

Disease

SLC25A32 associated riboflavin transporter deficiency^{1, 2}

Gene

SLC25A32

Inheritance	AR
Current Treatments	riboflavin
Evidence for Treatment	expert opinion
Non-Molecular Testing Availability	yes
Non-Molecular Test	plasma acylcarnitine profile, urine organic acids ¹
Experimental / Investigational Therapies	https://clinicaltrials.gov/search?cond=SLC25A32

References

¹ <https://pubmed.ncbi.nlm.nih.gov/34764427/>

² <https://omim.org/entry/616839>