DISEASE:
Syndactyly type 4

NAME:	Syndactyly type 4
DESCRIPTION:	A rare non-syndromic syndactyly characterized by complete bilateral cutaneous fusion of all fingers, frequently associated with polydactyly (usually involving six digits and six metacarpals). Phalanges may fuse as a conglomerate mass of bones. Feet are occasionally affected.
ORPHACODE:	93405
SYNONYMS:	Polysyndactyly, Haas type
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	SHH LMBR1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Brain malformations (gene panel)

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB

Related Analytes

- limb development membrane protein 1
- sonic hedgehog signaling molecule