

Open Peer Review on Qeios

## 17p13.3 microduplication syndrome

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>17p13.3</u> microduplication syndrome. ORPHA:217385

17p13.3 microduplication syndrome is characterized by variable psychomotor delay and dysmorphic features.

Qeios ID: PMS0YM · https://doi.org/10.32388/PMS0YM