Case Report

Extadigits: an unusual presentation

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INTRODUCTION

Polydactyly means the presence of more than the normal number of fingers or toes. It can vary from unnoticeable rudimentary finger or toe to fully develop extra digit, which occur as an isolated congenital anomaly or as one aspect of multi-symptom disease or syndrome.¹ Several syndromes were reported to be associated with polydactyly and geneticist often play roles in examination of children with this deformity for other congenital anomalies.²

The presentation of polydactyly can be inform of thumb polydactyly.³ Macroductyly of the left second toe with duplication of the nail beds,⁴ postaxial duplication of the fifth digit,⁵ preaxial duplication of the great toe,⁶⁷ Post-axial polydactyly in both hands which is very rare, an extra digit within the hand and not along its borders and along with one foot is even rarer.⁹ Studies across different population indicate variation in frequency of polydactyly, with 75% seen as Hand post axial, 15% as foot post axial polydactyly whip is less often found with other anomalies (7% versus 15%) and 10% as bilateral hand and foot post axial polydactyly.⁵ The incidence was reported to be 2.3 per 1000 in white males, 0.6 per 1000 in white females and 13.5 in black males and 11.1 in black females.⁶

Here we report a very rare case of polydactyly with two extra digits on the left hand and one on the right hand, right foot and left foot. The extra digits were well developed, with normal range of motion, good capillary refill and intact sensation. The treatment modality used was surgical removal of the extra digits and reconstruction of any associated anomalies in the remaining ray such as longitudinal epiphyseal bracket. After the surgery the patients is no longer experience difficulty with fitting gloves and shoes as well as discrimination among peer groups in his future life.

CASE REPORT

The patient is a 5 months old male infant who presented with accessory digits of the both hands and feet, with two extra digits on the left hand and one on the right hand, right foot and left foot. The extra digits were well developed, with normal range of motion, good capillary refill and intact sensation.

Keywords: Extra digits, Malformation, Surgery, Motion
Patient is a product of full term gestation, index pregnancy was booked at three months and there was no history of radiation exposure or use of unprescribed medications, no history of maternal febrile illness during pregnancy. Delivery was uneventful via spontaneous vertex delivery.

The patient’s mother is para4 +0, 4 alive there is no family history of extra digits or other congenital anomalies.

The patient has no other congenital malformations.

A general examination revealed an otherwise healthy child and a musculoskeletal system examination showed that he had two additional digits on the left hand with 1 digit on both feet and right hand respectively (Figure 1-4). The extra digits were well developed, with normal range of movement, with good capillary refill and intact sensation. There were no other anomalies detected.

An assessment of polydactyly of both hands and feet was made. The patient had two stage excisions of the extra digits, with excision of the extra digits of both feet and hands at the first stage and second stage respectively and there were no post-operative complications.

DISCUSSION

The management of polydactyly may appear simple, but careful consideration before and during surgical correction are needed. Several presentations were reported in the literature. The close presentation of case of polydactyly in relation to our case report documented in the literature were functional extra digit in both feet and non-functional extra digit in both hand, six digits in each limb with non functional extra digit in the hands, bilateral postaxial (one extra digit on each limb) functional polydactyly of both hands and feet. Polydactyly was also demonstrated as one of the presentation of autosomal recessive and dominant condition such as Acrocallosal syndrome, post axial polydactyly-progressive myopia syndrome, cleft lip and cleft palate. Variability in genetic expression. Other associated anomaly of bilateral polydactyly of hands and feet associated was supernumerary renal vessels in right kidney. Abnormalities involving polydactyly are usually bilateral, although, few studies revealed unilateral involvement being more common than bilateral involvement. In our case report the rarest form of presentation was seen in which two additional digits on the left hand (Figure 1) was observed making the total number of digit to be 25, and normal range of motion, good capillary refill and intact sensation was observed in the extradigit. A general examination of patient revealed an otherwise healthy child with no associated congenital anomalies.

The treatment modality of polydactyly in our patient involved surgical removal of the extra digits (Figure 5-8)
and reconstruction of any associated anomalies in the remaining ray such as longitudinal epiphyseal bracket. This in agreement with the basic goal for surgical management of patients with polydactyly which is removal of the most medial or most lateral digit so as to gain the normal contour of the hand as well as maintaining the maximum functional and cosmetic outcome.\(^{19-25}\)

Generally, it is desirable to treat polydactyly of the hands and toes because untreated patients experience difficulty with fitting gloves and shoes with discrimination among peer groups. These reasons probably explain why the parents of our index patient opted for surgical excision of the digits.

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