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# Wagner disease

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Wagner disease. ORPHA:898*

Wagner disease is a rare hereditary vitreoretinopathy characterized by an anomalous vitreous associated with myopia, cataract, chorioretinal atrophy, and peripheral tractional or rhegmatogenous retinal detachment.