

Open Peer Review on Qeios

## Acrodysostosis

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Acrodysostosis. ORPHA:950

Acrodysostosis (ACRDYS) is a rare primary bone dysplasia characterized by severe brachydactyly, peripheral dysostosis with facial dysostosis, nasal hypoplasia, and developmental delay.

Qeios ID: 04KW1A · https://doi.org/10.32388/04KW1A