Open Peer Review on Qeios

# Craniomicromelic syndrome

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Craniomicromelic syndrome. ORPHA:1524*

Craniomicromelic syndrome is a very rare disorder characterized by intrauterine growth retardation, underossification of the skull with large fontanels, short limbs with absent phalanges and finger and toe syndactyly.