Acrocephalosyndactyly

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Acrocephalosyndactyly. ORPHA:946

Acrocephalosyndactyly (ACS) syndromes represent a group of inherited congenital malformation disorders characterized by craniosynostosis and fusion or webbing of the fingers or toes, often with other associated manifestations.