

## Client

Jeevan Jyoti HLM

Pathkind Diagnostics Pvt. Ltd.

162, Lowther Road, Bai Ka Bagh, Prayagraj

## Processed By

Pathkind Diagnostics Pvt. Ltd.

162, Lowther Road, Bai Ka Bagh, Prayagraj

Uttar Pradesh-211003

Name	: Mr. SIDDHARTHA KUMAR MISHRA REG-307287	Billing Date	: 09/07/2022 10:42:28
Age	: 32 Yrs	Sample Collected on	: 09/07/2022 17:06:38
Sex	: Male	Sample Received on	: 09/07/2022 17:32:24
P. ID No.	: P121218819	Report Released on	: 09/07/2022 17:55:38
Accession No	: 12122204310	Barcode No.	: 1212028088
Referring Doctor	: SELF	Ref no.	:
Referred By	:		

## Report Status - Final

Test Name	Result	Biological Ref. Interval	Unit
<b>HAEMATOLOGY</b>			
<b>Complete Blood Count (CBC)</b>			
<b>Haemoglobin (Hb)</b> <i>Sample: Whole Blood EDTA Method: Photometric measurement</i>	<b>12.8 L</b>	13.0 - 17.0	gm/dL
<b>Total WBC Count / TLC</b> <i>Sample: Whole Blood EDTA Method: Impedance</i>	<b>6.4</b>	4.0 - 10.0	thou/ $\mu$ L
<b>RBC Count</b> <i>Sample: Whole Blood EDTA Method: Impedance</i>	<b>6.2 H</b>	4.5 - 5.5	million/ $\mu$ L
<b>PCV / Hematocrit</b> <i>Sample: Whole Blood EDTA Method: Impedance</i>	<b>42.2</b>	40.0 - 50.0	%
<b>MCV</b> <i>Sample: Whole Blood EDTA Method: Calculated</i>	<b>68.3 L</b>	83.0 - 101.0	fL
<b>MCH</b> <i>Sample: Whole Blood EDTA Method: Calculated</i>	<b>21.2 L</b>	27.0 - 32.0	pg
<b>MCHC</b> <i>Sample: Whole Blood EDTA Method: Calculated</i>	<b>31.1 L</b>	31.5 - 34.5	g/dL
<b>RDW (Red Cell Distribution Width)</b> <i>Sample: Whole Blood EDTA Method: Calculated</i>	<b>12.0</b>	11.8 - 15.6	%
<b>DLC (Differential Leucocyte Count)</b> <i>Method: Flowcytometry/Microscopy</i>			
<b>Neutrophils</b> <i>Sample: Whole Blood EDTA Method: VCS Technology &amp; Microscopy</i>	<b>67</b>	40 - 80	%
<b>Lymphocytes</b> <i>Sample: Whole Blood EDTA Method: VCS Technology &amp; Microscopy</i>	<b>26</b>	20 - 40	%

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<b>Eosinophils</b> <i>Sample: Whole Blood EDTA</i> <i>Method: VCS Technology &amp; Microscopy</i>	03	01 - 06	%
<b>Monocytes</b> <i>Sample: Whole Blood EDTA</i> <i>Method: VCS Technology &amp; Microscopy</i>	04	02 - 10	%
<b>Basophils</b> <i>Sample: Whole Blood EDTA</i> <i>Method: VCS Technology &amp; Microscopy</i>	00	00 - 02	%
<b>Absolute Neutrophil Count</b> <i>Sample: Whole Blood EDTA</i>	4288	2000 - 7000	/ $\mu$ L
<b>Absolute Lymphocyte Count</b> <i>Sample: Whole Blood EDTA</i>	1664	1000 - 3000	/ $\mu$ L
<b>Absolute Eosinophil Count</b> <i>Sample: Whole Blood EDTA</i>	192	20 - 500	/ $\mu$ L
<b>Absolute Monocyte Count</b> <i>Sample: Whole Blood EDTA</i>	256	200 - 1000	/ $\mu$ L
<b>Absolute Basophil Count</b> <i>Sample: Whole Blood EDTA</i>	0 L	20 - 100	/ $\mu$ L
<b>DLC Performed By</b> <i>Sample: Whole Blood EDTA</i>	EDTA Smear		
<b>Platelet Count</b> <i>Sample: Whole Blood EDTA</i> <i>Method: Impedance</i>	170	150 - 410	thou/ $\mu$ L
<b>MPV (Mean Platelet Volume)</b> <i>Sample: Whole Blood EDTA</i> <i>Method: Calculated</i>	10.9	6.8 - 10.9	fL
<b>Erythrocyte Sedimentation Rate (ESR)</b> <i>Sample: Whole Blood EDTA</i> <i>Method: Modified Westergren Method</i>	06	<10	mm 1st Hour

