-line alone</td>
<td>Reduction of mast cell activity, reactive thickening of the epidermis</td>
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<tr>
<td>Topical barrier creams (eg, petrolatum)</td>
<td>First line or adjuvant</td>
<td>Hydrophobic effect prevents water penetration into skin</td>
</tr>
<tr>
<td>Anabolic androgenic steroid (eg, stanozolol)</td>
<td>Case-specific, used in HIV patient with AU</td>
<td>Increases C1 esterase inhibitor synthesis</td>
</tr>
<tr>
<td>Selective serotonin reuptake inhibitors (eg, fluoxetine)</td>
<td>Case-specific, used in patient with extracutaneous symptom of AU</td>
<td>Unknown</td>
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Abbreviations: AU, aquagenic urticaria; HIV, human immunodeficiency virus, UV, ultraviolet.

Finally, there are case reports supporting the use of stanozolol and selective serotonin reuptake inhibitors in the treatment of AU. Stanozolol is an anabolic steroid without significant androgenic side effects; it has been shown to increase the levels of normal C1 esterase inhibitors in the management of hereditary angioedema. In one male patient with AU and associated systemic symptoms, 10 mg of stanozolol per day was successful at controlling his symptoms.8 Another patient with migraine-like headache upon exposure to water along with urticaria was able to achieve symptomatic control with the addition of a selective serotonin reuptake inhibitor to her medication regimen of antihistamines and anticholinergics. This raises an interesting possibility of serotonin involvement in the pathogenesis of AU.

Conclusion

While water may be a seemingly innocuous substance to most of us, there are patients who suffer from side effects of exposure to water. A clinical history of hives in response to water exposure, as well as a positive result of a water challenge test, are keys to diagnosing AU. It is critical to differentiate AU from other physical urticarias. Therefore, careful evaluation is necessary to rule out other physical stimuli that can mimic the effects of water alone. Patients may need to undergo other tests described in Table 1, depending on the clinical scenario, for a clinician to arrive at the diagnosis of AU.

There is still room for improvement in understanding the mechanism of AU, which could lead to more evidence-based, efficacious treatment strategies for patients. At this time, however, the cornerstone therapy for these patients is...
nonsedating, second-generation H₁ antihistamines. While there are several therapeutic options currently available, including other oral agents, topical agents, and phototherapy, there is a lack of strong evidence of their effectiveness. Some patients may be refractory to the medical management described earlier and still need to rely on minimizing water exposure by limiting bathing time and avoiding certain water-based activities.

**Future directions**

A familial case of AU associated with Bernard–Soulier syndrome (with a known genetic locus) represents a potential avenue for genetic linkage studies that may uncover the genomic alterations involved in this disease. More research is certainly needed to delineate the pathogenesis of AU, which will in turn help us develop effective therapies for AU. A large-scale study will be instrumental in this effort, yet has been difficult to execute due to the limited number of AU patients reported thus far. It is possible that AU is underrecognized and under reported in the general population. By raising awareness of this condition among health care providers, it may be possible to identify a larger group of patients to assist in further study.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**