

VITAMIN B12 METABOLIC DEFECT WITH METHYLMALONICACIDEMIA AND HOMOCYSTINURIA (MUTASE) REFERENCES

(COMBINED DEFICIENCY OF METHYLMALONYL CoA MUTASE AND HOMOCYSTEINE: METHYLTETRAHYDROFOLATE METHYLTRANSFERASE; *cb1C*; VITAMIN B12 METABOLIC DEFECT, TYPE 2; METHYLMALONICACIDEMIA AND HOMOCYSTINURIA; *cb1D*; VITAMIN B12 LYSOSOMAL RELEASE DEFECT; COBALAMIN, DEFECT IN LYSOSOMAL RELEASE OF VITAMIN B12 STORAGE DISEASE; COBALAMIN F DISEASE; *cb1F*; METHYLMALONICACIDURIA DUE TO VITAMIN B12-RELEASE DEFECT)

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