Ulnar/fibula ray defect-brachydactyly syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Ulnar/fibula ray defect-brachydactyly syndrome. ORPHA:52056

Ulnar/fibula ray defect - brachydactyly syndrome is a very rare malformation syndrome characterized by ulnar hypoplasia associated with hypoplastic to absent fourth and/or fifth digits, fibular hypoplasia, short stature and facial dysmorphism.