

Open Peer Review on Qeios

## Hall-Riggs syndrome

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Hall-Riggs</u> <u>syndrome</u>. ORPHA:2107

Hall-Riggs syndrome is a very rare syndrome consisting of microcephaly with facial dysmorphism, spondylometaepiphyseal dysplasia and severe intellectual deficit.

Qeios ID: TPSL1D · https://doi.org/10.32388/TPSL1D