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# Alpha-N-acetylgalactosaminidase deficiency type 2

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Alpha-N-acetylgalactosaminidase deficiency type 2. ORPHA:79280*

Alpha-N-acetylgalactosaminidase (NAGA) deficiency type 2 is a very rare mild adult type of NAGA deficiency (see this term) with the features of angiokeratoma corporis diffusum (see this term) and mild sensory neuropathy.