

Open Peer Review on Qeios

Craniodigital-intellectual disability syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Craniodigital-intellectual disability syndrome</u>. ORPHA:1514

Craniodigital syndrome - intellectual deficit is characterised by syndactyly of the fingers and toes, characteristic facies (`startled' facial expression with a small pointed nose, micrognathia, long dark eyelashes and prominent eyebrows) and intellectual deficit.

Qeios ID: LJKFU4 · https://doi.org/10.32388/LJKFU4