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Bullous eruption on the posterior thigh

A healthy 11-year-old girl visited her family physician with a lesion on her right posterior thigh. The lesion was a 1-cm plaque that was tender, firm, erythematous, and indurated, with a central pustule. It had been present for 3 days; it was noticed by the patient after returning from a camping trip in southeastern Pennsylvania. The pustular area was incised, drained, and cultured, and the patient was started on cephalexin.

Two days later, the lesion did not improve, showing increased induration, erythema, and blistering. The patient went to the emergency department with an 8 cm by 6 cm coalescence of thin-walled vesicles and bullae with surrounding erythema (FIGURES 1 AND 2). A thick, honey-yellow adherent crust covered the eroded center of the lesion. The girl’s temperature was 37.1°C, and she reported no burning, pain, or pruritus. She had full range of motion of her right hip and knee, and no lymphadenopathy was detected. Her white blood cell count was normal; blood and wound cultures were taken.

■ What is the most likely diagnosis?
■ How would you empirically treat this condition?
PHOTO ROUNDS

- **Diagnosis: Bullous impetigo, caused by methicillin-resistant S aureus**
  Impetigo is a highly contagious superficial skin infection, with peak incidence among children aged 2 to 6 years. Nonbullous impetigo (70% of cases) is caused by *Staphylococcus aureus* or betahemolytic *Streptococcus*. Bullous impetigo is almost always caused by *S aureus*. Epidermolytic toxins produced by phage group II strains cause loss of cell adhesion in the stratum granulosum due to proteolytic attack of desmoglein 1, resulting in bullae.

  Bullous impetigo may occur after minor skin injury, such as an insect bite, abrasion, or dermatitis. Lesions generally start as small vesicles on the face, buttocks, extremities, or perineum, and may progress to a coalescence of thin-roofed bullae. The flaccid bullae rupture easily, draining serous or purulent fluid.

  Lesions are usually painless, and systemic findings are rare. Lymphadenopathy is rare in bullous impetigo but common in nonbullous impetigo. The disease is generally self-limited and complications are uncommon. However, ecthyma (ulcerative impetigo) may result from an untreated impetigo infection.

- **Differential diagnosis**
  The differential diagnosis for bullous impetigo is broad, and may include allergic contact dermatitis, herpes simplex, herpes zoster, pemphigus foliaceous, bullous pemphigoid, pemphigus vulgaris, and (in this case specifically) erythema migrans. **Allergic contact dermatitis** is a delayed hypersensitivity reaction, usually caused by skin contact with an allergen. Lesions can be vesicular, edematous, erythematous, and pruritic. In this case, the patient did not have allergen exposure or a pruritic lesion.

  **Herpes zoster** is a reactivation of the varicella zoster virus, characterized by stabbing, neuritic pain in a dermatomal distribution. Clear vesicles on an erythematous, edematous base distributed along a dermatome constitutes the classic appearance. This was not the case with this patient.

  **Pemphigus foliaceous** is an autoimmune intraepidermal blistering disease with lesions occurring on the face, scalp, chest, and upper back. Intact blisters are not commonly seen. The vesicle roof is very thin and ruptures easily, forming broad areas of crust. Skin biopsy reveals intraepidermal bulla or acantholysis in the upper epidermis.

  **Pemphigus vulgaris** is also an autoimmune blistering disease that affects the skin and mucous membranes. It is generally seen among patients aged >40 years.

  **Bullous pemphigoid** is an autoimmune disorder presenting with chronic eruption of erythematous, papular, urticaria lesions often evolving into bullae. Childhood cases are rare. Biopsy of the lesions demonstrates subepidermal bulla with an infiltration of eosinophils within the dermis.

  **Erythema migrans** with central vesiculation must be considered given the patient's camping trip. Recent evidence shows that erythema migrans with central redness accounts for most cases in areas endemic for Lyme disease. Only 10% of the patients with early Lyme disease show the classic bulls-eye lesion with concentric erythematous rings and central clearing. Vesiculation can occur in up to 30% of lesions.

- **Staphylococcus aureus and antibiotic resistance**
  As many as 61% of community-acquired methicillin-resistant *S aureus* (MRSA) infections are initially treated only with beta-lactam antibiotics, to which they are resistant. Risk factors for community-acquired MRSA infection include day-care attendance, recent hospitalization, recent antibiotic use, chronic illness, and frequent health care visits. A growing number of cases are reported among patients without risk factors.

FAST TRACK

Risk factors for community-acquired MRSA infections include attending day care, recent hospital visit or antibiotic use, or chronic illness.
Community-acquired MRSA isolates are usually genetically different from nosocomial isolates, and have been relatively susceptible to non–beta-lactam antibiotics. These strains vary substantially, however, and it is important to check the susceptibility of the isolate.

Virulent new strains of \( \text{S aureus} \) are infecting children—these strains have a novel transpeptidase, which offers them a mechanism of resistance to beta-lactams different from hospital- and community-acquired types.

Awareness of the local antimicrobial susceptibility patterns of community \( \text{S aureus} \) isolates is also helpful. Oral antibiotics that have been successful include clindamycin, minocycline, doxycycline, and trimethoprim-sulfamethoxazole. Cephalexin has no therapeutic value in treating community-acquired MRSA.

**Empiric treatment of impetigo: Consider a culture for MRSA**

For localized impetigo, topical therapy with mupirocin 2% ointment 3 times a day for 10 days is usually adequate. A 10-day course of oral antibiotic therapy with dicloxacillin or cephalexin is indicated in more widespread impetigo presumed to be methicillin-sensitive \( \text{S aureus} \). Azithromycin (Zithromax) or clarithromycin (Biaxin) may be given to patients allergic to penicillin.

However, it is becoming increasingly important to consider community-acquired methicillin-resistant \( \text{S aureus} \) species in cases such as this that do not respond to traditional therapy. Hence, culture and sensitivity of all suspicious lesions is highly suggested.

**Patient’s treatment and recovery**

In this case, the patient was diagnosed with bullous impetigo and admitted to the hospital. She was started on intravenous clindamycin at 380 mg (30 mg/kg) every 8 hours. Clindamycin was chosen because most cases of community-acquired MRSA in this geographic area are resistant to trimethoprim-sulfamethoxazole and susceptible to clindamycin.

Although doxycycline would have covered both community-acquired MRSA and Lyme disease, we were less suspicious of Lyme given the physical exam of the patient, and we were reluctant to start this patient on doxycycline due to the fact she did not have complete maturation of her dentition.

Within 24 hours of intravenous clindamycin, the lesion was markedly improved and the culture confirmed that the MRSA was sensitive to clindamycin. She was discharged on oral clindamycin at 375 mg 3 times daily, to complete a 14-day course of therapy. The lesion was completely resolved without recurrence within 2 weeks.
REFERENCES


