

Comment on “Folliculitis decalvans: Effectiveness of therapies and prognostic factors in a multicenter series of 60 patients with long-term follow-up”



To the Editor: We read with interest the article entitled “Folliculitis decalvans: Effectiveness of therapies and prognostic factors in a multicenter series of 60 patients with long-term follow-up” by Miguel-Gómez et al.¹ Their article is one of the largest published series of folliculitis decalvans (FD), enrolling 60 patients with a minimum follow-up period of 5 years. The most frequently used treatments were antibiotics (doxycycline/minocycline and the association of rifampicin and clindamycin in 36 [60%] and 21 [35%] patients, respectively). Oral isotretinoin was given to 15 (25%) patients. Although no precise remission or relapse data were reported with the different treatments, Miguel-Gómez et al¹ stated that there was no significant difference between treatment regimen with rifampicin and clindamycin versus isotretinoin by multivariate analysis.

Miguel-Gómez et al¹ did a great job by proposing a practical algorithm for the treatment of FD. In severe cases, the association of rifampicin and clindamycin was suggested as a first-line treatment, and oral isotretinoin is only advised if the response is not maintained with antimicrobials and systemic corticosteroids.

There is, however, increasing evidence supporting oral isotretinoin as a first-line treatment for severe forms of FD.^{2,3} In a series of 49 men with FD who were treated with isotretinoin, Aksoy et al² reported complete and partial response rates of 82% and 10%, respectively. The optimal treatment regimen was yet to be determined, and patients who received oral isotretinoin (≥ 0.4 mg/kg/day) for at least 3 months had the best response, with two-thirds of these patients never experiencing a relapse.

Tietze et al³ retrospectively reviewed the outcome of different treatment options for FD. Full remission was obtained in 9 of 10 patients who were treated with isotretinoin. After stable remission was achieved, treatment was tapered slowly to a minimal dose for several months with good outcomes. In contrast, the long-term remission rate associated with the combination of clindamycin and rifampicin was low (17%).³

Rambhia et al⁴ performed a systematic review of the current published treatment efficacy for FD and found that the level of evidence was low overall, with studies lacking randomization and control

groups.⁵ Therefore, treatment of FD remains mostly empirical. A combination of clindamycin and rifampicin was indeed the most frequently used treatment.⁴

One of the major concerns about prescribing antimicrobials for FD is the emergence of antibiotic resistance, especially after a prolonged course, reaching up to 50% in relapsing disease.³ Therefore, in our opinion, isotretinoin should be considered a solid first-line treatment option of severe forms of FD.

The efficacy of isotretinoin could be explained by its direct inhibitory effect on the immune system.^{2,3} It inhibits the migration of neutrophils into the skin and modulates the innate immune system defenses against Gram-positive bacteria through decreasing the Toll-like receptor 2 level.^{2,3}

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