Stickler syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Stickler syndrome. ORPHA:828

Stickler syndrome is an inherited vitreoretinopathy characterized by the association of ocular signs with more or less complete forms of Pierre-Robin sequence (see this term), bone disorders, and sensorineural deafness (10% of cases).