

### 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
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#### Biotinidase Deficiency (Quantitative) Heparin (L)\*

Biotinidase	4.03	nmol/min/ml
Enzyme Assay		

#### Comment Ref. Range:

Normal >5.0

Deficient <5.0

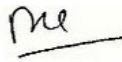
Biotinidase deficiency is autosomal recessive disorder. Its incidence is > 1:75,000. Major clinical features include Alopecia, periorificial skin rash, conjunctivitis, developmental delay, hypotonia. Other than low blood biotinidase activity, urine 3-OH-isovaleric acid, 3-methylcrotonylglycine and blood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

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*



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