Methylmalonic acidemia with homocystinuria, type cblD

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Methylmalonic acidemia with homocystinuria, type cblD. ORPHA:79283

cblD type methylmalonic acidemia with homocystinuria is a form of methylmalonic acidemia with homocystinuria (see this term), an inborn error of vitamin B12 (cobalamin) metabolism characterized by variable biochemical, neurological and hematological manifestations.