

**DEPARTMENT OF BIOCHEMISTRY**

<b>Patient Name</b> :	Mr. RAMESH KUMAR	<b>Bill Date</b> :	23/03/2024
<b>MR No</b> :	38448	<b>Reporting Date</b> :	23/03/2024
<b>Age/Sex</b> :	59 Years / Male	<b>Sample ID</b> :	204097
<b>Type</b> :	OPD	<b>Bill/Req. No.</b> :	24384167
<b>TPA/Corporate</b> :	MEDIWHEEL	<b>Ref Doctor</b> :	Dr. EMO
<b>IP No.</b> :			

Test	Result	Bio. Ref. Interval	Units
<b>BLOOD GLUCOSE FASTING AND PP</b>			
PLASMA GLUCOSE (FASTING)	<b>243</b> <i>H</i>	70 - 110	mg/dl
PLASMA POST-GLUCOSE	<b>294</b> <i>H</i>	80 - 150	mg/dL

**BLOOD GROUP**

BLOOD GROUP      " O " RH POSITIVE

**COMPLETE HAEMOGRAM**

<b>CBC</b>			
HAEMOGLOBIN	<b>17.1</b> <i>H</i>	12.0 - 16.5	g/dL
TOTAL LEUCOCYTE COUNT	9400	4000 - 11000	/cumm
RED BLOOD CELL COUNT	5.82	4.0 - 6.0	millions/cumm
PCV (HAEMATOCRIT)	51.8	40.0 - 54.0	%
MEAN CORPUSCULAR VOLUME	87.2	78 - 98	fL
MEAN CORPUSCULAR HAEMOGLOBIN	29.4	26.5 - 32.5	Picogrames
MEAN CORPUSCULAR HB CONC	33.0	32 - 37	g/dL
PLATELET COUNT	1.89	1.50 - 4.50	Lakh/cumm
NEUTROPHILS	71	40 - 73.0	%
LYMPHOCYTES	24	20 - 40	%
EOSINOPHILS	01	0.0 - 6.0	%
MONOCYTES	04	2.0 - 10.0	%
BASOPHILS	00	0.0 - 1.0	%
ABSOLUTE NEUTROPHIL	6674	2000 - 7000	cells/cumm
ABSOLUTE LYMPHOCYTE	2256	1000 - 3000	cells/cumm
ABSOLUTE EOSINOPHIL	94	20 - 500	cells/cumm
ABSOLUTE MONOCYTES	376	200 - 1000	cells/cumm
ABSOLUTE BASOPHILS	<b>0</b> <i>L</i>	20 - 100	cells/cumm

Checked By

Dr. Pradip Kumar  
(Consultant Microbiologist)

Dr. Nisha Rana  
(Consultant Pathologist)

**DEPARTMENT OF HAEMATOLOGY**

<b>Patient Name</b> :	Mr. RAMESH KUMAR	<b>Bill Date</b> :	23/03/2024
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<b>TPA/Corporate</b> :	MEDIWHEEL	<b>Ref Doctor</b> :	Dr. EMO
<b>IP No.</b> :			

Test	Result	Bio. Ref. Interval	Units
RDW-CV	12.9	11.5 - 14.5	%
E.S.R.	15	0 - 15	mm/hr

**HBA1C**

HBA1C	<b>10.0</b>	<i>H</i>	%
ESTIMATED AVERAGE GLUCOSE (EAG)	240.3		mg/dL

**Note :** HBA1c result is suggestive of Diabetes/ higher than glycemic goal in a known Diabetic patient.  
Please note, glycemic goal should be individualized based on duration of diabetes, age/life expectancy, comorbid conditions, known CVD or advanced microvascular complications, hypoglycaemia unawareness, and individual patient considerations.  
Please Correlate Clinically.

**KFT(KIDNEY FUNCTION TEST)/RFT/Renal Profile**

SERUM UREA	32	13.0 - 45.0	mg/dL
SERUM CREATININE	1.2	0.5 - 1.4	mg/dL
SERUM URIC ACID	4.7	3.6 - 7.2	mg/dL
SERUM SODIUM	133	130 - 149	mmol/L
SERUM POTASSIUM	4.8	3.5 - 5.5	mmol/L

**LFT(LIVER FUNCTION TEST)**

<b>LFT</b>			
TOTAL BILIRUBIN	0.6	0.1 - 1.2	mg/dL
DIRECT BILIRUBIN	0.2	0.00 - 0.30	mg/dL
INDIRECT BILIRUBIN	0.4	Adult: 0 - 0.8	mg/dL
SGOT (AST)	28	0.0 - 45	IU/L
SGPT (ALT)	40	00 - 45.00	IU/L
ALP	123	41 - 137	U/L

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## DEPARTMENT OF BIOCHEMISTRY

**Patient Name** : Mr. RAMESH KUMAR  
**MR No** : 38448  
**Age/Sex** : 59 Years / Male  
**Type** : OPD  
**TPA/Corporate** : MEDIWHEEL  
**IP No.** :

**Bill Date** : 23/03/2024  
**Reporting Date** : 23/03/2024  
**Sample ID** : 204097  
**Bill/Req. No.** : 24384167  
**Ref Doctor** : Dr. EMO

Test	Result	Bio. Ref. Interval	Units
TOTAL PROTEINS	7.6	6.0 - 8.2	g/dL
ALBUMIN	4.8	3.20 - 5.00	g/dL
GLOBULIN	2.8	2.0 - 3.50	g/dL
A/G RATIO	1.71		

