Aspartylglucosaminuria

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Aspartylglucosaminuria. ORPHA:93

Aspartylglycosaminuria (AGU) is an autosomal recessive lysosomal storage disease belonging to the oligosaccharidosis group (also called glycoproteinosis).