

Frontal Fibrosing Alopecia Successfully Treated with Intralesional Betamethasone and Oral Hydroxychloroquine: A Case Report

A. Moussa^{1*}, H. Hassan¹, M. El Amraoui¹, Y. Zemmez¹, R. Frikh¹, N. Hjira¹

¹Department of Dermatology, Mohammed V Military Teaching Hospital, Rabat

DOI: <https://doi.org/10.36347/sjmcr.2026.v14i05.018>

Received: 07.03.2026 | Accepted: 21.04.2026 | Published: 06.05.2026

*Corresponding author: A. Moussa

Department of Dermatology, Mohammed V Military Teaching Hospital, Rabat

Abstract

Case Report

Frontal fibrosing alopecia (FFA) is a chronic, lymphocytic, scarring alopecia that affects hair follicles of the frontal scalp, eyebrows, eyelashes, and body hair. We report the case of a 36-year-old woman with progressive FFA resistant to conventional therapies. Clinical examination, trichoscopy, and histopathology confirmed the diagnosis. The patient was treated with a combination of intralesional betamethasone injections and oral hydroxychloroquine. After six months of treatment, a significant clinical improvement was observed, with stabilization of hairline recession and partial eyebrow regrowth. This case highlights the potential benefit of combination therapy in controlling disease progression in FFA.

Keywords: Frontal Fibrosing Alopecia; Combination Therapy; Trichoscopy.

Copyright © 2026 The Author(s): This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International License (CC BY-NC 4.0) which permits unrestricted use, distribution, and reproduction in any medium for non-commercial use provided the original author and source are credited.

INTRODUCTION

Frontal fibrosing alopecia (FFA) is a chronic, lymphocytic, scarring alopecia that affects hair follicles of the frontal scalp, eyebrows, eyelashes, and body hair.

It is closely related to lichen planopilaris (LPP), from which it differs mainly by its distribution on the scalp. First described in the literature in 1994, both conditions have become increasingly common, and genetic, immunological, and environmental pathogenic factors have been proposed [1].

FFA is characterized by a progressive recession of the frontal, temporal, or frontotemporal hairline. The clinical pattern is distinctive and typically includes eyebrow loss, along with other associated features. In addition to frontotemporal hairline recession and eyebrow loss, symptoms such as pruritus, facial papules, eyelash loss, involvement of body hair, and trichodynia may also occur. Flattening of the frontal veins, hypopigmentation along the frontal hairline, and the presence of “lonely hairs” are other frequently reported clinical features [2,3].

Case reports and cohort studies have described clinical improvement and remission with various therapeutic approaches, including 5-alpha-reductase inhibitors, topical and intralesional corticosteroids,

topical and systemic immunomodulators, hydroxychloroquine, isotretinoin, antibiotics, and pioglitazone. Although treatment outcomes remain variable and evidence regarding their efficacy is still limited, combination therapies—particularly those including systemic agents—appear to be beneficial in slowing disease progression and maintaining stabilization [4,5].

We report the case of a patient with frontal fibrosing alopecia who showed a favorable clinical response to intralesional Diprosan® (betamethasone) injections combined with oral hydroxychloroquine.

CASE REPORT

A 36-year-old woman with no significant past medical history presented with a progressive hair loss affecting the frontotemporal scalp and eyebrows, associated with pruritus and scalp and eyebrow tenderness, evolving over one year.

Initial treatment included topical clobetasol propionate (Dermovate® gel) applied once daily, doxycycline 100 mg twice daily, 5% minoxidil, and a Ducray® rubefacient lotion for three months. Despite these treatments, the patient experienced disease progression.

Given the lack of clinical improvement and the impact on her quality of life, the patient was referred to our department. Clinical examination revealed a bilateral and symmetrical recession of the frontal hairline with progressive evolution (Figures 1A and 2A). Eyebrow loss was also noted, predominantly affecting the lateral portions (Figure 3). Trichoscopy showed perifollicular erythema and loss of follicular openings (Figure 4).

