

Risankizumab for the Treatment of Palmoplantar Pustular Psoriasis: a Report of Two Cases

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Introduction

Palmoplantar pustular psoriasis (PPPP), according to the ERASPEN (European Rare And Severe Psoriasis Expert Network) guidelines, is defined as primary, persistent (>3 months), sterile, macroscopically visible pustular eruption on palms and/or soles [1,2]. It can present with or without concomitant plaque psoriasis [2].

The pathophysiology of this condition is not fully understood, both the innate and adaptive immune systems are involved [2]. A primary role is played by the increased expression of interleukin-8 (IL-8) and by IL-17 related cytokines (including L-17A/F, IL-23A, IL-23 receptor). A role has been proposed also for the antimicrobial peptide LL-37, which may contribute to neutrophil recruitment by upregulating IL-8, IL-23, IL-17C, IL-1 [2]. Only limited data are available regarding the treatment of PPPP with biologics and no standardized guidelines have been published yet [3].

Case Presentation

We present 2 patients with PPPP who were successfully treated with risankizumab, an anti-IL-23 monoclonal antibody.

The first patient is a 32-year-old woman, affected by PPPP and plaque psoriasis since 2020. On clinical examination, we observed pustular lesions on the palms and soles (Figure 1) along with erythematous and scaly plaques on the scalp. Palmoplantar investigator global assessment (ppIGA) score was 3 on a 5-point scale. Psoriasis Area Severity Index (PASI) was 5.2. As topical corticosteroids had previously been ineffective, given the age of the patient, acitretin was not recommended. Because of the impact of the disease on her quality of life (Dermatology life quality index [DLQI] was 12), we prescribed risankizumab 75mg, two subcutaneous injections at weeks 0, 4, and then every 12 weeks.

After 16 weeks the patient returned showing only a slight improvement, but at week 28 we observed complete skin

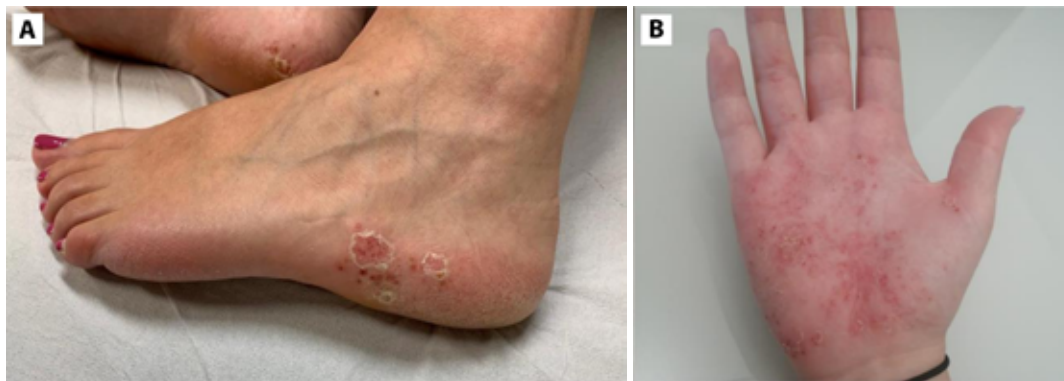


Figure 1. (A,B) Sterile pustules on erythematous skin with moderate scaling on the left foot (A) and on the right palm (B) of a 32-year-old woman, before the start of the therapy.

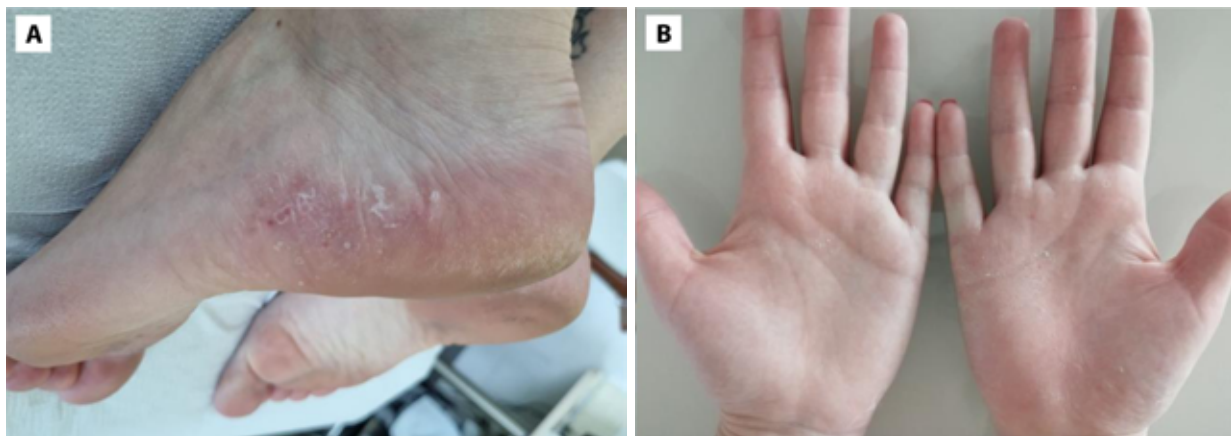


Figure 2. (A,B) Clinical appearance of the patient soles (A) and palms (B) after 28 weeks of therapy with risankizumab. Complete resolution of the pustules is observed, with the persistence of only slight erythema and scaling.

clearance, with a ppIGA of 0 (Figure 2). The patient to date has completed a year of therapy and she has maintained the remission.

The second patient is a 47-year-old woman, affected from PPPP since 2018, previously unsuccessfully treated with acitretin. She had a history of hepatitis B with positive anti-HBc and anti-HBs. On physical examination we observed erythematous, scaly patches on the palms and palmoplantar pustules (ppIGA=3, PASI=11.3). Given her comorbidities and the concomitant presence of plaque psoriasis, with the consensus of the hepatologist, we started risankizumab. At week 16 she achieved complete skin clearance and she has maintained the remission after two years (ppIGA=3). Both patients had provided written consent for retrospective study of data collected during routine clinical practice (demographics, clinical scores).

Conclusions

In scientific literature there is paucity of data regarding the efficacy of biologics in pustular psoriasis, especially for palmoplantar subtype. A few data from real-life experiences

and case reports are available for guselkumab, secukinumab and apremilast, which have demonstrated a moderate efficacy [4]. We decided to prescribe risankizumab for multiple reasons: our favorable experience with this drug, including a patient with a flare of generalized pustular psoriasis [5]; the high safety profile, even in patients with serological evidence of viral hepatitis [6]; the high effectiveness on difficult-to-treat areas [5]. In the first patient, we observed a slower response compared to our experience with risankizumab in psoriasis vulgaris, however this is expectable given the resistance of these areas to treatment [5].

To our knowledge, the experience on the effectiveness of risankizumab in PPPP is limited. Further data, corroborated by longitudinal studies with higher numbers of patients, are needed to assess the role of risankizumab and other biologic drugs for the treatment of PPPP.

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