Anticonvulsant hypersensitivity syndrome (AHS) is a nondose-related idiosyncratic reaction to aromatic antiepileptic drugs and is a cause of drug discontinuation. Pseudomonas aeruginosa is a gram-negative bacillus that can produce infections in many different organs, including the skin and soft tissue. We report a patient with erythroderma and AHS who developed a pseudomonal infection. Green staining of the underwear served as a diagnostic clue for severe P aeruginosa infection that had developed because of a local flexural skin infection that spread due to a damaged skin barrier. Inspection of the patient’s clothes may give information about any exudate from the skin and should be done routinely as part of the physical examination.

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Anticonvulsant hypersensitivity syndrome (AHS) is a cutaneous drug eruption to aromatic antiepileptic drugs and consists of cutaneous eruption; fever; facial edema; lymphadenopathy; leukocytosis; and internal organ involvement such as nephritis, hepatitis, or pneumonia.\(^1\)\(^2\) Usually 2 to 8 weeks after initiation of antiepileptic therapy, a cutaneous eruption ranging from mild morbilliform lesions to severe toxic epidermal necrolysis can be seen.\(^3\)\(^4\) Among the antiepileptics, carbamazepine, phenytoin, and phenobarbital are the common causes of AHS.\(^5\)\(^6\) Erythroderma is one of several cutaneous manifestations of AHS. Erythroderma is defined as extensive erythema and/or scaling of the skin, and the clinical diagnosis is based on involvement of 90% body surface area. Because of disruption of skin integrity, erythroderma places the patient at high risk for secondary infections.\(^7\) Early published case series of erythroderma reported a notable mortality rate from complications including sepsis.\(^8\) Pseudomonas aeruginosa is a gram-negative bacillus and can produce infections in many different organs, including the skin and soft tissue. As an opportunistic pathogen, P aeruginosa seldom causes disease in healthy patients. For an infection to occur, some disruption of the physical barriers (skin or mucous membranes) and/or an underlying dysfunction of the immune system generally is necessary.\(^9\) Pseudomonas aeruginosa may cause severe infections that can be difficult to treat if not diagnosed early. Pseudomonas wound infections often are indistinguishable from similar infections caused by other etiologic agents. Pseudomonas aeruginosa is capable of producing multiple hydrosoluble pigments, most commonly pyocyanin (blue) and pyoverdin (nonfluorescent greenish color). On occasion, by producing these pigments, patients with Pseudomonas infection may present with a green exudate with a characteristic grapefruit odor. These signs are important clues for prompt diagnosis and treatment of potentially fatal severe Pseudomonas infections.\(^10-12\)

We present the case of a 31-year-old woman with erythroderma and green discoloration of clothing to emphasize that these findings can be important indicators for the diagnosis of Pseudomonas infection.

Case Report
A 31-year-old woman was admitted to the emergency department and reported a high fever, redness of skin, and facial puffiness of 5 days’ duration. She had developed generalized erythema with edema and pruritus 7 weeks after initiation of carbamazepine therapy for
epilepsy. She had been previously hospitalized with an unknown diagnosis, treated with broad-spectrum antibiotics (ie, vancomycin, cefazolin), and discharged within a couple of days. With the deterioration of her symptoms, the patient presented to our emergency department.

Upon admission, the patient’s temperature was 39°C. Her heart rate was normal but tachycardic. Skin examination revealed diffuse erythema, induration, and desquamation including the face, trunk, and limbs (Figure 1). More than 90% body surface area was involved. Axillary and inguinal skin was macerated and had fissures. Thick hyperkeratosis of the palmoplantar regions was observed. Facial edema was present with hyperemic conjunctivae. A painless, mobile, 0.5×0.5-cm left postauricular lymph node was detected. The remaining systemic examination did not reveal any abnormalities. During physical examination, green staining of the patient’s underpants and axillary part of her undershirt also were noted (Figure 2). She had neither a history of preexisting dermatoses prior to admission nor a history of malignancy.

Laboratory examination revealed the following values: hemoglobin, 11.1 g/dL (reference range, 14.0–17.5 g/dL); hematocrit, 34.9% (reference range, 41%–50%); white blood cell count, 27,600/μL (reference range, 4500–11,000/μL); platelet count, 659,000/μL (reference range, 150,000–350,000/μL); erythrocyte sedimentation rate, 20 mm/h (reference range, 0–20 mm/h); and C-reactive protein, 36.41 mg/L (reference range, 0.08–3.1 mg/L). In a peripheral blood smear, 40% eosinophils, 30% neutrophils, 16% lymphocytes, and 10% monocytes were detected. Except for low blood protein and albumin levels, other biochemical parameters were within reference range. Pseudomonas aeruginosa was isolated from the hemocultures taken from the patient during the febrile period.

