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CASE AND RESEARCH LETTER

Linear Alopecia Areata: An Uncommon Clinical Variant. Presentation of Three Cases and Review of the Literature

Alopecia areata lineal: una variante clínica infrecuente. Presentación de 3 casos y revisión de la literatura

To the Editor,

Alopecia areata (AA) is a frequent autoimmune non-scarring hair-loss disorder which can have a significant impact on quality of life. It usually presents with sharply demarcated alopecic patches, although multiple clinical variants of AA have been described. Linear alopecia areata (LAA) is exceedingly rare. ¹⁻⁴ Here we present a series of patients (one adult and two children) with LAA.

The first case was an otherwise healthy 26-year-old male who presented with a history of 8 months of hair loss. Physical examination revealed a 23 cm linear alopecic patch extending from the vertex to the left occipital area (Fig. 1A), following Blaschko's lines (Fig. 1B). The skin was normal in appearance in the alopecic area. Trichoscopy showed black and yellow dots, short vellus hairs and exclamation mark hairs. A diagnosis of LAA was made. Topical 5% minoxidil, mini-pulse therapy with prednisone (20 mg on two consecutive days weekly) and intralesional triamcinolone acetonide injections every six weeks were started. In the following 4 months, a mild response was observed, and topical 1% anthralin lotion was initiated with partial response after 3 months. The patient was lost to follow-up.

Our second case, a 9-year-old girl with no previous comorbidities presented with scalp hair loss during the previous 3 months. On physical examination a linear alopecic patch was observed along the interparietal area (Fig. 1C). The pull test was positive. Trichoscopy showed black dots, pigtail hairs and exclamation mark hairs (Fig. 1D). No erythema, atypical vessels or perifollicular scaling were seen. These findings were consistent with LAA. Oral prednisone 15 mg/day was prescribed with progression of the alopecia during the following month. Dexamethasone 0.1 mg/kg twice a week was then initiated, with complete response

after 6 months. Treatment was then stopped. No recurrences have been seen after 2 years of follow-up.

The third case, an otherwise healthy 5-year-old girl presented with hair loss during the previous month. Physical examination showed a linear alopecic patch with normal skin appearance on the frontal and left parietal area (Fig. 1E). The hair pull test was positive. Trichoscopy revealed yellow and black dots and exclamation mark hairs. A diagnosis of LAA was made. Topical 0.05% Clobetasol foam was prescribed and complete response was observed during the following three weeks, and topical corticosteroid was then suspended. The patient was lost to follow-up.

The clinical presentation of AA can be highly variable, even annular or rectangular-pattern variants have been described. Only four cases (two adults and two children) presenting with LAA have been reported (Table 1). Hall While the gold standard in the diagnosis of hair-loss disorders is the histopathology, in real-life practice the diagnosis of AA is generally based on clinical and trichoscopic findings, and histologic study is only performed in challenging cases. Trichoscopy of AA is characterized by the presence of black and yellow dots, short vellus hairs, pigtail (circle) hairs, upright regrowing hairs, tapered hairs, Pohl-Pinkus constrictions and exclamation mark hairs.

Differential diagnoses of alopecia with linear distribution include linear epidermal nevi, linear scleroderma, linear lichen planus, linear folliculitis decalvans, linear cutaneous lupus erythematosus (LCLE),⁶ LAA and trichotillomania.³ An accurate diagnosis is important since management and clinical implications can be different. Trichoscopy can be very useful in this context: in LCLE it may reveal thick or prominent arborizing blood vessels, brown or blue-grey speckled pigmentation and black dots; in linear scleroderma, pili torti, broken hairs, and black dots on a whitish skin surface and thick telangiectatic vessels; and in trichotillomania, broken hair, black dots, flame hair, and V-hair.³

As we all know, the prognosis of AA is extremely variable and difficult to predict. We have found no reports suggesting that LAA can have different prognosis or outcomes.

We have presented a series of three patients with LAA and highlighted the importance of trichoscopy in the differential diagnosis of linear-pattern hair loss. Dermatologists should be aware of this in order to avoid performing invasive diagnostic procedures.

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+Model AD-3626; No. of Pages 4

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J. Piquero-Casals, D. Morgado-Carrasco, D. Saceda-Corralo et al.

