Homocystinuria-Megaloblastic Anemia, cblE Complementation Type

National Cancer Institute

Source

National Cancer Institute. Homocystinuria-Megaloblastic Anemia, cblE Complementation Type. NCI Thesaurus. Code C142173.

An autosomal recessive condition caused by mutation(s) in the MTRR gene, encoding methionine synthase reductase. It is characterized by homocystinuria and megaloblastic anemia.