Alpers-Huttenlocher syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Alpers-Huttenlocher syndrome. ORPHA:726

Alpers Huttenlocher syndrome (AHS) is a cerebrohepatopathy and a rare and severe form of mitochondrial DNA (mtDNA) depletion syndrome characterized by the triad of progressive developmental regression, intractable seizures, and hepatic failure.