CASE REPORT

Late onset Ito’s nevus

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SUMMARY
Dermal melanocytes include a variety of congenital and acquired melanocytic lesions characterised by the presence of multiple spindle-shaped dendritic melanocytes in the dermis. These lesions are commonly found in the skin of Asians, but they can also appear in Caucasians. The Mongolian spot, nevi of Ota and Ito are the most common morphological forms. We report a case of a 24-year-old Caucasian woman presented with a 10-months history of progressive darkening of the right side of her upper back. Cutaneous examination revealed a macular blue-grey hyperpigmentation of the right side of her upper back. Biopsy specimen from the macule showed multiple darkly pigmented, spindle-shaped dendritic melanocytes in the superficial dermis, interstitially arranged between collagen bundles. The diagnosis of nevus of Ito was therefore established.

BACKGROUND
Dermal melanocytosis include a variety of congenital and acquired melanocytic lesions characterised by the presence of multiple spindle-shaped dendritic melanocytes in the dermis, migrating from the neural crest to the epidermis.1–3 These lesions are typically associated with Asians, but can also appear in Caucasians patients.4–5 Because of their similar histology, differentiating dermal melanocytic lesions is most often supported on clinical features and anatomic distribution.1–6

Dermal melanocytosis results in a brown or blue pigmentation, depending whether melanin is mainly in the upper or lower dermis, respectively.7 The Mongolian spot, nevi of Ota and Ito are the most common morphological forms of dermal melanocytosis.6 Nevus of Ito is often unilateral and clinically similar to nevus of Ota,6 except in anatomic location, which corresponds to the distribution of the posterior supravacular and cutaneous brachial nerves.6–7

We report this case to salient the rarity of acquired Ito’s nevus, which appeared in a Caucasian adult and the importance of adequate follow-up.

CASE PRESENTATION
A 24-year-old Caucasian woman presented with a 10-month history of progressive darkening of the right side of her upper back. The patient denied any prior inflammation, significant sun exposure or trauma to the area. She also denied using any medications or metals that could possibly alter skin pigmentation. Her medical history was unremarkable and there was no family history of cutaneous diseases.

Cutaneous examination revealed a macular blue-grey hyperpigmentation of the right side of her upper back and shoulder (figure 1). There was no associated hypertrichosis. Examinations of the rest of the skin, mucous membranes, hair, nails and the remainder of the physical examination were normal.

Biopsy specimen from the macule showed multiple darkly pigmented, spindle-shaped dendritic melanocytes in the superficial dermis, interstitially arranged between collagen bundles (figure 2). The diagnosis of nevus of Ito was therefore established.

OUTCOME AND FOLLOW-UP
Our patient refused to undergo any therapy. She is maintaining vigilance in dermatology consultation.

DISCUSSION
Nevus of Ito occurs most frequently in Asian populations and the true incidence is unknown.4 The majority of cases are diagnosed in early infancy and in early adolescence.4–9 Few cases of acquired dermal melanocytosis after puberty have been reported.2–7 A late-onset Ito’s nevus may be seen as in our patient.

There are several hypotheses to explain the pathogenesis of acquired dermal melanocytosis.1–7 One hypothesis is that dermal melanocytes migrating from the neural crest during the embryological development fail to reach their location in the basal layer of epidermis. In addition, dermal melanocytes may migrate from the basal layer of the epidermis or from hair bulb to dermis.1–7–10 Other hypothesis is that reactivation of pre-existing, latent dermal melanocytosis, may be activated by inflammation, trauma, pregnancy, hormone replacement therapy.
or some unknown aging stimuli. Genetic factors are also important in the pathogenesis of dermal melanocytosis and a positive family history has been reported in literature.

After the onset, nevus of Ito may progressively enlarge and darken in colour and its appearance usually remains stable once adulthood is reached. All dermal melanocytosis are generally considered benign, although rare cases of malignant transformation have been described in literature. Although nevus of Ito is much more prevalent in Asian populations, it is generally recognised that the risk of malignant transformation appears to be much more common in Caucasian individuals. Consequently, vigilance may be necessary, particularly, when these lesions occur in Caucasian individuals. Skin biopsies after the diagnosis are necessary only if clinical changes or malignant transformation are suspected within the involved skin, ocular or mucosal tissues.

Pulsed Q-switched laser surgery is the current treatment for nevus of Ito and it works via selective photothermal and photomechanical destruction of dermal melanocytes and melanosomes. High success rates have been described with the Q-switched ruby, Q-switched alexandrite and Q-switched Nd:YAG lasers. After 4–8 treatments, skin pigmentation is reduced or removed in almost 100% of cases, with a less than 1% risk of scarring. Second-line therapies like dermabrasion or cosmetic camouflage can also be used. Other treatment modalities that have been tried without any success are chemical peels, carbon dioxide laser and topical bleaching. In the present case, the patient is maintaining vigilance in dermatology consultation and she refused to undergo any therapy.

Learning points

- Nevus of Ito occurs most frequently in Asian populations, but it can also occur in Caucasian patients.
- Nevus of Ito is usually diagnosed in early life, but it can be diagnosed only in the adult life.
- There are several hypotheses to explain the pathogenesis of acquired dermal melanocytosis.

Contributors CR contributed with the bibliographic research, the diagnosis and treatment of the patient and the elaboration of the article. CA contributed with the bibliographic research. APV helped in the diagnosis and treatment of the patient. CB also contributed to the bibliographic research.

Competing interests None.

Provenance and peer review Not commissioned; externally peer reviewed.

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