# SHORT COMMUNICATION

## Rolled Hairs in Two Cases of Naevus Comedonicus of the Scalp

Umi TAHARA<sup>1</sup>, Sachiko ISHIHARA<sup>2</sup>, Satomi AOKI<sup>1</sup>, Manabu OHYAMA<sup>3</sup> and Akiharu KUBO<sup>1\*</sup>

<sup>1</sup>Department of Dermatology, Keio University School of Medicine, 35 Shinanomachi, Shinjuku-ku, Tokyo 160-8582, <sup>2</sup>Department of Dermatology, National Hospital Organization Tokyo Medical Center, and <sup>3</sup>Department of Dermatology, Kyorin University Faculty of Medicine, Tokyo, Japan. \*E-mail: akiharu@keio.jp

Accepted Sep 2, 2021; Epub ahead of print Sep 7, 2021

Spirally twisted rolled hairs have been reported in association with follicular hyperkeratotic conditions, including keratosis pilaris and ichthyosis (1). Rolled hairs appear mostly on the extensor extremities and trunk, but have not been reported on the scalp (1). Naevus comedonicus (NC) is a rare naevus caused by somatic *NEK9* mutations, which shows multiple comedo formation within the affected skin (2). We report here 2 cases of NC of the scalp presenting with rolled hairs, detected by trichoscopy.

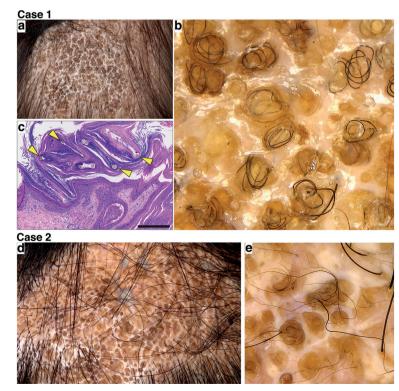
### **CASE REPORTS**

*Case 1*. Case 1 was a 1-year-old Japanese girl with a congenital naevus on the parietal region of the scalp. The patient was otherwise healthy and had an unremarkable family history. The naevus presented a cluster of hyperkeratotic, brownish papules with alopecia, showing a cobblestone appearance (**Fig. 1**a). Trichoscopic examination revealed heavily rolled hairs within papules (Fig. 1b).

Skin biopsy of the lesion revealed rolled hairs embedded in follicular plugs and a dilated hair follicle infundibulum (Fig. 1c).

*Case 2.* Case 2 was a 25-year-old Japanese man with multiple grouped comedo-like pits, cyst formation, and scarring over his face, neck, trunk, and limbs since childhood. Notably, skin lesions on the limbs were distributed along Blaschko's lines. His family history was unremarkable. Physical examination of the scalp revealed a well-demarcated hypotrichotic lesion with a cluster of hyperkeratotic, brownish papules in the parietal region, similar to case 1 (Fig. 1d). The lesional scalp was atrophic and fibrotic, probably due to extensive inflammation during adolescence. Trichoscopic examination revealed rolled hairs within papules, but the detection rate was lower than that observed in case 1 (Fig. 1e).

After obtaining written informed consent and approval from the Institutional Review Board, genetic analyses of genomic DNA isolated from peripheral blood and the lesional epidermis and dermis, which were separated by dispase treatment, were performed (3). Somatic heterozygous mutations in *NEK9* were identified; c.1718T>C (p.1573T) in case 1 and c.1755\_1757delAAC (p.T586del) in case 2, in an epidermis-specific manner (**Fig. 2**). The somatic mutations of *NEK9* identified in the current cases have



**Fig. 1. Clinical, trichoscopic, and histopathological features of the patients.** The features of case 1 and case 2 are shown in upper panels (a–c) and lower panels (d–e), respectively. (a, d) A cluster of hyperkeratotic, brownish papules with alopecia on the scalps. (b, e) Trichoscopic examination of the patient's scalp showing rolled hairs within papules. (c) Rolled hairs (*yellow arrowheads*) embedded in follicular plugs with a dilated hair follicle infundibulum (haematoxylin-eosin; scale bars, 250 µm).

been previously reported in NC (2, 4). The diagnosis of NC was therefore confirmed in both cases.

## DISCUSSION

NC is an uncommon malformation of the pilosebaceous unit characterized by dilated follicular orifices filled with follicular plugs. In NC, massive cystic inflammation and fibrosis sometimes occur during adolescence (5), as happened in case 2.

