

Open Peer Review on Qeios

## Angelman syndrome

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Angelman syndrome</u>. ORPHA:72

Angelman syndrome (AS) is a neurogenetic disorder characterized by severe intellectual deficit and distinct facial dysmorphic features.

Qeios ID: LJYH5V · https://doi.org/10.32388/LJYH5V