Jackson-Weiss syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Jackson-Weiss syndrome. ORPHA:1540

Jackson-Weiss syndrome (JWS) is a rare genetic disorder characterized by foot malformations (tarsal and metatarsal fusions; short, broad, medially deviated great toes) and in some patients craniosynostosis with facial anomalies. Hands are normal in affected patients.