

Open Peer Review on Qeios

## Microduplication Xp11.22p11.23 syndrome

INSERM

## Source

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

<u>Microduplication Xp11.22p11.23 syndrome</u>. ORPHA:217377

Familial and de novo recurrent Xp11.22-p11.23 microduplication has been recently identified in males and females.

Qeios ID: 4VDE91 · https://doi.org/10.32388/4VDE91