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Test Name	Result	Biological Ref. Interval	Unit
<b>Blood Group</b>			
<b>Blood Grouping</b> <i>Sample: Whole Blood EDTA</i>	" O "		
<b>Rh (D) Typing</b> <i>Sample: Whole Blood EDTA</i>	Positive		
<b>BIOCHEMISTRY</b>			
<b>Fasting Plasma Glucose</b> <i>Sample: Fluoride Plasma - F</i>	90	74 - 106	mg/dl
<b>Glucose Post-Prandial</b> <i>Sample: Fluoride Plasma - PP</i> <i>Method: Hexokinase</i>	101	70 - 140	mg/dl
<b>Liver Function Extended Panel</b>			
<b>Bilirubin Total</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	0.9	<1.1	mg/dL
<b>Bilirubin Direct</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	0.3 H	<0.2	mg/dL
<b>Serum Bilirubin (Indirect)</b> <i>Sample: Serum</i> <i>Method: Calculated</i>	0.6	<0.90	mg/dL
<b>SGOT / AST</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	22	<37	U/L
<b>SGPT / ALT</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	18	<41	U/L
<b>Alkaline Phosphatase (ALP)</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	97	<128	U/L
<b>Lactate Dehydrogenase (LDH)</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	206	<232	U/L

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<b>Referred By</b> :	

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Test Name	Result	Biological Ref. Interval	Unit
<b>Gamma-Glutamyl Transferase (GGT)</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	15	<71	U/L
<b>Total Protein</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	7.1	6.4 - 8.3	g/dL
<b>Albumin</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	4.8	4.0 - 4.9	g/dL
<b>Globulin</b> <i>Sample: Serum</i> <i>Method: Calculated</i>	2.3	1.9 - 3.7	g/dL
<b>Albumin Globulin A/G Ratio</b> <i>Sample: Serum</i> <i>Method: Calculated</i>	2.1	1.0 - 2.1	
<b>Thyroid Profile Total</b>			
<b>Total T3 (Triiodothyronine)</b> <i>Sample: Serum</i> <i>Method: ECLIA</i>	1.54	0.80 - 2.00	ng/mL
<b>Total T4 (Thyroxine)</b> <i>Sample: Serum</i> <i>Method: ECLIA</i>	9.69	5.10 - 14.10	µg/dL
<b>TSH 3rd Generation</b> <i>Sample: Serum</i> <i>Method: ECLIA</i>	1.940	0.270 - 4.200	µIU/mL

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**CLINICAL PATHOLOGY****Stool Routine & Microscopic Examination****Physical Examination**

<b>Colour</b> <i>Sample: Stool</i>	Brownish	Yellowish Brown
<b>Consistency</b> <i>Sample: Stool</i>	Semi Solid	Semi Solid
<b>Mucus</b> <i>Sample: Stool</i>	Absent	Absent
<b>Blood</b> <i>Sample: Stool</i>	Absent	Absent
<b>Odour</b> <i>Sample: Stool</i>	Fecal	Fecal

