

[Open Peer Review on Qeios](#)

# Mucopolysaccharidosis type 7

INSERM

## Source

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

*Mucopolysaccharidosis type 7. ORPHA:584*

Mucopolysaccharidosis type VII (MPS VII) is a very rare lysosomal storage disease belonging to the group of mucopolysaccharidoses.