

A case of secondary syphilis mimicking palmoplantar psoriasis in HIV infected patient*

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Abstract: Due to diverse clinical and histopathological presentations, diagnosis of secondary syphilis can occasionally prove challenging. Variable clinical presentations of secondary syphilis in HIV disease may result in an incorrect diagnosis and an inappropriate treatment regimen. Similarly, the histology of secondary syphilitic lesions may show considerable variation, depending on the clinical morphology of the eruption. We report a case of secondary syphilis in an HIV infected patient with cutaneous palmoplantar lesions simulating palmoplantar psoriasis.

Keywords: Histology; Nails; Psoriasis; Syphilis; Treponemal infections; Treponema pallidum

INTRODUCTION

Syphilis is a sexually transmitted disease caused by the *Treponema pallidum*, whose prevalence has been increasing worldwide. It is considered "the great imitator" due to its varied presentations. The HIV virus has many effects on clinical presentation, diagnosis, disease progression and treatment of syphilis. ¹ Even though syphilis presentation in these patients is for the most part similar to that of patients without HIV, atypical manifestations of the disease may be present, including severe and rare forms. ²⁻⁵

CASE REPORT

Male patient, 32 years old, has been presenting bilateral asymptomatic erythematous-scaly plaques on palms and soles for three months, associated with periungual erythema, subungual hyperkeratosis and onychodystrophy of toenails (Figures 1 and 2). He had no previous history of skin disease or other comorbidities and was not using any medication. Direct mycological examination of cutaneous lesions and of all nails involved resulted negative. Biopsy of left palm lesion showed epidermis with

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FIGURE 1: A. Erythematous-scaly plaque on soles. B. Erythematous-scaly plaque on palms



Figure 2: A. Periungual erythema and edema, subungueal hyperkeratosis and onychodystrophy of toenails.

B. infiltrated erythematous-scaly plate, circinated, on the sole and dorsal part of right foot

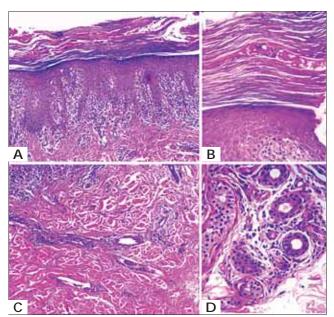


FIGURE 3: A. Psoriasiform hyperplasia and lichenoid infiltrate which darkens the dermal-epidermal interface, a mix of lymphocytes, histiocytes, plasmocytes and neutrophils (H&E, x10). B. Compact hyperkeratosis, parakeratosis and intracorneal neutrophils (H&E, x10). C. Lymphohistiocytic interstitial infiltrate, superficial and deep (H&E, x40). D. Inflammatory infiltrate involving deep annexes (H&E, x40)

compact hyperkeratosis, parakeratosis, intracorneal neutrophils, hypergranulosis, psoriasiform acanthosis, vacuolar alteration/interface squamotization and focal apoptosis. Dermis showed a mixed inflammatory infiltrate of lymphocytes, histiocytes, plasma cells and neutrophils in a subepidermal band, in the interstices of connective tissue and around deep vessels and appendages (Figure 3).

The diagnosis of psoriasiform as well as

superficial and deep interface dermatitis suggested the inclusion of treponematosis among the diagnostic possibilities. The patient returned with the following laboratory results: VDRL 1/256 and anti HIV 1 and 2 antibodies: 1003.37, confirming the diagnosis of secondary syphilis and HIV coinfection. CD4 levels were 185 and viral load 800,000 copies/ml. Treatment with benzathine penicillin 2,400,000 U per week was prescribed for three weeks, with complete remission

