

Open Peer Review on Qeios

## Cohen syndrome

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Cohen syndrome</u>. ORPHA:193

Cohen syndrome (CS) is a rare genetic developmental disorder characterized by microcephaly, characteristic facial features, hypotonia, non-progressive intellectual deficit, myopia and retinal dystrophy, neutropenia and truncal obesity.

Qeios ID: CG8EJ2 · https://doi.org/10.32388/CG8EJ2