The remainder of the clinical examination was unremarkable. A scalp biopsy demonstrated a regular orthokeratotic epidermis, while the underlying dermis

appeared fibrotic with a reduced number of hair follicles (Figure 5). A diagnosis of frontal fibrosing alopecia was established. Laboratory investigations and ophthalmologic examination were within normal limits.

Therapeutically, all previous treatments were discontinued, and intralesional corticosteroid therapy with betamethasone (Diprospan®) was initiated at a dose of 1 mL per monthly session for six months, in combination with oral hydroxychloroquine (200 mg twice daily). A satisfactory clinical improvement was observed (Figures 1B and 2B).



Figure 1: Clinical image of the right frontotemporal scalp.

(A) Before treatment: decreased hair density with recession of the right frontotemporal hairline.

(B) After treatment: stabilization of the right frontotemporal hairline recession, with improved hair density after 6 months of treatment with intralesional Diprospan® (betamethasone) injections combined with oral hydroxychloroquine.



Figure 2: Clinical image of the left frontotemporal scalp

(A) Before treatment: decreased hair density with recession of the left frontotemporal hairline.

(B) After treatment: stabilization of the left frontotemporal hairline recession, with improved hair density after 6 months of treatment with intralesional Diprospan® (betamethasone) injections combined with oral hydroxychloroquine.



Figure 3: Clinical image of the eyebrows.
(A) Before treatment: eye-brow hair loss.
(B) After treatment: eye-brow regrowth



Figure 4: Trichoscopic findings in the patient showing perifollicular erythema associated with loss of follicular openings

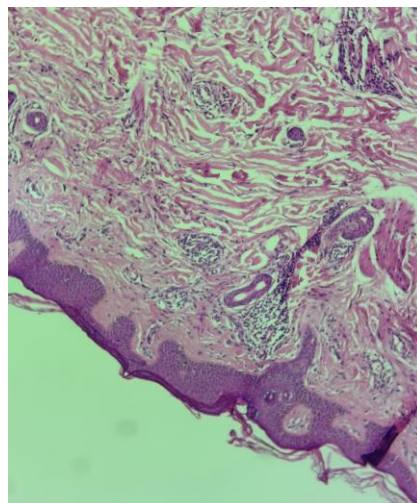


Figure 5. Scalp histopathology: skin biopsy revealed a regular orthokeratotic epidermis, while the underlying dermis appeared fibrotic with a reduced number of hair follicles

DISCUSSION

Frontal fibrosing alopecia (FFA) does not exclusively affect postmenopausal women; it has also been reported in premenopausal women as well as in men [5].

The etiology and pathogenesis of FFA are not yet fully understood and remain the subject of ongoing research. Autoimmune, environmental, hormonal, and genetic factors have all been implicated in its development [6].

Although treatment outcomes are variable and evidence regarding their efficacy remains limited, combination therapies—particularly those including systemic agents—appear to be beneficial in slowing disease progression and maintaining stabilization [7]. Intralesional corticosteroids (ILCs) are often considered a cornerstone of FFA management, as they suppress the inflammatory process, thereby reducing T lymphocyte-mediated destruction of follicular stem cells [8–10].

Hydroxychloroquine is regarded as a first-line systemic therapy in FFA due to its anti-inflammatory and immunomodulatory (anti-lymphocytic) effects, along with a generally favorable safety profile, particularly in cases resistant to topical treatments [11,12].

In our patient, the combination of intralesional corticosteroid therapy and oral hydroxychloroquine resulted in a satisfactory clinical improvement, with a reduction in disease impact on quality of life.

CONCLUSION

In light of these findings, early diagnosis of frontal fibrosing alopecia is essential to enable appropriate management and prevent disease progression.

Trichoscopy represents a valuable tool for both positive diagnosis and differential diagnosis. Combination therapies may help slow disease progression and maintain disease stabilization.

