Microduplication Xp11.22p11.23 syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Microduplication Xp11.22p11.23 syndrome. ORPHA:217377

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