

Open Peer Review on Qeios

3M syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>3M</u> syndrome. ORPHA:2616

3M syndrome is a primordial growth disorder characterized by low birth weight, reduced birth length, severe postnatal growth restriction, a spectrum of minor anomalies (including facial dysmorphism) and normal intelligence.

Qeios ID: V56ZOV · https://doi.org/10.32388/V56ZOV