REFERENCES

1. De Souza B, Burns L, Senna MM. Frontal fibrosing alopecia preceding the development of vitiligo : A case report. *JAAD Case Rep.* 2020 Jan 30 ;6(2):154-155. doi: 10.1016/j.jcdr.2019.12.011.
2. Vañó-Galván S, Molina-Ruiz AM, Serrano-Falcón C, et al. Frontal fibrosing alopecia: a multicenter review of 355 patients. *J Am Acad Dermatol.* 2014 ;70(4):670–678. doi: 10.1016/j.jaad.2013.12.003

3. Vañó-Galván S, Rodrigues-Barata AR, Urech M, et al. Depression of the frontal veins: a new clinical sign of frontal fibrosing alopecia. *J Am Acad Dermatol.* 2015 ;72(6):1087–1088. doi: 10.1016/j.jaad.2015.02.1129
4. Imhof RL, Chaudhry HM, Larkin SC, Torgerson RR, Tolkachjov SN. Frontal fibrosing alopecia in women: the mayo clinic experience with 148 patients, 1992–2016. *Mayo Clin Proc.* 2018 ;93(11):1581–1588. doi: 10.1016/j.mayocp.2018.05.036
5. Tan KT, Messenger AG. Frontal fibrosing alopecia: clinical presentations and prognosis. *Br J Dermatol.* 2009 Jan;160(1):75-9. doi: 10.1111/j.1365-2133.2008.08861. x. Epub 2008 Sep 22. PMID : 18811690.
6. Tavakolpour S, Mahmoudi H, Abedini R, Kamyab Hesari K, Kiani A, Daneshpazhooch M. Frontal fibrosing alopecia: An update on the hypothesis of pathogenesis and treatment. *Int J Womens Dermatol.* 2019 Jan 23;5(2):116-123. doi: 10.1016/j.ijwd.2018.11.003. PMID : 30997385 ; PMCID : PMC6451751.
7. Imhof R, Tolkachjov SN. Optimal Management of Frontal Fibrosing Alopecia: A Practical Guide. *Clin Cosmet Investig Dermatol.* 2020 Dec 1 ;13 :897-910. doi: 10.2147/CCID.S235980. PMID : 33293846 ; PMCID : PMC7718862.
8. Gamret AC, Potluri VS, Krishnamurthy K, Fertig RM. Frontal fibrosing alopecia: efficacy of treatment modalities. *Int J Womens Health.* 2019 Apr 29 ;11 :273-285. doi: 10.2147/IJWH.S177308.
9. Donovan JC, Samrao A, Ruben BS, Price VH. Eyebrow regrowth in patients with frontal fibrosing alopecia treated with intralesional triamcinolone acetonide. *Br J Dermatol.* 2010 Nov ;163(5):1142-4. doi: 10.1111/j.1365-2133.2010.09994. x. PMID : 20716217.
10. Martínez-Pérez M, Churruca-Grijelmo M. Frontal Fibrosing Alopecia: An Update on Epidemiology and Treatment. *Actas Dermosifiliogr.* 2015 Nov;106(9):757-8. English, Spanish. doi: 10.1016/j.ad.2014.12.002. Epub 2015 Feb 18. PMID : 25701899.
11. Ho A, Shapiro J. Medical therapy for frontal fibrosing alopecia: A review and clinical approach. *J Am Acad Dermatol.* 2019 Aug ;81(2):568-580. doi: 10.1016/j.jaad.2019.03.079. Epub 2019 Apr 3. PMID : 30953702.
12. Meah N, Li J, Wall D, York K, Bhojru B, Bokhari L, Coulthard L, Asfour L, Abraham LS, Asz-Sigall D. Statement from the frontal fibrosing alopecia international expert alliance : SOFFIA 2024. *J Eur Acad Dermatol Venereol.* 2026 Feb;40(2):210-223. doi: 10.1111/jdv.20833.