

Open Peer Review on Qeios

## Alpha-mannosidosis

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Alphamannosidosis</u>. ORPHA:61

Alpha-mannosidosis is an inherited lysosomal storage disorder characterized by immune deficiency, facial and skeletal abnormalities, hearing impairment, and intellectual deficit.

Qeios ID: P4UBCW · https://doi.org/10.32388/P4UBCW