## LIPID PROFILE

Test	Result	Remarks	Bio. Ref. Interval	Units
SERUM CHOLESTROL	234	H	0 - 200	mg/dl
SERUM TRIGLYCERIDES	127		Up to 150	mg/dl
HDL CHOLESTEROL	49		30 - 60	mg/dl
VLDL CHOLESTEROL	25.4		*Less than 30	mg/dL
LDL CHOLESTEROL	159.6		Optimal <100, Above Opt. 100-129 -high 160-189	mg/dl
LDL CHOLESTEROL/HDL RATIO	3.26		Desirable Level : 0.5 - 3.0 Borderline Risk : 3.0 - 6.0 High Risk : > 6.0	

## URINE ROUTINE EXAMINATION

### PHYSICAL EXAMINATION

Test	Result	Reference	Units
VOLUME	15		ml
COLOUR	Pale Yellow	Pale Yellow	
APPEARANCE	Clear	Clear	
SPECIFIC GRAVITY	1.015		

### CHEMICAL EXAMINATION

REACTION	Acidic	
BLOOD	NIL	
ALBUMIN	NIL	NIL
GLUCOSE	Traces	NIL
PH	6.5	

### MICROSCOPIC EXAMINATION

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(Consultant Microbiologist)

Dr. Nisha Rana  
(Consultant Pathologist)

**DEPARTMENT OF CLINICAL PATHOLOGY**

**Patient Name** : Mr. RAMESH KUMAR  
**MR No** : 38448  
**Age/Sex** : 59 Years / Male  
**Type** : OPD  
**TPA/Corporate** : MEDIWHEEL  
**IP No.** :  
**Bill Date** : 23/03/2024  
**Reporting Date** : 23/03/2024  
**Sample ID** : 204111  
**Bill/Req. No.** : 24384167  
**Ref Doctor** : Dr. EMO

Test	Result	Bio. Ref. Interval	Units
PUS CELL	3-4	2-4	/HPF
EPITHELIAL CELLS	1-2	2-4	/HPF
RED BLOOD CELLS	Nil	NIL	/HPF
CASTS	NIL	NIL	
CRYSTALS	NIL	NIL	

**Note** : Albumin test positive by Multistrip Method is confirmed by Sulphosalicylic acid method.

\*\*\*\*\* END OF THE REPORT \*\*\*\*\*

Checked By :

**Dr. Pradip Kumar**  
(Consultant Microbiologist)

**Dr. Nisha Rana**  
(Consultant Pathologist)



<b>Lab No.</b>	012403230733	<b>Age/Gender</b>	59 YRS/MALE	<b>Coll. On</b>	23/Mar/2024 04:24PM
<b>Name</b>	Mr. RAMESH KUMAR 38448			<b>Reg. On</b>	23/Mar/2024
<b>Ref. Dr.</b>				<b>Approved On</b>	23/Mar/2024 06:39PM
<b>Rpt. Centre</b>	Self,undefined			<b>Printed On</b>	29/Mar/2024 08:48AM

Test Name	Value	Unit	Biological Reference Interval
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**Thyroid profile, Total (T3,T4,TSH)**

T3 (Triiodothyronine) , serum Method : ECLIA	0.73	ng/mL	0.80 - 2.0
T4 (Thyroxine) , serum Method : ECLIA	6.12	ug/dL	5.1 - 14.1
TSH (Thyroid Stimulating Hormone) , serum Method : ECLIA	0.83	uIU/ml	0.5 - 8.9

**Interpretation:**

- Primary hyperthyroidism is accompanied by elevated serum T3 and T4 values alongwith depressed TSH levels
- Primary hypothyroidism is accompanied by depressed serum T3 and T4 values and elevated serum TSH levels.
- High T3 levels coupled with normal T4 and suppressed TSH may be seen in T3 toxicosis.

**Note:** Total T3 and total T4 are highly bound to plasma proteins and are amenable to fluctuations with plasma protein content as well as due to binding defects in the thyroid hormone binding proteins.

The following ranges are recommended for pregnant females:

Gestation period	TSH (uIU/ml)
First trimester	0.1 - 2.5
Second trimester	0.2 - 3.0
Third trimester	0.3 - 3.0

<b>PSA Total, serum</b> Method : ECLIA	0.34	ng/mL	0 - 3.1
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**Interpretation:**

Prostate-specific antigen (PSA) is a glycoprotein that is produced by the prostate gland, the lining of the urethra, and the bulbourethral gland. Normally, very little PSA is secreted in the blood. Increases in glandular size and tissue damage caused by benign prostatic hypertrophy, prostatitis, or prostate cancer may increase circulating PSA levels.

In patients with previously diagnosed prostate cancer, PSA testing is advocated as an early indicator of tumor recurrence and as an indicator of response to therapy.

The test is also useful for initial screening for prostate cancer:

Total PSA levels < 2 ng/ml almost rule out the possibility of prostatic malignancy.

Total PSA levels between 2 and 10 ng/ml lie in the grey zone. Such values may be obtained in prostatitis, benign hyperplasia and malignancy. Further testing including a free PSA/PSA ratio and prostate biopsy is recommended for these patients for confirmation of the diagnosis.

Total PSA values >10 ng/ml are highly suspicious for prostate cancer but further testing, such as prostate biopsy, is needed to diagnose the exact pathology.

\*Disclaimer: This is an electronically validated report. If any discrepancy is found, it should be confirmed by the user.  
Processing Centre : Prognosis Laboratories,515-516, Sector-19, Dwarka, Behind Gupta Properties.

\*\*\* Partial Report \*\*\*



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