MAX Lab

DEEP HOSPITAL
LAMP OF LIFE

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	

Outsourced

SIN No:B2B4308198

Test Name	Result	Unit	Bio Ref Interval
Biotinidase Deficiency (Quantitative) Heparin (L)*			
Biotinidase Enzyme Assay	**4.03**	nmol/min/ml	

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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The authenticity of the report can be verified by scanning the Q R Code on top of the page

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