

Open Peer Review on Qeios

Hydrolethalus

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hydrolethalus. ORPHA:2189

Hydrolethalus (HLS) is a severe fetal malformation syndrome characterized by craniofacial dysmorphic features, central nervous system, cardiac, respiratory tract and limb abnormalities.

Qeios ID: E4UQIR · https://doi.org/10.32388/E4UQIR