**Microscopic Examination**

<b>Cyst</b> <i>Sample: Stool</i>	Not Detected	Not Detected
<b>Trophozoites</b> <i>Sample: Stool</i>	Not Detected	Not Detected
<b>Charcot - Leyden Crystals</b> <i>Sample: Stool</i>	Not Detected	Not Detected
<b>Ova</b> <i>Sample: Stool</i>	Not Detected	Not Detected
<b>Adult Parasite</b> <i>Sample: Stool</i>	Not Detected	Not Detected

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Test Name	Result	Biological Ref. Interval	Unit
<b>RBC</b> <i>Sample: Stool</i>	Not Detected	0 - 0	/hpf
<b>Pus Cells</b> <i>Sample: Stool</i>	0 - 2	0 - 5	/HPF
<b>Lipid Profile</b>			
<b>Total Cholesterol</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	150	No risk : < 200 Moderate risk : 200-239 High risk : =240	mg/dL
<b>Triglycerides</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	79	Desirable : < 150 Borderline High : 150 - 199 High : 200 - 499 Very High : >= 500	mg/dL
<b>LDL Cholesterol (Calculated)</b> <i>Sample: Serum</i> <i>Method: Calculated</i>	92	Optimal : <100 Near Optimal : 100 - 129 Borderline High : 130 - 160 High : 161 - 189 Very High : >=190	mg/dL
<b>HDL Cholesterol</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	42	Low : < 40 Optimal : 40 - 60 High : > 60	mg/dL
<b>Non HDL Cholesterol</b> <i>Sample: Serum</i>	108	< 130	mg/dL
<b>VLDL Cholesterol</b> <i>Sample: Serum</i> <i>Method: Calculated</i>	15.8	Desirable 10 - 35	mg/dL
<b>Total Cholesterol / HDL Ratio</b> <i>Sample: Serum</i> <i>Method: Calculated</i>	3.57	Low Risk : 3.3 - 4.4 Average Risk : 4.5 - 7.0 Moderate Risk : 7.1 - 11.0 High Risk : > 11.0	
<b>LDL / HDL Ratio</b> <i>Sample: Serum</i> <i>Method: Calculated</i>	2.2	0.5 - 3.0	

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Low Risk : 0.5 - 3.0  
Moderate Risk : 3.1 - 6.0  
High Risk : > 6.0

## Kidney Profile (KFT)

## Blood Urea

## Blood Urea Nitrogen (BUN)

Sample: Serum

Method: Spectrophotometry-Urease / GLDH

7.31 L

8.87 - 20.50

mg/dL

## Urea

Sample: Serum

Method: Spectrophotometry

15.64 L

17.00 - 43.00

mg/dL

## Creatinine

Sample: Serum

Method: Spectrophotometry

0.84

0.70 - 1.30

mg/dL

## BUN Creatinine Ratio

Sample: Serum

Method: Calculated

9 L

10 - 20

## Calcium

Sample: Serum

Method: Spectrophotometry

9.9

8.6 - 10.0

mg/dL

## Uric Acid

Sample: Serum

Method: Spectrophotometry

6.7

3.4 - 7.0

mg/dL

## Electrolytes (Na/K/Cl)

## Sodium

Sample: Serum

Method: ISE

141

136 - 145

mmol/L

## Potassium

Sample: Serum

Method: ISE

4.3

3.5 - 5.1

mmol/L

## Chloride

Sample: Serum

Method: ISE

106

97 - 107

mmol/L

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<b>Total Protein</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	7.1	6.4 - 8.3	g/dL
<b>Albumin</b> <i>Sample: Serum</i> <i>Method: Spectrophotometry</i>	4.8	4.0 - 4.9	g/dL
<b>Globulin</b> <i>Sample: Serum</i> <i>Method: Calculated</i>	2.3	1.9 - 3.7	g/dL
<b>Albumin/Globulin (A/G) Ratio</b> <i>Sample: Serum</i> <i>Method: Calculated</i>	2.1	1.0 - 2.1	g/dL

