Alpha-N-acetylgalactosaminidase deficiency type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Alpha-N-acetylgalactosaminidase deficiency type 1. ORPHA:79279

Alpha-N-acetylgalactosaminidase (NAGA) deficiency type 1 is a very rare and severe type of NAGA deficiency (see this term) characterized by infantile neuroaxonal dystrophy.