

[Open Peer Review on Qeios](#)

Aphalangy-syndactyly-microcephaly syndrome

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base. Aphalangy-syndactyly-microcephaly syndrome. ORPHA:1113*

Aphalangy-syndactyly-microcephaly is an extremely rare malformation syndrome characterized by the association of partial distal aphalangia with syndactyly, duplication of metatarsal IV, microcephaly, and mild intellectual disability.