

Open Peer Review on Qeios

## Infantile glycine encephalopathy

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Infantile</u> <u>glycine encephalopathy</u>. ORPHA:289860

Infantile glycine encephalopathy is a mild to severe form of glycine encephalopathy (GE; see this term), characterized by early hypotonia, developmental delay and seizures.

Qeios ID: C7HXWR · https://doi.org/10.32388/C7HXWR