

Open Peer Review on Qeios

## Glycogen storage disease due to glycogen branching enzyme deficiency

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Glycogen</u> storage disease due to glycogen branching enzyme deficiency. ORPHA:367

Glycogen branching enzyme (GBE) deficiency (Andersen's disease or amylopectinosis), or glycogen storage disease type 4 (GSD4), is a rare and severe form of glycogen storage disease which accounts for approximately 3% of all the glycogen storage diseases (see these terms).

Qeios ID: VHL78D · https://doi.org/10.32388/VHL78D