DPM3-CDG

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. DPM3-CDG. ORPHA:263494

DPM3-CDG is an extremely rare form of CDG syndrome (see this term) characterized clinically in the single reported case by muscle weakness, waddling gait and dilated cardiomyopathy (see this term).