10q22.3q23.3 microduplication syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. 10q22.3q23.3 microduplication syndrome. ORPHA:276422

10q22.3q23.3 microduplication syndrome is a rare, chromosomal anomaly characterized by variable clinical features that may include developmental delay, mild intellectual disability and dysmorphic facial features. In some cases, microcephaly, growth retardation and congenital heart defects have been reported.