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A multicenter descriptive analysis of 270 men with frontal fibrosing alopecia and lichen planopilaris in the United States

Permalink https://escholarship.org/uc/item/4ms4p7pj

Journal Journal of the American Academy of Dermatology, 88(4)

ISSN

0190-9622

Authors

Pathoulas, James T Flanagan, Kelly E Walker, Chloe J <u>et al.</u>

Publication Date

2023-04-01

DOI

10.1016/j.jaad.2022.10.060

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Peer reviewed

| | Treatment delay (n = 79) N (%) | No treatment delay $(n = 70) N (\%)$ | Total (N = 149) N (%) |
|--------------------------------|-----------------------------------|--------------------------------------|--------------------------|
| Age | | | |
| Mean | 64 | 59 | |
| Median (IQR) | 65 (53, 78) | 62 (48, 72) | |
| Range | 28, 95 | 22, 88 | |
| Biological sex | | | |
| Male | 43 (54.4) | 40 (57.1) | 83 (55.7) |
| Female | 36 (45.6) | 30 (42.9) | 66 (44.3) |
| Race | | | |
| White | 60 (75.9) | 44 (62.9) | 104 (69.8) |
| Black/African American | 8 (10.1) | 19 (27.1) | 27 (18.1) |
| Other/unknown/declined | 11 (13.9) | 7 (10.0) | 18 (12.1) |
| Stage | | | |
| IA-IIA | 40 (50.6) | 37 (52.9) | 77 (51.6) |
| IIB | 11 (13.9) | 12 (17.1) | 23 (15.4) |
| IIIA-IIIB | 4 (5.1) | 2 (2.8) | 6 (4.0) |
| IVA1 (SS) | 19 (24.1) | 16 (22.9) | 35 (23.5) |
| IVA2-IVB | 3 (3.8) | 3 (4.3) | 6 (4.0) |
| PC CD30+ LPD | 2 (2.5) | 0 (0.0) | 2 (1.3) |
| COVID-19 status | | | |
| Positive | 18 (22.8) | 10 (14.3) | 28 (18.8) |
| Negative | 61 (77.2) | 60 (85.7) | 121 (81.2) |
| Length of treatment delay (mo) | | | |
| Mean | 3.20 | 0.00 | |
| Median (IQR) | 3.00 [1.05, 4.00] | 0.00 [0.00, 0.00] | |
| Range | 0.30, 10.00 | 0.00, 0.00 | |
| Unknown | 2 | 0 | |
| Outcomes | | | |
| Improvement of disease | 5 (6.3) | 10 (14.3) | 15 (10.1) |
| Stable disease | 15 (19.0) | 36 (51.4) | 51 (34.2) |
| Disease progression/relapse | 52 (65.8) | 22 (31.4) | 74 (49.7) |
| Deceased | 7 (8.9) | 2 (2.9) | 9 (6.0) |

Table I. Demographic and outcomes by treatment delay status

Correspondence to: Larisa J. Geskin, MD, Department of Dermatology, Columbia University, Herbert Irving Pavilion, 161 Fort Washington Ave, 12th Floor, New York, NY 10032

E-mail: ljg2045@cumc.columbia.edu

Conflicts of interest

None disclosed.

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https://doi.org/10.1016/j.jaad.2022.11.016

A multicenter descriptive analysis of 270 men with frontal fibrosing alopecia and lichen planopilaris in the United States

To the Editor: Lichen planopilaris (LPP) and frontal fibrosing alopecia (FFA) are forms of primary lymphocytic cicatricial alopecia characterized by inflammation and fibrosis of the follicular unit causing irreversible hair loss if untreated.¹ Epidemiologic studies of LPP/FFA are lacking but it is thought that the incidence is increasing globally.² Men with LPP/FFA are increasingly reported in the literature, challenging the paradigm that LPP/FFA is a disease of postmenopausal women.^{3,4} This multicenter descriptive study is the first to characterize LPP/FFA among men in the United States.

