Chronic superficial ulcers on the trunk



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Key words: chronic ulcer; pyoderma gangrenosum; superficial granulomatous pyoderma.





CASE

A 62-year-old man presented with several painful superficial ulcers on the trunk that began to expand slowly 1 year after inguinal hernia surgery. Physical examination demonstrated 5 ulcerative plaques on a violaceous base with overlying crust (Figs 1 and 2). A punch biopsy was performed, and tissue bacterial, fungal, and acid-fast bacilli cultures demonstrated no growth.

Question 1: Which of the following is the best diagnosis?

- **A.** Superficial granulomatous pyoderma (SGP)
- B. Ecthyma
- C. Blastomycosis-like pyoderma
- D. Chromomycosis

E. Factitious ulcer

Answer:

A. SGP – Correct. SGP is an uncommon variant of pyoderma gangrenosum (PG) characterized by slowly progressive superficial ulcers with vegetative margins and a clean granulating base. Unlike classic PG, SGP most commonly occurs on the

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Funding sources: None.

IRB approval status: Not applicable.

Patient consent: Consent for the publication of all patient photographs and medical information was provided by the authors at the time of article submission to the journal stating that all patients gave consent for their photographs and medical information to be published in print and online and with the understanding that this information may be publicly available.

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JAAD Case Reports 2022;27:121-3.

2352-5126

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https://doi.org/10.1016/j.jdcr.2022.07.023

trunk and is not significantly associated with systemic conditions.¹

- **B.** Ecthyma Incorrect. Ecthyma is a deep, ulcerative version of impetigo that is characterized by punched-out ulcers, most commonly observed on the lower extremities. The most commonly implicated bacteria include *Streptococcus pyogenes* and *Staphylococcus aureus*.
- **C.** Blastomycosis-like pyoderma Incorrect. Blastomycosis-like pyoderma is characterized by large verrucous plaques with pustules and an elevated border and is most commonly observed on the distal extremities.
- **D.** Chromomycosis Incorrect. Chromomycosis most often presents as a solitary verrucous nodule or plaque on an acral anatomic site.
- **E.** Factitious ulcer Incorrect. Factitious ulcers often demonstrate irregular or geographic borders and are induced by patients who assume the sick role without secondary gain.

Question 2: What is the most common anatomic site observed in this disorder?

- A. Trunk
- **B.** Extremities
- C. Face
- **D.** Anogenital region
- E. Head and neck

Answer:

- **A.** Trunk Correct. SGP most commonly occurs on the trunk. Disseminated disease affects multiple anatomic sites.¹
- **B.** Extremities Incorrect. Classic ulcerative PG typically affects the lower extremities.
- **C.** Face Incorrect. While SGP is rarely reported on the face, this is not the most common anatomic site. SGP on the face appears to be refractory to topical and systemic corticosteroids.²
- **D.** Anogenital region Incorrect. The anogenital region is rarely described in SGP.³
- **E.** Hands and neck Incorrect. This is not a typical location for SGP.

Question 3: Which of the following descriptions reflects the histopathologic findings in this disorder?

- **A.** Necrotic epidermis, ulcer, and sparse inflammation
- **B.** Pigmented yeast with internal transverse septae
- **C.** Ulceration with dense neutrophilic infiltrate and dermal necrosis
- **D.** Layered suppurative granuloma with necrosis and suppuration surrounded by histiocytes and plasma cells, along with overlying epidermal hyperplasia
- **E.** Pseudoepitheliomatous hyperplasia with neutrophilic abscesses and scar-like fibrosis

Answer:

- **A.** Necrotic epidermis, ulcer, and sparse inflammation Incorrect. These are nonspecific findings typically observed in factitious ulcer.
- **B.** Pigmented yeast with internal transverse septae Incorrect. Medlar or sclerotic bodies undergo binary fission and are characteristic of chromomycosis.
- **C.** Ulceration with dense neutrophilic infiltrate and dermal necrosis Incorrect. This describes findings present in classic ulcerative PG, which lacks the layered suppurative granuloma typical of SGP.¹⁻⁴
- **D.** Layered suppurative granuloma with necrosis and suppuration surrounded by histiocytes and plasma cells, along with overlying epidermal hyperplasia Correct. While nonspecific, these findings are consistent in SGP.¹⁻⁴
- **E.** Pseudoepitheliomatous hyperplasia with neutrophilic abscesses and scar-like fibrosis Incorrect. Blastomycosis-like pyoderma demonstrates these features; isolation of a pathogenic organism such as *Staphylococcus aureus* is required for diagnosis.

Abbreviations used:

PG: pyoderma gangrenosum SGP: superficial granulomatous pyoderma

Conflicts of interest

None disclosed.

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