

Congenital Hand Anomalies - An Overview



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Melbourne, 2014

Presentation outline

- Incidence
- Embryology
- Classification of Congenital Hand Anomalies
- Splinting
- Surgery

Incidence

- 1 in 600 live births
- When 3 or more minor anomalies exist there is 90% chance of a major anomaly in one of the critical organ systems
 - Eg. cardiac, pulmonary, gastrointestinal, genitourinary & haematological
 - Usually in combinations identified as syndromes eg. VACTERL syndrome

Embryology

Congenital hand anomalies may result from:

- Chromosomal abnormalities
- Inborn errors of metabolism
- Infections or other environmental factors
- Other unknown influences

Classification of Congenital Hand Anomalies

- I. Failure of formation of parts (arrest of development)
- II. Failure of differentiation (separation) of parts
- III. Duplication
- IV. Overgrowth
- V. Undergrowth
- VI. Congenital constriction ring syndrome
- VII. Generalised “skeletal” abnormalities & syndromes

I. Failure of formation of parts



1

Radial deficiency

1. Radial dysplasia (radial club hand)
2. Hypoplastic thumb



2



3

Central deficiency (involves rays 2,3 & 4)

3. Symbrachydactyly
4. Ectrodactyly



4



5

5. **Ulnar deficiency** (Little & ring finger may be absent +/- ulna & carpal bones)

II. Failure of Differentiation of Parts

Soft Tissue involvement

1. Arthrogryposis
2. Syndactyly
3. Camptodactyly
4. Clasp thumb
5. Trigger thumb

2



1



3



5



Skeletal involvement

6. Clinodactyly

6



4



III. Duplication

Digit

1. Polydactyly



2. Duplicate thumb



IV. Overgrowth

Digits

1. Macrodactyly



V. Undergrowth

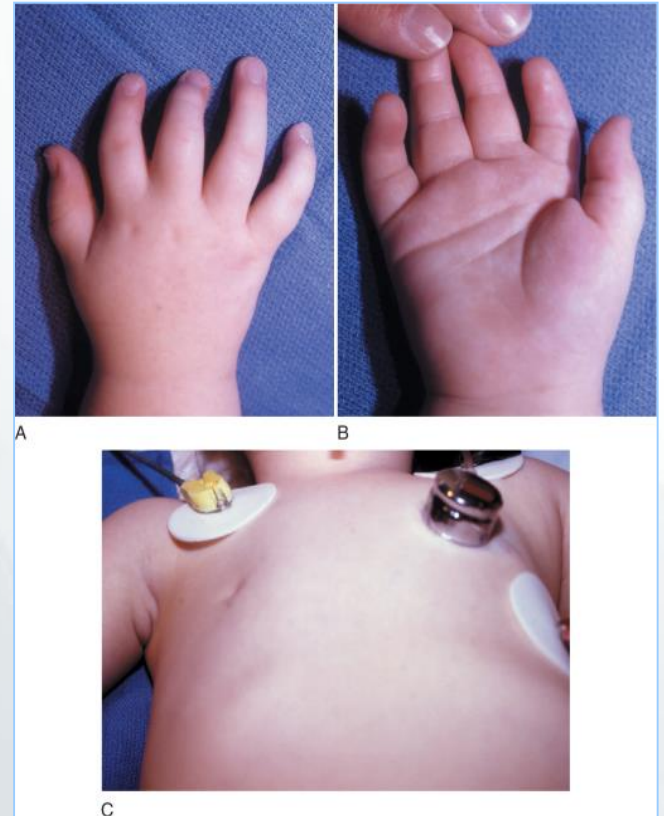
Digits

1. Brachysyndactyly
2. Poland syndrome

1



2



VI. Constriction Ring Syndrome

- Amniotic bands



VII. Generalised Skeletal Anomalies

- Chromosomal & others

General Splinting for Congenital Hand Anomalies

Goals of treatment:

- Maximise range of motion
- Maximise functional hand use and independence in occupational performance
- Minimise secondary consequences
- Protection of structures post surgery



Splinting for Congenital Contractures

Timing and Protocol

- As early as possible after birth - Most successful correction before 4-6months
- Rigid thermoplastic splinting
- Ideal: >8-12 hours daily
- Frequent serial adjustment
- When resolved: splint overnight to maintain
- Consider developmental implications

In older children & adolescents:

- More time is required to serially correct
- Some residual contracture is more likely to remain
- Psychosocial issues have greater impact on compliance

Splinting for Contracture Prevention

Timing and Protocol

- From neonate
- Overnight, rest periods
- Rigid thermoplastic splinting if required
- Neoprene, lycra or elastic may be sufficient
- Functional splints may also be required

Splinting for Function

Timing and Protocol

- From neonate
- During functional hand use
- Rigid thermoplastic splinting if stability required
- Neoprene, lycra or elastic if dynamic assistance required
- Wear during functional tasks
- Not required overnight / at rest
- Consider developmental implications

Surgery for Congenital Hand Anomalies

With **early surgery**

- Joints are more amenable to remoulding
- Child will develop & cortically imprint fewer bad habits
- Plasticity of the central nervous system (motor & sensory cortex) is greater

With **delayed surgery**

- Task of surgery is technically easier
- Child's functional needs are more obvious
- Child is potentially more co-operative

RCH OT Post-surgery protocols

- Hypoplastic thumb reconstruction
- Pollicisation
- Duplicate thumb reconstruction
- Toe-to-hand transfer
- Syndactyly release

*At RCH most procedures of this type will be performed on children between the ages of six months & two years



Protocols available for post-surgical management from RCH

References of interest

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With thanks to Tanya Cole and Josie Duncan