

Tests you can trust

Name : Rajlaxmi Rajlaxmi(26Y/F)

Date : 03 Mar 2025

Test Asked: Mediwheel Health Checkup Below 40

Report Status: Complete Report



First National Diagnostic Chain to have 100% of its Labs with NABL Accreditation*



















Your reports are digitally verifiable

Scan the QR code inside the report to check authenticity of reported values

QR code will remain active for 30 days from report release date

Accredited by







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First National Diagnostic Chain to have 100% of its Labs with NABL Accreditation[#]

NAME : RAJLAXMI RAJLAXMI(26Y/F)

REF. BY : SELF

TEST ASKED : MEDIWHEEL HEALTH CHECKUP BELOW 40

HOME COLLECTION:

B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, Mumbai,

Report Availability Summary

Note: Please refer to the table below for status of your tests.

• Ready with Cancellation



O Processing



(X) O Cancelled in Lab

Т	EST DETAILS	REPORT STATUS
M	IEDIWHEEL HEALTH CHECKUP BELOW 40	Ready 🔗
	LIPID PROFILE	Ready ⊗
	ERYTHROCYTE SEDIMENTATION RATE (ESR)	Ready ⊘
	HEMOGRAM - 6 PART (DIFF)	Ready 🔗
	T3-T4-USTSH	Ready ⊘
	FASTING BLOOD SUGAR(GLUCOSE)	Ready 🔗
	HbA1c	Ready ⊘
	COMPLETE URINE ANALYSIS	Ready 🔗
	VITAMIN B-12	Ready ⊘
	LIVER FUNCTION TESTS	Ready ⊗
	PHOSPHOROUS	Ready 📀
	SERUM ELECTROLYTES	Ready 📀
	KIDPRO	Ready 🛇
	25-OH VITAMIN D (TOTAL)	Ready 🕢

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: RAJLAXMI RAJLAXMI(26Y/F) NAME

HOME COLLECTION:

REF. BY : SELF B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel

: MEDIWHEEL HEALTH CHECKUP BELOW 40 **TEST ASKED**

Deeepak Talkies, Mumbai, Maharashtra, 400013

Summary Report

Cumilary Report						
Tes	Tests outside reference range					
TEST NAME	OBSERVED VALUE	UNITS	Bio. Ref. Interval.			
COMPLETE HEMOGRAM						
MEAN PLATELET VOLUME(MPV)	13	fL	6.5-12			
MONOCYTES - ABSOLUTE COUNT	0.13	$X~10^3$ / μL	0.2 - 1.0			
PLATELET DISTRIBUTION WIDTH(PDW)	17.5	fL	9.6-15.2			
PLATELET TO LARGE CELL RATIO(PLCR)	50	%	19.7-42.4			
COMPLETE URINE ANALYSIS						
APPEARANCE	SLIGHT CLOUDY	-	Clear			
BACTERIA	PRESENT	-	Absent			
EPITHELIAL CELLS	6	cells/HPF	0-5			
RED BLOOD CELLS	7	cells/HPF	0-5			
URINE BLOOD	PRESENT	-	Absent			
LIPID						
LDL CHOLESTEROL - DIRECT	109	mg/dL	< 100			
LIVER						
ALKALINE PHOSPHATASE	129.6	U/L	45-129			
SERUM GLOBULIN	3.48	gm/dL	2.5-3.4			
OTHER COUNTS						
ERYTHROCYTE SEDIMENTATION RATE (ESR)	38	mm / hr	0 - 20			
RENAL						
BLOOD UREA NITROGEN (BUN)	6.31	mg/dL	7.94 - 20.07			
UREA (CALCULATED)	13.5	mg/dL	Adult : 17-43			
VITAMIN						
25-OH VITAMIN D (TOTAL)	5.25	ng/mL	30-100			

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HOME COLLECTION:

