Lathosterolosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Lathosterolosis. ORPHA:46059

Lathosterolosis is an extremely rare inborn error of sterol biosynthesis characterized by facial dysmorphism, congenital anomalies (including limb and kidney anomalies), failure to thrive, developmental delay and liver disease.