Rubinstein-Taybi syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Rubinstein-Taybi syndrome. ORPHA:783

Rubinstein-Taybi syndrome is a rare malformation syndrome characterized by congenital anomalies (microcephaly, specific facial characteristics, broad thumbs and halluces and postnatal growth retardation), short stature, intellectual disability and behavioural characteristics.