Paternal uniparental disomy of chromosome 6

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Paternal uniparental disomy of chromosome 6. ORPHA:96191

Paternal uniparental disomy of chromosome 6 is an uniparental disomy of paternal origin characterized by intrauterine growth retardation, transient neonatal diabetes mellitus, and macroglossia.