B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, Mumbai, Maharashtra,

VALUE TEST NAME TECHNOLOGY UNITS 25-OH VITAMIN D (TOTAL) C.L.I.A 5.25 ng/mL Bio. Ref. Interval. :-

DEFICIENCY: <20 ng/ml || INSUFFICIENCY: 20-<30 ng/ml SUFFICIENCY: 30-100 ng/ml || TOXICITY: >100 ng/ml

Clinical Significance:

Vitamin D is a fat soluble vitamin that has been known to help the body absorb and retain calcium and phosphorous; both are critical for building bone health. Decrease in vitamin D total levels indicate inadequate exposure of sunlight, dietary deficiency, nephrotic syndrome. Increase in vitamin D total levels indicate Vitamin D intoxication.

Specifications: Precision: Intra assay (%CV):5.3%, Inter assay (%CV):11.9%; Sensitivity:3.2 ng/ml.

Kit Validation Reference: Holick MF. Vitamin D Deficiency. N Engl J Med. 2007;357:266-81.

Please correlate with clinical conditions.

Method:-Fully Automated Chemi Luminescent Immuno Assay

Sample Collected on (SCT)

Sample Received on (SRT)

Report Released on (RRT)

Sample Type

Labcode **Barcode**

: 03 Mar 2025 08:07

: 03 Mar 2025 10:19

: 03 Mar 2025 14:37

: SERUM

: 0303040177/DS853 Dr.Samrita Samaddar MD (Path)Dr Sumanta Basak, DPB

: DT280269

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HOME COLLECTION:

B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, Mumbai, Maharashtra,

VALUE TEST NAME TECHNOLOGY UNITS VITAMIN B-12 C.L.I.A 244 pg/mL Bio. Ref. Interval. :-

Normal: 211 - 911 pg/ml

Clinical significance:

Vitamin B12 or cyanocobalamin, is a complex corrinoid compound found exclusively from animal dietary sources, such as meat, eggs and milk. It is critical in normal DNA synthesis, which in turn affects erythrocyte maturation and in the formation of myelin sheath. Vitamin-B12 is used to find out neurological abnormalities and impaired DNA synthesis associated with macrocytic anemias. For diagnostic purpose, results should always be assessed in conjunction with the patients medical history, clinical examination and other findings.

Specifications: Intra assay (%CV):5.0%, Inter assay (%CV):9.2 %; Sensitivity:45 pg/ml

Kit Validation reference:

Chen IW, Sperling MI, Heminger LA. Vitamin B12. In: Pesce AJ, Kaplan LA, eds. Methods in Clinical Chemistry. St. Louis: CV Mosby; 1987:569-73.

Please correlate with clinical conditions.

COMPETITIVE CHEMI LUMINESCENT IMMUNO ASSAY Method:-

Sample Collected on (SCT) : 03 Mar 2025 08:07

Sample Received on (SRT) : 03 Mar 2025 10:19

Report Released on (RRT) : 03 Mar 2025 14:37

: SERUM Sample Type

: 0303040177/DS853 Dr.Samrita Samaddar MD (Path)Dr Sumanta Basak, DPB Labcode

Barcode : DT280269 Page: 2 of 13

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: SELF REF. BY

TEST ASKED : MEDIWHEEL HEALTH CHECKUP BELOW 40

HOME COLLECTION:

B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, Mumbai, Maharashtra, 400013

TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
TOTAL CHOLESTEROL	PHOTOMETRY	170	mg/dL	< 200
HDL CHOLESTEROL - DIRECT	PHOTOMETRY	45	mg/dL	40-60
LDL CHOLESTEROL - DIRECT	PHOTOMETRY	109	mg/dL	< 100
TRIGLYCERIDES	PHOTOMETRY	107	mg/dL	< 150
TC/ HDL CHOLESTEROL RATIO	CALCULATED	3.8	Ratio	3 - 5
TRIG / HDL RATIO	CALCULATED	2.37	Ratio	< 3.12
LDL / HDL RATIO	CALCULATED	2.4	Ratio	1.5-3.5
HDL / LDL RATIO	CALCULATED	0.41	Ratio	> 0.40
NON-HDL CHOLESTEROL	CALCULATED	124.9	mg/dL	< 160
VLDL CHOLESTEROL	CALCULATED	21.32	mg/dL	5 - 40

Please correlate with clinical conditions.

