Alpha-N-acetylgalactosaminidase deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Alpha-N-acetylgalactosaminidase deficiency. ORPHA:3137

Alpha-N-acetylgalactosaminidase (NAGA) deficiency is a very rare lysosomal storage disease that is clinically and pathologically heterogeneous and is characterized by deficient NAGA activity.