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PHAVER syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. PHAVER syndrome. ORPHA:2876

Phaver syndrome is a very rare syndrome characterized by the association of limb Pterygia, Heart anomalies, Autosomal recessive inheritance, Vertebral defects, Ear anomalies and Radial defects.