Medical records from adult men seen at specialty hair clinics in the U.S. between January 2010 and

| Item | All (<i>n</i> = 270) | LPP (<i>n</i> = 215) | FFA (<i>n</i> = 37) | LPP + FFA (n = 18) |
|--|-----------------------|-----------------------|----------------------|--------------------|
| Current mean age \pm SD | 52.54 ± 14.72 | 52.03 ± 14.85 | 56.92 ± 12.79 | 49.66 ± 15.90 |
| Mean age at diagnosis \pm SD | 45.77 ± 14.63 | 45.00 ± 14.60 | 50.68 ± 13.63 | 45.11 ± 15.87 |
| Race, race, ethnicity, or origin, n (%) | | | | |
| American Indian or Alaska Native | 1 (0.37) | 1 (0.47) | 0 (0.00) | 0 (0.00) |
| Asian | 12 (4.44) | 8 (3.72) | 2 (5.41) | 2 (11.11) |
| Black and/or African American | 14 (5.19) | 10 (4.65) | 2 (5.41) | 2 (11.11) |
| Non-White Hispanic, Latino, or Spanish origin | 9 (3.33) | 9 (4.19) | 0 (0.00) | 0 (0.00) |
| Middle Eastern or North African | 3 (1.11) | 3 (1.40) | 0 (0.00) | 0 (0.00) |
| Native Hawaiian or other Pacific Islander | 0 (0.00) | 0 (0.00) | 0 (0.00) | 0 (0.00) |
| White (Hispanic and Non-Hispanic) | 191 (70.74) | 147 (68.37) | 31 (83.78) | 13 (72.22) |
| Other race, ethnicity, or origin | 40 (14.81) | 37 (17.21) | 2 (5.41) | 1 (5.56) |
| History of scalp biopsy consistent with disease, n (%) | | | | |
| Yes, biopsy confirmed | 186 (69.89) | 155 (72.09) | 26 (70.27) | 5 (27.78) |
| No, clinical diagnosis only | 74 (27.41) | 52 (24.19) | 11 (29.73) | 11 (61.11) |
| Unable to determine | 10 (3.70) | 8 (3.27) | 0 (0.00) | 2 (11.11) |

Table I. Demographic and diagnostic history of men with lichen planopilaris, frontal fibrosing alopecia, and both disorders (LPP + FFA)

FFA, Frontal fibrosing alopecia; LPP, lichen planopilaris.

September 2021 were reviewed. Men with a clinical or biopsy-confirmed diagnosis of LPP, FFA, or overlap LPP + FFA were included in the analysis. Patients with a concomitant cicatricial alopecia other than LPP or FFA were excluded (see the Supplement for complete methods).

A total of 270 patients met criteria for inclusion, with 215 having a diagnosis of LPP (79.6%), 37 with FFA (13.7%), and 18 with overlap LPP + FFA (6.7%) (Table I). The average age was 52.5 ± 14.7 years and was similar between groups (P = .11). History of sunscreen application to the face, most days of the week, year-round for at least 5 years was recorded in a minority (14.8%) of charts, and was most prevalent among men with FFA (27.0%) and overlap LPP + FFA (16.7%) compared to men with LPP (6.7%) (Supplementary Table I, available via Mendeley at https://doi.org/10.17632/shw6684cfh.1). Hair loss affecting the beard (21.6%) and sideburns (16.2%) at disease onset was most common among patients with FFA (Fig 1). The frequency of hair loss affecting the arms, chest, and legs at disease onset was 10.4% at each location in men with FFA and less than 1.0% at each location in LPP. Most patients with LPP (61.9%), FFA (54.1%), and overlap LPP + FFA (50.0%) had a history of scalp pruritus (Supplementary Table II, available via Mendeley at https://doi.org/10.17632/shw6684cfh.1). A minority (27.8%) had concomitant androgenetic alopecia. Nearly one-third (29.3%) had been incorrectly diagnosed before seeing a hair specialist (Supplemental Table 3).

This multicenter study of 270 men with LPP/FFA in the United States is, to our knowledge, the largest descriptive study of men with LPP/FFA published to date. A recent meta-analysis reported that use of facial sunscreens increased the likelihood of developing FFA.⁵ We found that long-term consistent use of facial products with sunscreen was more common among men with FFA than those with LPP. However, there was insufficient data to establish causality. Nearly 1 out of 3 men in our cohort had been incorrectly diagnosed before seeing a hair specialist. LPP was commonly misdiagnosed as seborrheic dermatitis. FFA and overlap LPP + FFA were both commonly misdiagnosed as alopecia areata. Loss of beard hair at FFA onset was reported by 1 in 5 patients. Our findings highlight that a history of symptomatic scalp, beard loss, and body hair loss in men presenting with frontal hairline recession should raise suspicion for a cicatricial process. Increased awareness of cicatricial disease in men could promote care-seeking behavior and timely diagnosis.

