Telecanthus-hypertelorism-strabismus-pes cavus syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Telecanthus-hypertelorism-strabismus-pes cavus syndrome. ORPHA:3293

Telecanthus-hypertelorism-strabismus-pes cavus syndrome is characterized by telecanthus, hypertelorism, strabismus, pes cavus and other variable anomalies. It has been described in a father and his son. The son also had hypospadias, bilateral inguinal hernia, clinodactyly and camptodactyly of the fingers, and radiographic findings including flared metaphyses of the long bones and osteopenia.