

Open Peer Review on Qeios

Fountain syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Fountain</u> <u>syndrome</u>. ORPHA:3219

Fountain syndrome is an extremely rare multi-systemic genetic disorder characterized by intellectual disability, deafness, skeletal abnormalities and coarse facial features.

Qeios ID: KQS1LY · https://doi.org/10.32388/KQS1LY