

Hidradenitis Suppurativa-Like Tuberculosis: A Warning for Dermatologists

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Introduction

Tuberculosis (TB) ranks among the leading causes of death worldwide and because of this alarming statistic, its extrapulmonary forms are also becoming more prevalent [1,2]. The cutaneous presentations are highly variable, ranging from papules to ulcero-vegetative lesions, posing a challenge for dermatologists who may confuse it with other common conditions such as hidradenitis suppurativa (HS) [2,3]. This paper aims to describe the correlation between these often neglected diseases and their potential diagnostic confusion.

Case Presentation

A 17-year-old male student without comorbidities, presented to the Dermatology Outpatient Clinic with painful nodules that had been abscessing and draining purulent discharge in the thoracic and axillary regions for the past 8 months. Despite previous antibiotics prescribed by general practitioners, there had been no improvement. He denied fever, weight loss

or any decline in general health. On examination, he had a painful nodule in the right axilla and ulcerations in the anterior axillary line, symmetrically located on both sides. In the sternal region, four ulcerated plaques with granulation tissue and undermined and violaceous borders (Figure 1). The hypothesis of HS was considered but due to atypical presentation, differential diagnoses with ulcerovegetative pattern diseases were regarded. The patient returned with negative cultures but with a positive molecular rapid test (MRT) for tuberculosis, confirming the diagnosis of cutaneous TB. Standard anti-tuberculosis chemotherapy was initiated and maintained for six months, resulting in the complete resolution (Figure 2).

Conclusions

The mimicking of cutaneous TB as HS was also noted by other authors [4,5]. However, in this case, further investigation was carried out at the initial assessment due to the presence of advancing ulcerations, the lack of involvement



Figure 1. A 17-year-old man presenting ulcers with purulent exudate drainage. (A) Nodule and ulcer on right armpit. (B) Ulcer on left armpit. (C) Ulcers on sternal region.

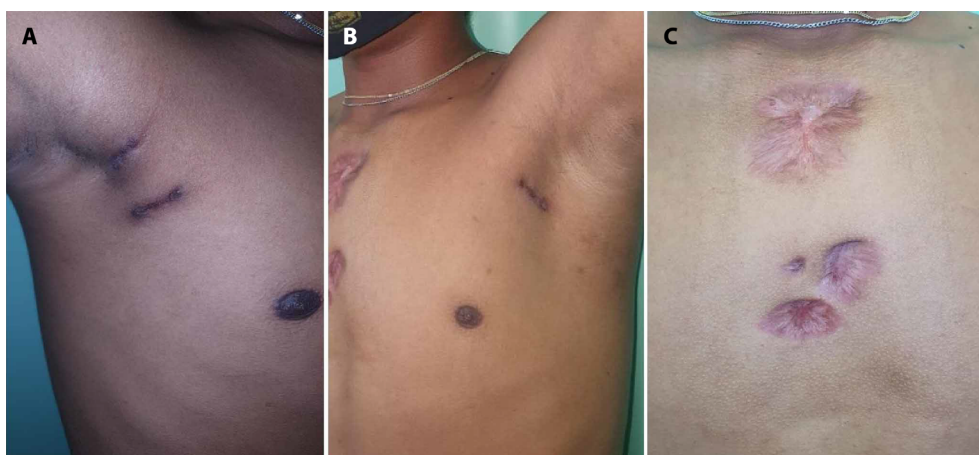


Figure 2. A 17-year-old man treated for cutaneous tuberculosis. Scars from ulcers after 3 months of standard polychemotherapy. (A) Right armpit. (B) Left armpit. (C) Sternal region.

of other cutaneous folds, and no improvement with tetracyclines. The clinical manifestation of the disease was consistent with the scrofuloderma subtype, characterized by nodules that coalesce into fistulas, draining purulent material and lead to ulceration [6]. However, it drew attention due to its unusual presentation with multiple and symmetrical lesions, an atypical manifestation for an infectious disease, resembling inflammatory conditions. Another key aspect is that the patient, a healthy young individual, has not exhibited pulmonary or systemic symptoms, distinguishing this case from previously reported ones.

Melo and colleagues also documented scrofuloderma following therapeutic attempts for pyoderma gangrenosum and HS [4]. A case involving deeper tissues has also been presented by Ermertcan and colleagues who described a severe case of Pott disease (vertebral tuberculosis) that was mistakenly treated as HS [5]. The diagnostic confusion can be explained by the clinical similarities, highlighted by the HS criteria, which include morphology, distribution of lesions

(intertriginous areas), and a protracted course. Given the multiple presentations, tuberculosis may mimic more prevalent diseases, including HS but also bacterial abscesses, tertiary syphilis, and tumor metastases [2,6].

This study emphasizes the need to reconsider atypical HS cases, regardless of systemic symptoms or immunocompromised status. Due to the surgical nature of HS treatment and the emerging immunosuppressive therapies, which can exacerbate and spread pre-existing TB, assertiveness is critical. Whilst we do not recommend biopsy in all cases of HS, caution is advised, emphasizing that diagnostic testing for TB should be performed in atypical presentations.

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