Based on the patient’s history as well as clinical and laboratory findings, carbamazepine-dependent AHS was diagnosed. The patient was hospitalized and carbamazepine treatment was immediately stopped. Emollients constituted the topical treatment of the erythroderma. Systemic antihistamines were administered to relieve pruritus. To manage pseudomonal infection, intravenous cefepime hydrochloride 2 g twice daily was started. Fever subsided within 72 hours of antibiotic therapy. By the end of the second week of treatment, all dermatologic findings had subsided with the exception of palmoplantar hyperkeratosis. Eosinophil and white blood cell leukocyte counts returned to normal and antibiotic therapy was stopped by the 14th day of hospitalization.

**Comment**

Anticonvulsant hypersensitivity syndrome is a nondose-related idiosyncratic reaction to aromatic antiepileptic drugs and is a cause of drug discontinuation. This syndrome also can be seen with the use of sulfonamides, allopurinol, gold salts, and dapsone. The most common features are cutaneous eruption, prominent eosinophilia, fever, and multisystemic involvement. Cutaneous manifestations associated with this syndrome range from a mild morbilliform eruption to severe toxic epidermal necrolysis or erythroderma. The variability in the clinical presentation of AHS is likely to reflect host-specific metabolic and immunologic reactions associated
with T-cell–mediated responses.\textsuperscript{15} Onset of the syndrome usually occurs 2 to 8 weeks after initiation of antiepileptic agents. The mortality rate of AHS ranges from 10% to 20%. Prompt diagnosis of the syndrome is essential to prevent mortality. The most important therapeutic approach is discontinuation of the causal drug.\textsuperscript{16-18}

Erythroderma is characterized by generalized erythema, induration, and scaling of the skin. Secondary infections are potential complications in erythrodermic cases.\textsuperscript{15} 

\textit{Pseudomonas aeruginosa} is an opportunistic, aerobic, gram-negative bacillus that rarely causes infection in healthy individuals but is a frequent cause of severe nosocomial infections in immunocompromised patients.\textsuperscript{11} Occasionally it can colonize human body sites, with a preference for moist areas such as the perineum, axillae, ears, nasal mucosa, and throat. The prevalence of \textit{P aeruginosa} colonization in healthy patients usually is low, but higher colonization rates can be encountered following hospitalization, especially among patients treated with broad-spectrum antimicrobial agents.\textsuperscript{12} 

A defective skin barrier usually is caused by destruction of intact stratum corneum due to scratching, dermatitis, maceration, heat, or ulcerations. Primary skin infections may show greenish color and give fluorescence by Wood lamp examination. Our patient with erythroderma had been previously hospitalized and had used broad-spectrum antibiotics, which are both risk factors for the development of \textit{Pseudomonas} infection. Green staining on her underwear was noticed during physical examination by chance, which was the most important clue for the diagnosis of \textit{P aeruginosa} infection.\textsuperscript{9,12} Kaya et al\textsuperscript{19} reported a similar case involving blue underpants sign in the inguinal area, which was indicative of \textit{P aeruginosa} infection. In our patient, this sign was seen on multiple areas of the underwear in contact with the inguinal and axillary regions. Green discoloration of the underwear served as a diagnostic clue for severe \textit{P aeruginosa} infection that had developed because of a local flexural skin infection that spread due to a damaged skin barrier.

Anticonvulsant hypersensitivity syndrome may necessitate systemic glucocorticosteroid treatment.\textsuperscript{3} However, caution is warranted when considering use of corticosteroids. Glucocorticosteroid treatment of patients with a pseudomonal infection may increase the rate of morbidity and mortality.\textsuperscript{20} Thus, in patients with erythroderma and AHS, it is imperative to exclude the additional diagnosis of infection prior to initiation of corticosteroid treatment, as both conditions present with fever, malaise, tachycardia, hepatic dysfunction, and leukocytosis.\textsuperscript{4} Our patient was not started on corticosteroids due to initial suspicion of systemic pseudomonal infection following inspection of her green underwear.

**Conclusion**

The presence of green staining on the underwear should be a warning sign of the presence of \textit{Pseudomonas} infection. All physicians should be familiar with this sign, especially in patients with erythroderma because its recognition may enable early treatment, thereby improving the chances of survival. The prognosis of \textit{Pseudomonas} infection is poor. Inspection of the patient’s clothes may give information about any exudate from the skin and should be done routinely as part of the physical examination. In summary, examination of not just the patient but also the attire is a rewarding medical practice.

**REFERENCES**