

Laboratory Investigation Report

Patient Name	Centre	n Hospital
Age/Gender	OP/IP No/UHID	
MaxID/Lab ID	Collection Date/Time	
Ref Doctor	Reporting Date/Time	

Test Name	Result	Unit	Bio Ref Interval
Outsourced			
SIN No: B2B4308198			

Biotinidase Deficiency (Quantitative) Heparin (L)*

Biotinidase Enzyme Assay	4.03	nmol/min/ml
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Comment Ref. Range:

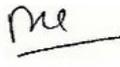
Normal >5.0
Deficient <5.0

Biotinidase deficiency is autosomal recessive disorder. Its incidence is > 1:75,000. Major clinical features include Alopecia, periorificial skin rash, conjunctivitis, developmental delay, hypotonia. Other than low blood biotinidase activity, urine 3-OH-isovaleric acid, 3-methylcrotonylglycine and blood C5-OH-carnitine may be elevated.

Kindly correlate with clinical findings

*** End Of Report ***


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