Neonate born to hepatitis B carrier mother presenting with congenital onychodysplasia of the index finger (Iso–Kikuchi syndrome)

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To The Editor,

Congenital onychodysplasia of the index fingers, also known as Iso–Kikuchi syndrome, is characterized using five diagnostic criteria: congenital occurrence, unilateral or bilateral involvement of the index fingers, variation in nail appearance, possible hereditary involvement and frequently associated bone abnormalities1–5. Herein, we present the case of a newborn with congenital onychodysplasia of the left index finger.

A 2,480 g male neonate was born to a hepatitis B carrier mother at 38 weeks gestation by cesarean section. Hepatitis B immune globulin in combination with the recombinant hepatitis B vaccine was given at the time of birth. Two additional doses of the recombinant hepatitis B vaccine were scheduled over the first 6 months to prevent vertical transmission of hepatitis B virus (HBV). At birth, physical examination showed micronychia on the ulnar aspect of the left index finger (Fig. 1b). No limitation in clasping the left hand or spreading out the palm of the left hand was observed. A lateral X-ray showed a Y-shaped bifurcation of the distal phalanx of the left index finger (Fig. 1a). A diagnosis of congenital onychodysplasia of the index finger (COIF) was confirmed. No other family member had similar features. The parents were unrelated.

COIF is characterized by the full spectrum of nail dysplasia, including anonychia, micronychia and polyonychia (rudimentary splitting), hemi-onychocryptosis, irregular lunula and nail misalignment1–5. Both sexes are affected equally, with international incidence of 4.2 cases per 100,000 live births5. COIF, which is usually congenital, can be transmitted in an autosomal or recessive dominant inheritance pattern, although sporadic cases are described and are regarded as de novo mutations1,5. Our newborn filled four of the five criteria, with no reported positive family history.

The exact pathogenesis remains unknown. In utero, ischemia of the palmar artery and a dysplastic change in the crescent-shaped cap of the distal phalanx have been proposed as the two main candidate pathogenetic mechanisms1–3,5. Exposure to teratogens during pregnancy, especially to antidepressants, antiepileptics and abortifacient drugs, and to insulin have been described in the literature as causes5. The mother never took any medication that might have influenced the onset of COIF during pregnancy.

Skin and nail abnormalities are frequently associated with liver disease6. Although HBV-associated COIF has not yet been reported in the literature, HBV infection might cause COIF, because various nail abnormalities have been associated with HBV infection6. Transplacental transmission of HBV might occur, especially if mothers are positive for both hepatitis B surface antigen (HBsAg) and hepatitis B envelope antigen (HBeAg) and have high viral loads7. In our case, transplacental transmission was unlikely because of the HBeAg-negative HBsAg carrier mother.

The absence of an index fingernail rarely affects a patient’s ability to perform functions. COIF is of cosmetic importance, but its prognosis is excellent. Identification of the syndrome and reassurance of the patient and parents are all that is necessary.

Key words: congenital onychodysplasia of the index fingers, hepatitis B virus, Iso–Kikuchi syndrome, nail dysplasia.
REFERENCES


Fig. 1. a. Radiograph shows Y-shaped bifurcation on lateral view on the left index digit, marked by an arrow. b: Micronychia on the ulnar aspect of the left index finger.