Onychomatricoma on the fourth toenail: A rare tumor in a rare localization

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**Abstract**

Onychomatricoma is a rare benign tumor of the nail matrix, characterized by finger-like projections that invade the nail plate. The fingernails of Caucasian women are most commonly affected. Because this tumor can easily mimic other more prevalent ungual diseases, it is crucial to be acquainted with its characteristic clinical and histopathologic features. The authors present a case of a 40-year-old man with an onychomatrichoma in the fourth left toenail, which was initially misdiagnosed and treated as onychomycosis.

**Key words:** Nail disorders, nail tumors, onychomatrichoma, onychomycosis

**Introduction**

Onychomatricoma is a rare and benign tumor of the nail matrix. It predominantly affects the fingernails of Caucasian women during the fifth decade of life and typically is slow-growing and painless. Although onychomatricoma has a unique set of clinical and histopathologic features, its rarity and the lack of knowledge about this disease contribute to a frequent mistaken and late diagnosis. Herein, we report a case of a 40-year-old man with an onychomatricoma in an unusual localization, the fourth left toenail which was initially misdiagnosed and treated as onychomycosis.

**Case Report**

A 40-year-old Caucasian man presented to our outpatient department with a two-year history of asymptomatic discoloration and thickening of the fourth left toenail. He denied any previous local trauma. For the clinical suspicion of onychomycosis, he was treated by his primary care physician with a three-month regimen of oral terbinafine, and topical amorolfine for eight months, without improvement.

Dermatologic examination revealed a hypertrophic and dystrophic nail with longitudinal ridging, transverse and longitudinal over-curvature and brown-yellowish discoloration [Figures 1a and b]. Nail plate dermoscopy demonstrated white longitudinal grooves. All the other toenails and fingernails were unaffected.

The ventral view of the extracted nail plate evidenced multiple parallel cavities [Figure 2b], fitting perfectly in the tumor surface. The tumor was completely excised [Figure 2c], and histopathological examination demonstrated a fibroepithelial neoplasm with filiform projections [Figure 3], corroborating onychomatricoma’s clinical diagnosis. Periodic acid-Schiff histochemical staining was negative for fungal organisms.

**Discussion**

Since the first description of onychomatricoma by Baran and Kint in 1992, fewer than 100 cases have been reported. Although etiology remains unknown, nail plate trauma and onychomycosis have been identified as possible predisposing factors of this tumor.3

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Fingers are more commonly affected than toes with a prevalence of 64%. When the onychomatricoma reaches toenails, most cases are found in the big toe and sometimes in the second or the third toe. In 2015, Di Chiacchio et al. reported a large series of 30 onychomatricoma, where 11 cases were found on the toenails: seven in the big toenail, two in the second toe and two in the third toe. In 2016, Coutellier et al. were the first to describe an onychomatricoma on the fifth toe of the left foot in a 68-year-old female. We were unable to find any previous reports of onychomatricoma on the fourth toenail.

The classic tetrad signs of onychomatricoma include thickening of the nail plate, transverse or longitudinal over-curvature, xanthonychia and multiple splinter hemorrhages. Other possible presentations are nail dystrophy, erythronychia, melanonychia, plate fracture, paronychia and dorsal pterygium. The clinical diagnosis of onychomatricoma can be challenging because it can easily mimic many other nail tumors and diseases, including the most prevalent ungual disorder—onychomycosis. Consequently, onychomatricoma should be considered in the differential diagnosis of patients with onychomycosis-like changes failing to respond to proper antifungal treatment, as in this case. The index of suspicion should increase when a single dystrophic nail is involved. Notably, cases of onychomatricoma with concomitant onychomycosis can also occur, as the channels within the nail plate created in onychomatricoma provide a favorable environment for fungal invasion.

Onychoscopy, nail clipping, ultrasound and confocal microscopy may aid in onychomatricoma’s diagnosis. Dermatoscopy criteria include longitudinal parallel white lines, parallel lesion edges, splinter hemorrhages, dark dots, nail pitting and thickening of the free edge. However, the gold standard diagnostic method is surgical biopsy with histopathological examination. Intraoperatively, a filamentous tumor with digitiform projections that penetrate the nail plate leading to woodworm-like cavities is usually found. Histopathological evaluation characteristically exhibits a fibroepithelial tumor composed of a proximal pedunculated base and a distal zone with multiple epithelial digitations. A biphasic fibrous stroma, superficially cellular and deeply collagenized and relatively hypocellular, is present throughout the tumor.

The treatment of choice for onychomatricoma is complete surgical excision and is typically curative. Secondary ungual dystrophy and recurrences have been rarely reported.

**Conclusion**

Onychomatricoma should be kept in mind in the differential diagnosis of nail apparatus changes to avoid misdiagnosis and proper treatment delay. The current case illustrates the
importance of suspecting an underlying onychomatricoma toward a single long-standing distorted nail unresponsive to fungicidal treatment. It well highlights onychomatricoma’s described clinical and histopathological features and is of particular interest due to its unusual localization on the fourth toenail.

Declaration of patient consent
The authors certify that they have obtained all appropriate patient consent.

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Conflicts of interest
There are no conflicts of interest.

References