Aicardi syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Aicardi syndrome. ORPHA:50

Aicardi syndrome is a rare neurodevelopmental disorder defined by the triad of agenesis of the corpus callosum (total or partial), typical chorioretinal lacunae and infantile spasms that affect almost exclusively females.