ADan amyloidosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. ADan amyloidosis. ORPHA:97346

ADan amyloidosis is a rare, neurodegenerative disease characterized by progressive cataracts, hearing loss, cerebellar ataxia, paranoid psychosis and dementia. Neuropathological features are diffuse atrophy of all parts of the brain, chronic diffuse encephalopathy and the presence of extremely thin and almost completely demyelinated cranial nerves.