

Open Peer Review on Qeios

## Smith-Lemli-Opitz syndrome

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Smith-Lemli-Opitz syndrome</u>. ORPHA:818

Smith-Lemli-Opitz syndrome (SLOS) is characterized by multiple congenital anomalies, intellectual deficit, and behavioral problems.

Qeios ID: XIZ1BQ · https://doi.org/10.32388/XIZ1BQ