NC is most commonly located on the face. neck, trunk and arms. While several cases of NC have been reported to develop on the scalp, trichoscopic findings were described for only one adult patient with follicular plugs and vellus hair (6). To the best of our knowledge, this is the first report describing a unique trichoscopic finding of NC of the scalp. Both case 1 and case 2 exhibited rolled hairs of the scalp embedded within follicular plugs. The waxy follicular plugs caused by abnormal keratinization were speculated to have hindered the emergence of hair, which resulted in the spiral transverse hair growth within the semilucent structures, probably via slower growth of follicular plugs than hairs (7). Most of the lesional hairs were rolled hairs in case 1, which could be associated

This is an open access article under the CC BY-NC license. www.medicaljournals.se/acta Society for Publication of Acta Dermato-Venereologica

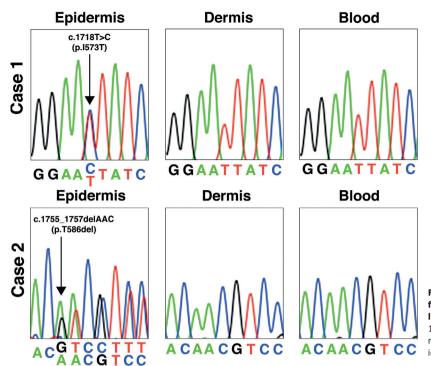


Fig. 2. Analysis of NEK9 in genomic DNA isolated from the biopsied skin and peripheral blood leukocytes. Sanger sequencing chromatograms of cases 1 and 2 are shown in upper panels and lower panels, respectively. Arrows indicate the mutations of NEK9 identified in an epidermis-specific manner.

with the softness of babyhood hairs. The lesional hairs of case 1 were considered not hard enough to penetrate the follicular plugs.

It is noteworthy that differential diagnosis of NC of the scalp includes naevus sebaceous, within which various neoplasms can occur (8). Cerebriform patterns and vellowish globules in cobblestone patterns are unique trichoscopic features of naevus sebaceous in adulthood and childhood, respectively (9). Our observation suggested that rolled hairs within follicular plugs are potential diagnostic clues for NC of the scalp, which can be useful for differentiation from naevus sebaceous. Further accumulation of cases with NC of the scalp would enable the establishment of its trichoscopic features.

#### ACKNOWLEDGEMENTS

The patients in this manuscript have given written informed consent to publish their case details.

Conflicts of interest. M. Ohyama is a scientific advisor for Eli Lilly Japan, Pfizer Japan Inc., Taisho Pharmaceutical Co., and RHOTO Pharmaceutical Co. and received research grants not related to this study from Shiseido Co. The other authors have no conflicts of interest to declare.

## REFERENCES

- 1. Panchaprateep R, Tanus A, Tosti A. Clinical, dermoscopic, and histopathologic features of body hair disorders. J Am Acad Dermatol 2015: 72: 890-900.
- 2. Levinsohn JL, Sugarman JL, Yale Center for Mendelian Genomics, McNiff JM, Antava RJ, Choate KA, Somatic mutations in NEK9 cause nevus comedonicus. Am J Hum Genet 2016; 98: 1030-1037.
- 3. Kitano Y, Okada N. Separation of the epidermal sheet by dispase. Br J Dermatol 1983; 108: 555-560.
- 4. Sheppard SE, Smith A, Grand K, Pogoriler J, Rubin AI, Schindewolf E, et al. Further delineation of the phenotypic spectrum of nevus comedonicus syndrome to include congenital pulmonary airway malformation of the lung and aneurysm. Am J Med Genet A 2020; 182: 746-754.
- 5. Ferrari B, Taliercio V, Restrepo P, Luna P, Abad ME, Larralde M. Nevus comedonicus: a case series. Pediatr Dermatol 2015; 32: 216-219.
- 6. Mathachan SR, Arora P, Ahuja A, Gautam RK. Late onset nevus comedonicus of scalp: a rare site for an uncommon condition. Dermatol Online J 2019; 25: 13.
- 7. Durmaz EÖ, Demirkesen C. Circle hair: report of two cases and brief review of the literature. Dermatol Online J 2019; 25: 10.
- 8. Cribier B, Scrivener Y, Grosshans E. Tumors arising in nevus sebaceus: a study of 596 cases. J Am Acad Dermatol 2000; 42: 263-268.
- 9. Ankad BS, Beergouder SL, Domble V. Trichoscopy: the best auxiliary tool in the evaluation of nevus sebaceous. Int J Trichology 2016; 8: 5-10.