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**CLINICAL PATHOLOGY****Urine Routine & Microscopic Examination**

Method: Reflectance Photometry

**Physical Examination****Colour**

Sample: Urine

Method: Physical Examination

amber

Pale Yellow

**Appearance**

Sample: Urine

Method: Physical Examination

Slightly Hazy

Clear

**Specific Gravity**

Sample: Urine

Method: pKa change of pretreated polyelectrolytes

1.025

1.003 - 1.035

**pH**

Sample: Urine

Method: Double indicator principle

5.0

4.7 - 7.5

**Chemical Examination****Glucose**

Sample: Urine

Method: Glucose oxidase/peroxidase

Not Detected

Not Detected

**Protein**

Sample: Urine

Method: Protein-error-of-indicators principle

Not Detected

Not Detected

**Ketones**

Sample: Urine

Method: Sodium nitroprusside reaction

Not Detected

Not Detected

**Blood**

Sample: Urine

Method: Peroxidase

Not Detected

Not Detected

**Bilirubin**

Sample: Urine

Method: Diazo reaction

Not Detected

Not Detected

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<b>Urobilinogen</b> <i>Sample: Urine</i> <i>Method: Ehrlich's reaction</i>	Normal	Normal	
<b>Nitrite</b> <i>Sample: Urine</i> <i>Method: Nitrite Test</i>	Not Detected	Not Detected	
<b>Microscopic Examination</b> <i>Method: Microscopy</i>			
<b>Pus Cells</b> <i>Sample: Urine</i>	8-10	0 - 5	/hpf
<b>RBC</b> <i>Sample: Urine</i>	Not Detected	Not Detected	/hpf
<b>Epithelial Cells</b> <i>Sample: Urine</i>	3-5	0 - 5	/hpf
<b>Casts</b> <i>Sample: Urine</i>	Not Detected	Not Detected	/hpf
<b>Crystals</b> <i>Sample: Urine</i>	Not Detected	Not Detected	/hpf
<b>Bacteria</b> <i>Sample: Urine</i>	Not Detected	Not Detected	/hpf
<b>Remarks</b> <i>Sample: Urine</i>			

**Remarks** : Microscopic Examination is performed on urine sediment  
**Complete Blood Count (CBC)**

Clinical Significance :

CBC comprises of estimation of the cellular componenets of blood including RBCs, WBCs and Platelets. Mean corpuscular volume (MCV) is a

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measure of the size of the average RBC, MCH is a measure of the hemoglobin content of the average RBC and MCHC is the hemoglobin concentration per RBC. The red cell distribution width (RDW) is a measure of the degree of variation in RBC size (anisocytosis) and is helpful in distinguishing between some anemias. CBC examination is used as a screening tool to confirm a hematologic disorder, to establish or rule out a diagnosis, to detect an unsuspected hematologic disorder, or to monitor effects of radiation or chemotherapy. Abnormal results may be due to a primary disorder of the cell-producing organs or an underlying disease. Results should be interpreted in conjunction with the patient's clinical picture and appropriate additional testing performed.

**Erythrocyte Sedimentation Rate (ESR)**Clinical Significance :

The erythrocyte sedimentation rate (ESR) is a simple but non-specific test that helps to detect inflammation associated with conditions such as infections, cancers, and autoimmune diseases.

**Total T3 (Triiodothyronine)**Clinical Significance :

Thyroid hormones, T3 and T4, which are secreted by the thyroid gland, regulate a number of developmental, metabolic, and neural activities throughout the body. The thyroid gland synthesizes 2 hormones - T3 and T4. T3 production in the thyroid gland constitutes approximately 20% of the total circulating T3, 80% being produced by peripheral conversion from T4. T3 is more potent biologically. Total T3 comprises of Free T3 and bound T3. Bound T3 remains bound to carrier proteins like thyroid-binding globulin, prealbumin, and albumin). Only the free forms are metabolically active. In hyperthyroidism, both T4 and T3 levels are usually elevated, but in some rare cases, only T3 elevation is also seen. In hypothyroidism T4 and T3 levels are both low. T3 levels are frequently low in sick or hospitalized euthyroid patients.

