Moyamoya angiopathy-short stature-facial dysmorphism-hypergonadotropic hypogonadism syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Moyamoya angiopathy-short stature-facial dysmorphism-hypergonadotropic hypogonadism syndrome. ORPHA:280679

Moyamoya angiopathy - short stature - facial dysmorphism - hypergonadotropic hypogonadism is a very rare, hereditary, neurological, dysmorphic syndrome characterized by moyamoya disease, short stature of postnatal onset, and stereotyped facial dysmorphism.