Method:

CHOL - Cholesterol Oxidase, Esterase, Peroxidase

HCHO - Direct Enzymatic Colorimetric

LDL - Direct Measure

TRIG - Enzymatic, End Point

TC/H - Derived from serum Cholesterol and Hdl values

TRI/H - Derived from TRIG and HDL Values

LDL/ - Derived from serum HDL and LDL Values

HD/LD - Derived from HDL and LDL values.

NHDL - Derived from serum Cholesterol and HDL values

VLDL - Derived from serum Triglyceride values

*REFERENCE RANGES AS PER NCEP ATP III GUIDELINES:

TOTAL CHOLESTEROL	(mg/dl)	HDL	(mg/dl)	LDL	(mg/dl)	TRIGLYCERIDES	(mg/dl)
DESIRABLE	<200	LOW	<40	OPTIMAL	<100	NORMAL	<150
BORDERLINE HIGH	200-239	HIGH	>60	NEAR OPTIMAL	100-129	BORDERLINE HIGH	150-199
HIGH	>240			BORDERLINE HIGH	130-159	HIGH	200-499
				HIGH	160-189	VERY HIGH	>500
				VERY HIGH	>190		

Alert !!! 10-12 hours fasting is mandatory for lipid parameters. If not, values might fluctuate,

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Sample Received on (SRT) : 03 Mar 2025 10:19 Report Released on (RRT) : 03 Mar 2025 14:37

Sample Type : SERUM

Labcode : 0303040177/DS853

Barcode . DT280269

Dr.Samrita Samaddar MD (Path)

Dr Sumanta Basak, DPB

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REF. BY : SELF

TEST ASKED : MEDIWHEEL HEALTH CHECKUP BELOW 40

HOME COLLECTION:

B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, Mumbai, Maharashtra, 400013

TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
ALKALINE PHOSPHATASE	PHOTOMETRY	129.6	U/L	45-129
BILIRUBIN - TOTAL	PHOTOMETRY	0.55	mg/dL	0.3-1.2
BILIRUBIN -DIRECT	PHOTOMETRY	0.11	mg/dL	< 0.3
BILIRUBIN (INDIRECT)	CALCULATED	0.44	mg/dL	0-0.9
GAMMA GLUTAMYL TRANSFERASE (GGT)	PHOTOMETRY	14.9	U/L	< 38
ASPARTATE AMINOTRANSFERASE (SGOT)	PHOTOMETRY	20.1	U/L	< 31
ALANINE TRANSAMINASE (SGPT)	PHOTOMETRY	20	U/L	< 34
SGOT / SGPT RATIO	CALCULATED	1	Ratio	< 2
PROTEIN - TOTAL	PHOTOMETRY	7.74	gm/dL	5.7-8.2
ALBUMIN - SERUM	PHOTOMETRY	4.26	gm/dL	3.2-4.8
SERUM GLOBULIN	CALCULATED	3.48	gm/dL	2.5-3.4
SERUM ALB/GLOBULIN RATIO	CALCULATED	1.22	Ratio	0.9 - 2

Please correlate with clinical conditions.

Method:

ALKP - Modified IFCC method

BILT - Vanadate Oxidation

BILD - Vanadate Oxidation

BILI - Derived from serum Total and Direct Bilirubin values

GGT - Modified IFCC method

SGOT - IFCC* Without Pyridoxal Phosphate Activation

SGPT - IFCC* Without Pyridoxal Phosphate Activation

OT/PT - Derived from SGOT and SGPT values.