**Total T4 (Thyroxine)**Clinical Significance :

Total T4 is synthesized in the thyroid gland. About 0.05% of circulating T4 is in the free or biologically active form. The remainder is bound to thyroxine-binding globulin (TBG), prealbumin, and albumin. High levels of T4 (and FT4) causes hyperthyroidism and low levels lead to hypothyroidism.

**TSH 3rd Generation**Clinical Significance :

12122204310 Mr. SIDDHARTHA KUMAR MISHRA REG



**Client**

Jeevan Jyoti HLM

Pathkind Diagnostics Pvt. Ltd.

162, Lowther Road, Bai Ka Bagh, Prayagraj

**Processed By**

Pathkind Diagnostics Pvt. Ltd.

162, Lowther Road, Bai Ka Bagh, Prayagraj

Uttar Pradesh-211003

<b>Name</b>	: Mr. SIDDHARTHA KUMAR MISHRA REG-307287	<b>Billing Date</b>	: 09/07/2022 10:42:28
<b>Age</b>	: 32 Yrs	<b>Sample Collected on</b>	: 09/07/2022 17:06:38
<b>Sex</b>	: Male	<b>Sample Received on</b>	: 09/07/2022 17:32:24
<b>P. ID No.</b>	: P121218819	<b>Report Released on</b>	: 09/07/2022 17:55:38
<b>Accession No</b>	: 12122204310	<b>Barcode No.</b>	: 1212028086, 1212028088, 1212028085, 1212028089, 16835967, 1212028087
<b>Referring Doctor</b>	: SELF	<b>Ref no.</b>	:
<b>Referred By</b>	:		

**Report Status - Final**

Test Name	Result	Biological Ref. Interval	Unit
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TSH levels are elevated in primary hypothyroidism and low in primary hyperthyroidism. Evaluation of TSH is useful in the differential diagnosis of primary from secondary and tertiary hypothyroidism. In primary hypothyroidism, TSH levels are elevated, while in secondary and tertiary hypothyroidism, TSH levels are low or normal. High TSH level in the presence of normal FT4 is called subclinical hypothyroidism and low TSH with normal FT4 is called subclinical hyperthyroidism. Sick, hospitalized patients may have falsely low or transiently elevated TSH. Significant diurnal

**Stool Routine & Microscopic Examination**Clinical Significance :

Routine and microscopic examination of stool sample comprises of macroscopic as well as microscopic examination of the sample for presence of parasitic ova and cysts.

**Uric Acid**Clinical Significance :

Uric acid is the final product of purine metabolism. Serum uric acid levels are raised in case of increased purine synthesis, inherited metabolic disorder, excess dietary purine intake, increased nucleic acid turnover, malignancy and cytotoxic drugs. Decreased levels are seen in chronic renal failure, severe hepatocellular disease with reduced purine synthesis, defective renal tubular reabsorption, overtreatment of hyperuricemia with allopurinol, as well as some cancer therapies.

**Urine Routine & Microscopic Examination**Clinical Significance :

Urine routine examination and microscopy comprises of a set of screening tests that can detect some common diseases like urinary tract infections, kidney disorders, liver problems, diabetes or other metabolic conditions. Physical characteristics (colour and appearance), chemical composition (glucose, protein, ketone, blood, bilirubin and urobilinogen) and microscopic content ( pus cells, epithelial cells, RBCs, casts and crystals) are analyzed and reported.

**\*\* End of Report\*\*****Dr. Ankit Singh**

MBBS, MD (Pathologist)

Lab Head

12122204310 Mr. SIDDHARTHA KUMAR MISHRA REG



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