Infantile neuronal ceroid lipofuscinosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Infantile neuronal ceroid lipofuscinosis. ORPHA:79263

Infantile neuronal ceroid lipofuscinosis (INCL) is a form of neuronal ceroid lipofuscinosis (NCL; see this term) characterized by onset during the second half of the first year of life and rapid mental and motor deterioration leading to loss of all psychomotor abilities.