Microcephaly-glomerulonephritis-marfanoid habitus syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Microcephaly-glomerulonephritis-marfanoid habitus syndrome. ORPHA:2172

This syndrome is characterised by intellectual deficit, marfanoid habitus, microcephaly, and glomerulonephritis.