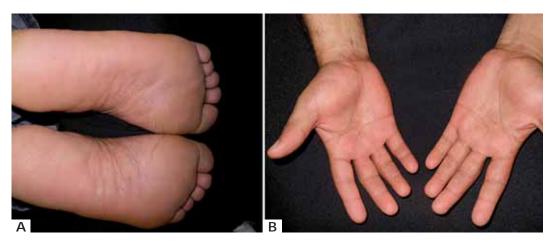


Figure 4: A and B. Clinical aspect of palmoplantar lesions 1 week after completion of treatment with benzathine penicillin





Figure 5: A and B. Clinical aspect of ungual lesions 1 week after completion of treatment with benzathine penicillin

of lesions; the patient was sent to a REFERENCES center for HIV virus treatment (Figures 4 and 5). New nail mycological examination was performed upon his return and the result was negative.

DISCUSSION

According to World Health Organization data, there are approximately 900,000 new cases of syphilis detected annually in Brazil, and this fact reiterates the need for vigilance. Interactions between syphilis and HIV virus arise from the fact that both are transmitted mainly through sexual relations and their importance is increased because ulcerated genital lesions augment the risk of contracting and transmitting the HIV virus; for this reason all patients with syphilis should be offered HIV tests and all HIV-positive patients must be regularly tested for syphilis.²⁶

In HIV-infected patients syphilis does not present opportunistic behavior, but has less usual clinical characteristics, more aggressive and exuberant, with more frequent and early involvement of the nervous system.^{3,4,5,7} Cutaneous lesions are the most common clinical manifestation of secondary syphilis (72%), and may be the only manifestation of

the disease.7

Cutaneous manifestations are so variable that syphilis must be considered in the diagnosis of all dermatoses with atypical presentation. The involvement of palmar and plantar regions is very characteristic and erythematous papules with a scaly collarette are the most common presentation. Sometimes scaling is intense, giving a psoriasiform aspect to lesions.6 Ungual involvement is rarely described in literature and includes nail plate and periungual tissue alterations.8 The negative result of direct mycological examination of nails, the absence of ungual alterations before the onset of cutaneous lesions and excellent improvement of nail plate and periungual tissue by the end of treatment suggested to the authors that these alterations were associated with syphilis.

Due to the several clinical presentations, a biopsy is usually performed and the diagnosis established, as in our case. Histological characteristics of secondary syphilis are very variable and show epidermal and/or dermal changes. In general, the inflammatory reaction acquires a lichenoid (lichen planus type) and/or psoriasiform (psoriasis type) pattern. Findings with

lichenoid psoriasiform aspect, as observed in our case, present a dense mononuclear infiltrate which extends itself to the deep dermis. ⁹ The infiltrate is composed of lymphocytes, macrophages, plasmocytes, neutrophils and eosinophils in variable proportions. Associated vascular alterations include vascular proliferation, prominence of endothelial cells under light, associated with perivascular infiltrate. Specific Wharthin-Starry staining (silver staining) can identify the treponema in the tissue. ⁴

Serological tests in the majority of HIV-infected patients are within the patterns found in non-infected patients. ⁶ However, atypical results may occur. Titration may be too high or too low, and false negativity may occur in some cases. ⁶ Several studies suggest that syphilis, like many other acute infections, leads to transient viral load increases and diminished CD4 cell count, which are resolved after the infection

is treated.¹⁰ It is possible that these transient viral load increases contribute to magnification of HIV transmission risk among patients with syphilis and HIV. ¹⁰ The treatment in the case presented was performed with intramuscular benzathine penicillin 7,200,000UI, in three weekly doses of 2,400,000UI, since it was a case of syphilis of undetermined duration. ²

Syphilis is still a serious public health problem and we must be alert to this diagnosis, especially in case of atypical cutaneous clinical pictures in which they simulate other cutaneous diseases. We present a case of secondary syphilis which clinically simulated psoriasis and dermatophytosis, showing ungual alterations that suggest syphilis involvement, due to the negative result of mycological examination and improvement after treatment. The association with the HIV virus must always be investigated, as in the case presented, due to its association with atypical manifestations of secondary syphilis. \square

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