Table 1 Cases with linear alopecia areata described in the literature.

Author/year	Age (years)/ sex	Disease evolution (months)	Location	Trichoscopy	Comorbidities/ family history of autoimmune disorders	Histology	Treatment	Clinical response
Shetty et al./2016 ¹	35/M	36	Vertex and bilateral temporal areas	Short vellus hair in centre and hairs with the tapering base at the periphery	No	Lymphocytic infiltrate in peribulbar area with an increase in telogen hair units. Negative staining for mucin	Monthly intralesional triamcinolone (5 mg/ml) injections and topical minoxidil 5% solution	NR
Yu et al./2018 ²	28/M	48	Interparietal	Short vellus hairs	No	Increased catagen and telogen follicles. Negative staining for mucin	Monthly intramuscular betamethasone and topical minoxidil 5% solution	NR
Mukherjee et al./2018 ³	7/F	1	Bilateral temporal area and vertex	Broken hair, hair dust, flame hair, and V-hair	No	"Swarm of bees" appearance	Topical 2% minoxidil solution and mometasone ointment	NR
Albaghsi et al./2022 ⁴	12/M	1	Right parieto- temporo- occipital	Diffuse yellow and black dots and a few exclamation mark hairs	Sickle cell disease	Dermis with a focal, moderate, superficial, perivascular, lymphocytic infiltrate, with scattered mast cells and rare eosinophils. Hair follicles thin and short, with their bulbs extending only to the mid-dermis (miniaturized/vellus hairs)	NP	Spontaneous healing after 6 months
Current report	26/M	8	Right parietal to left occipital area	Black dots, short vellus hairs, tapered hairs, broken hairs and exclamation mark hairs	No	NP	Topical 5% minoxidil solution, prednisone micro pulses two consecutive days weekly and intralesional triamcinolone every six weeks Topical 1% anthralin lotion	Mild response Partial response

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ACTAS Dermo-Sifiliográficas xxx (xxxx) xxx-xxx

Table 1 (Continued)

Author/year	Age (years)/ sex	Disease evolution (months)	Location	Trichoscopy	Comorbidities/ family history of autoimmune disorders	Histology	Treatment	Clinical response
Current report	9/F	3	Interparietal	Black dots and exclamation mark hairs	No	NP	Dexamethasone pulses 0.1 mg/kg twice weekly	Complete response after 6 months
Current report	5/F	1	Left parietal area	Yellow dots, black dots, broken hairs and exclamation mark hairs	No	NP	Topical 0.05% Clobetasol foam twice daily for three weeks	Complete response

Abbreviations: M, male; F, female; NP, not performed; NR, not reported.



Figure 1 Linear alopecia areata. (A) Linear alopecic patch extending from the vertex to the left occipital area, following Blashko's lines. (B) Blashko's lines on the head. (C) Linear alopecic patch along the interparietal area. (D) Trichoscopy shows black dots, pigtail hairs and exclamation mark hairs (FotoFinder® medicam 1000, Fotofinder systems, Germany). (E) Linear alopecic patch on the frontal and left parietal area.

+Model AD-3626; No. of Pages 4

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J. Piquero-Casals, D. Morgado-Carrasco, D. Saceda-Corralo et al.

Conflict of interest

The authors declare that they have no conflict of interest.

References

- 1. Shetty S, Rao R, Kudva RR, Subramanian K. Linear alopecia areata. Int J Trichology. 2016;8:144–5.
- 2. Yu L, Lu Z. Linear alopecia areata. JAAD Case Rep. 2018;4:1072-3.
- 3. Mukherjee SS, Chandrashekar BS. Linear alopecia areata versus trichotillomania: the game of time. Indian J Paediatr Dermatol. 2018;19:136.
- **4.** Albaghsi H, AlSaba'a S, Almustafa ZZ. Blaschko-linear alopecia areata in a child, an exceedingly rare variant. Int J Dermatol. 2022;61:e403-5.
- 5. Shin J, Jang HS, Cho SB. Rectangular-patterned occipital alopecia areata: a report of three cases. Int J Trichology. 2012;4:164–6.

- Rhee C-H, Kim S-M, Kim MH, Cinn YW, Ihm C-W. Two cases of linear alopecia on the occipital scalp. Ann Dermatol. 2009;21:159–63.
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