

**TEST REPORT**

Patient's Name : **Rahul M Parikh**  
Referred by : C/o. Dynatech Systems  
Date : 21/05/2024 15:15  
Mo. :

Ref. No. : 452  
Age : 28 Years  
Sex : Male

**LIPID PROFILE**

Test Name	Result	Units	Biological Reference Interval
Serum Cholesterol :	157.02	mg / dl	Desirable level/low risk : < 200 Borderline level/moderate risk : 200-239 Elevated level/ high risk : > 240
Serum Triglyceride :	93.29	mg / dl	Normal : <150 Borderline high :150-400 High : 400-1000
S. HDL Cholesterol :	41.01	mg / dl	Desirable level/low risk : >80 Borderline level/moderate risk : 35-80 Elevated level/ high risk : <35
S. LDL Cholesterol :	18.658	mg / dl	Desirable level/low risk : <130 Borderline level/moderate risk : 130-159 Elevated level/ high risk : >160
S. VLDL Cholesterol :	18.658	mg / dl	Upto 34
Total Lipids :	528.636	mg / dl	400 - 700
Chol./HDL Ratio :	3.8		
LDL/HDL Ratio :	0.5		Desirable level/low risk : 0.5-3.0 Borderline level/moderate risk : 3.0-6.0 Elevated level/ high risk : >6.0

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**Dr. M.P. Patel**  
M.D. Pathologist



5, Patel Chember, Near Scope CT Scan, Opp. HDFC Bank, Mansa Mo.: 8866364 222  
108, Shri Shakti Complex, Pethapur Cross To Pethapur Village Road, Pethapur, Gandhinagar Mo.: 63527 78001

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**ELECTROLYTES**

Test Name	Result	Units	Biological Reference Interval
Serum Sodium :	139.87	mmol /L	[137 - 145]
Serum Potassium :	3.9	mmol/L	[3.5 - 5.1]
Serum Chlorides :	102.32	mmol/L	[98 - 107]

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**FASTING BLOOD SUGAR**

Test Name	Result	Units	Biological Reference Interval
Fasting blood sugar :	102.40	mg/dl	Non Diabetic: 70 - 109 Impaired : 110 - 125 Diabetic: 126

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S.G.P.T

Test Name	Result	Units	Biological Reference Interval
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S.G.P.T :	24.44	U/L	0 - 40
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**ERYTHROCYTE SEDIMENTATION RATE (ESR)**

Test Name	Result	Units	Biological Reference Interval
After 1 hour :	6.00	mm	1 - 7

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**CREATININE**

Test Name	Result	Units	Biological Reference Interval
Serum creatinine :	0.88	mg/dl	0.4 - 1.5
Estimated GFR MALE :	109.599067785396		ml/min/1.73m [ > 60 ]
Estimated GFR FEMALE:	81.3225082967639		ml/min/1.73m [ > 60 ]

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**HEMOGRAM**

Test Name	Result	Units	Biological Reference Interval
Hemoglobin :	13.9	g/dl	[13.0-18.0]
Total RBC Count :	5.0	mill/cmm	[4.7-6.0]
Total WBC Count :	5400	/cmm	[4000-10000]
Platelet Count :	217000	/cmm	150000-450000
<b><u>Blood Indices</u></b>			
P.C.V. :	40.8	%	[42-52]
M.C.V. :	81.6	femtolitre	[78-100]
M.C.H. :	27.8	pg	[27-31]
M.C.H.C. :	34.1	g/dl	[32-36]
R.D.W. :	16.0	%	[11.5-14.0]
<b><u>Differential WBC Count</u></b>			
	<b><u>Expected value</u></b>	<b><u>Absolute Count</u></b>	<b><u>Expected value</u></b>
Polymorphs : <b>44</b>	[60 - 70]%	2376	1600 - 7000
Lymphocytes : <b>48</b>	[20 - 40 ]%	2592	800 - 4000
Eosinophils : 01	[1 - 4 ]%	54	00-450
Monocytes : 07	[2 - 6 ]%	378	200 - 1000
Basophils : 00	[0 - 1 ]%	0	0 - 100

**Smear Study**
 RBCs : RBCs are normocytic,normochromic.  
 WBCs : **WBCs series shows relative lymphocytosis.**  
 Platelets : Platelets are adequate in number.  
 PS For MP : No any parasites seen.


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### TSH

Test Name	Result	Units	Biological Reference Interval
Serum TSH :	1.53	μIU/ml	[0.34-5.60]

Note :

1. In patients receiving high dose Biotin therapy (>5mg/day) the specimen should be collected for at least 8 hours after the last biotin administration.
2. Sodium heparin therapy interferes with this assay hence sampling from these patients is not recommended.
3. Rarely high titers of antibodies to Streptavidin and Rubidium may also interfere with the assay.

Comments:

TSH Receptor stimulating antibodies are most closely associated with disease pathogenesis in all forms of Autoimmune thyrotoxicosis (Graves disease), Hashitoxicosis & Neonatal Thyroloxisosis. These antibodies may be detected before autoimmune thyrotoxicosis becomes biochemically or clinically manifest. Since treatments for Graves disease are not aimed at underlying disease process but deal with ablation of thyroid tissue, these antibodies may persist even after apparent clinical. This is specially relevant in pregnant women with Graves disease treated with thyroid ablative therapy who continue to produce thyroid receptor antibodies which can cross the placental barrier and cause neonatal thyrotoxicosis.

Uses:

- Differential diagnosis of etiology of Thyrotoxicosis in patients with ambiguous clinical findings non diagnostic thyroid radio-isotope scans & in pregnant or breast feeding females where thyroid radio-isotope scans are contraindicated.
- Diagnosis of clinically suspected Graves disease (Extra thyroidal manifestation of Graves disease Exophthalmos, Pretibial Myxedema, thyroid acropachy) in patients with normal thyroid function tests.
- Determining risk of Neonatal Thyrotoxicosis in a pregnant female with active or past history of Graves disease.
- Differential diagnosis of Gestational Thyrotoxicosis versus first trimester manifestation or recurrence of Graves disease.
- Assessing the risk of Graves disease relapse after antithyroid therapy.

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**URINE EXAMINATION****PHYSICAL EXAMINATION:**

Volume - 20 ml  
Colour - Pale Yellow  
Blood - Absent  
Appearance - Clear  
Deposit - Absent

**CHEMICAL EXAMINATION:**

Sp. Gravity - 1.020  
Protein - Absent  
Glucose - Absent  
Ketone - Absent  
Urobilinogens - Absent  
Bile Salts - Absent  
Bile Pigments - Absent  
Reaction - Acidic

**MICROSCOPIC EXAMINATION:** [ After centrifugation at 2000 r.p.m. for 5 minutes ]

Pus Cells - 1-2 /H.P.F  
Red Cells - Absent /H.P.F.  
Epithelial Cells - Occ. /H.P.F.  
Casts - Occ.  
Crystals - Absent  
Yeast Cells - Absent  
Trichomonas Vag. - Absent  
Bacteria - Absent

**NOTE: Done by Urometer 120 (Abott)**

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