James T. Pathoulas, MD,^a Kelly E. Flanagan, MD, MS,^a Chloe J. Walker, MD, MHS,^a Maya S. Collins, BS,^a Shaheir Ali, BS,^a Isabel M. Pupo Wiss, BS,^a George Cotsarelis, MD,^b Heather Milbar, MD, Huang, MD,^{c,d} Arash MPH,^b Kathie Mostaghimi, MD,^{c,d} Deborah Scott, MD,^{c,d} Jane J. Han, BS,^c Karen J. Lee, BS,^c Maria K. Hordinsky, MD,^e Ronda S. Farab, MD,^e Gretchen Bellefeuille, BS,^e Ora Raymond, BA,^e Wilma Bergfeld, MD,^f Geraldine Ranasinghe, MD,^f Jerry Shapiro, MD,^g Kristen I. Lo Sicco, MD,^g Daniel Gutierrez, MD,^g Justin Ko, MD,^b Paradi Mirmirani, MD,ⁱ Natasha Mesinkovska, MD, PhD,^j Katerina L. Yale, MD,^j Lynne J. Goldberg, MD,^k Antonella Tosti, MD,^l Eran C.



Fig 1. Perifollicular erythema and loss of facial hair in man with frontal fibrosing alopecia (FFA). This patient reported loss of beard hair at FFA onset.

Gwillim, MD,^l Carolyn Gob, MD,^m and Maryanne M. Senna, MD^{a,d}

From the Department of Dermatology, Massachusetts General Hospital, Boston, Massachusetts^a; Department of Dermatology, University of Pennsylvania, Pennsylvania, Pennsylvania^b; Department of Dermatology, Brigham and Women's Hospital, Boston, Massachusetts^c; Harvard Med*ical School, Boston, Massachusetts^d; Department* of Dermatology, University of Minnesota Medical School, Minneapolis, Minnesota^e; Department of Dermatology, Cleveland Clinic, Cleveland, Obio[†]; Department of Dermatology, NYU Langone Health, New York, New York^g; Department of Dermatology, Stanford University, Palo Alto, *California^b*; *Department of Dermatology, Kaiser* Permanente Vallejo Medical Center, Vallejo, Californiaⁱ; Department of Dermatology, University of California Irvine, Irvine, California^j; Department of Dermatology, Boston University Medical Center, Boston, Massachusetts^k; Department of Dermatology, University of Miami Health System, Miami, Florida¹; and Department of Dermatology, David Geffen School of Medicine at UCLA, Los Angeles, California.^m

Funding sources: None.

- *IRB approval status: Approved by Partners IRB* #2020P002456.
- Patient consent: Consent for the publication of recognizable patient photographs or other identifiable material was obtained by the authors and included at the time of article submission to

the journal stating that all patients gave consent with the understanding that this information may be publicly available.

- Key words: AGA; alopecia; androgenetic alopecia; autoimmune; beard loss; body hair; cicatricial; cicatricial alopecia; dermatology; eyebrows; facial hair; FFA; fibrosing; fibrosis; frontal fibrosing alopecia; hair; hair loss; hair loss in men; hair specialists; itch; lichen planopilaris; loss; LPP; male pattern hair loss; men; misdiagnosis; MPHL; multicenter; pattern hair loss; PHL; pruritus; retrospective review; scalp itch; scarring; scarring alopecia; scalp symptoms; sideburns; specialty hair clinic; sunscreen.
- Correspondence to: Maryanne M. Senna, MD, Department of Dermatology, Massachusetts General Hospital, 50 Staniford St, 2nd Floor, Boston, MA 02114

E-mail: msenna@partners.org

Conflicts of interest

None disclosed.

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https://doi.org/10.1016/j.jaad.2022.10.060

Popular sunscreens marketed to individuals with skin of color: Cost, marketing claims, and allergenic ingredients

To the Editor: There is increasing awareness of the negative effects of ultraviolet (UV) light in individuals with skin of color (SOC), especially in regards to pigmentation disorders induced and/or exacerbated

