Larsen syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Larsen syndrome. ORPHA:503

Larsen syndrome (LS) is a rare skeletal dysplasia characterized by congenital dislocation of large joints, foot deformities, cervical spine dysplasia, scoliosis, spatula-shaped distal phalanges and distinctive craniofacial abnormalities, including cleft palate.