

Disease

Succinyl-CoA:3-ketoacid CoA transferase (SCOT) deficiency^{1, 2}

Gene

OXCT1

Inheritance	AR
Current Treatments	IV glucose during acute episodes, avoid prolonged fasting and excessive fat intake, mildly restrict protein intake, carnitine supplements
Evidence for Treatment	expert opinion
Non-Molecular Testing Availability	no
Experimental / Investigational Therapies	https://clinicaltrials.gov/ct2/results?cond=OXCT1&term=&cntry=&state=&city=&dist=

References

¹ <https://omim.org/entry/245050>

² <https://www.ncbi.nlm.nih.gov/pubmed/30799594>