Tel Hashomer camptodactyly syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Tel Hashomer camptodactyly syndrome. ORPHA:3292

Tel Hashomer camptodactyly syndrome is a rare syndrome characterized by camptodactyly, muscle hypoplasia and weakness, skeletal anomalies, facial dysmorphism and abnormal dermatoglyphics.