Weill-Marchesani syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Weill-Marchesani syndrome. ORPHA:3449

Weill-Marchesani syndrome (WMS) is a rare condition characterized by short stature, brachydactyly, joint stiffness, and characteristic eye abnormalities including microspherophakia, ectopia of the lens, severe myopia, and glaucoma.