Williams syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Williams syndrome. ORPHA:904

Williams syndrome is a rare genetic multisystemic neurodevelopmental disorder characterized by a distinct facial appearance, cardiac anomalies (most frequently supravalvular aortic stenosis), cognitive and developmental abnormalities, and connective tissue abnormalities (such as joint laxity)