Camptodactyly syndrome, Guadalajara type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Camptodactyly syndrome, Guadalajara type 1. ORPHA:1327

Camptodactyly syndrome, Guadalajara type 1 is a rare syndrome consisting of growth retardation, facial dysmorphism, camptodactyly and skeletal anomalies.