Aicardi-Goutières syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Aicardi-Goutières syndrome. ORPHA:51

Aicardi-Goutières syndrome (AGS) is an inherited, subacute encephalopathy characterised by the association of basal ganglia calcification, leukodystrophy and cerebrospinal fluid (CSF) lymphocytosis.