

Disease

Combined methylmalonic acidemia and homocystinuria, cblX type^{1, 2}

Gene

HCFC1

Inheritance	XL
Current Treatments	cobalamin, carnitine & diet
Evidence for Treatment	expert opinion
Non-Molecular Testing Availability	yes
Non-Molecular Test	serum amino acids, urine organic acids ³
Disease Frequency (per 100,000)	1.5 ⁴
Genes Included in Disease Frequency	MMUT, MMAA, MMAB, MMADHC, MMACHC, MCEE, PRDX1, HCFC1, THAP11, ZNF143, LMBRD1, ABCD4, CD320
Experimental / Investigational Therapies	https://clinicaltrials.gov/ct2/results?cond=Methylmalonic+Acidemia&term=&cntry=&state=&city=&dist=

References

- ¹ <https://www.ncbi.nlm.nih.gov/books/NBK1328/>
- ² <https://pubmed.ncbi.nlm.nih.gov/36207831/>
- ³ <https://www.ncbi.nlm.nih.gov/books/NBK1328/#cbl.Diagnosis>
- ⁴ <https://ghr.nlm.nih.gov/condition/methylmalonic-acidemia#statistics>