PROT - Biuret Method

SALB - Albumin Bcg¹method (Colorimetric Assay Endpoint)

SEGB - DERIVED FROM SERUM ALBUMIN AND PROTEIN VALUES

A/GR - Derived from serum Albumin and Protein values

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Sample Type : SERUM

Labcode : 0303040177/DS853

Barcode . DT280269

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B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, Mumbai, Maharashtra, 400013

TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval
CALCIUM	PHOTOMETRY	9.04	mg/dL	8.8-10.6
URIC ACID	PHOTOMETRY	4.43	mg/dL	3.2 - 6.1
BLOOD UREA NITROGEN (BUN)	PHOTOMETRY	6.31	mg/dL	7.94 - 20.07
UREA (CALCULATED)	CALCULATED	13.5	mg/dL	Adult : 17-43
CREATININE - SERUM	PHOTOMETRY	0.58	mg/dL	0.55-1.02
UREA / SR.CREATININE RATIO	CALCULATED	23.28	Ratio	< 52
BUN / SR.CREATININE RATIO	CALCULATED	10.88	Ratio	9:1-23:1
PHOSPHOROUS	PHOTOMETRY	3.5	mg/dL	2.4 - 5.1
SODIUM	I.S.E - INDIRECT	137.5	mmol/L	136 - 145
POTASSIUM	I.S.E - INDIRECT	5.06	mmol/L	3.5 - 5.1
CHLORIDE	I.S.E - INDIRECT	101.2	mmol/L	98 - 107

Please correlate with clinical conditions.

Method:

CALC - Arsenazo III Method, End Point.

URIC - Uricase / Peroxidase Method

BUN - Kinetic UV Assay.

UREAC - Derived from BUN Value.

SCRE - Creatinine Enzymatic Method

UR/CR - Derived from UREA and Sr.Creatinine values.

B/CR - Derived from serum Bun and Creatinine values

PHOS - UNREDUCED PHOSPHOMOLYBDATE METHOD

SOD - ION SELECTIVE ELECTRODE - INDIRECT

POT - ION SELECTIVE ELECTRODE - INDIRECT CHL - ION SELECTIVE ELECTRODE - INDIRECT

Sample Collected on (SCT) : 03 Mar 2025 08:07

: 03 Mar 2025 10:19

Sample Received on (SRT) Report Released on (RRT)

: 03 Mar 2025 14:37

Sample Type

: SERUM

Labcode **Barcode** : 0303040177/DS853

. DT280269

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TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
TOTAL TRIIODOTHYRONINE (T3)	C.M.I.A	131	ng/dL	58-159
TOTAL THYROXINE (T4)	C.M.I.A	9.09	μg/dL	4.87-11.72
TSH - ULTRASENSITIVE	C.M.I.A	2.131	μIU/mL	0.35-4.94

The Biological Reference Ranges is specific to the age group. Kindly correlate clinically. Method:

T3,T4,USTSH - Fully Automated Chemi Luminescent Microparticle Immunoassay

Pregnancy reference ranges for TSH/USTSH:

Trimester || T3 (ng/dl) || T4 (µg/dl) || TSH/USTSH (µIU/ml)

|| 83.9-196.6 || 4.4-11.5 || 0.1-2.5 1st || 86.1-217.4 || 4.9-12.2 || 0.2-3.0 2nd 3rd || 79.9-186 || 5.1-13.2 || 0.3-3.5

References:

- 1. Carol Devilia, C I Parhon. First Trimester Pregnancy ranges for Serum TSH and Thyroid Tumor reclassified as Benign. Acta Endocrinol. 2016; 12(2): 242 - 243
- 2. Kulhari K, Negi R, Kalra DK et al. Establishing Trimester specific Reference ranges for thyroid hormones in Indian women with normal pregnancy: New light through old window. Indian Journal of Contemporary medical research.

Disclaimer: Results should always be interpreted using the reference range provided by the laboratory that performed the test. Different laboratories do tests using different technologies, methods and using different reagents which may cause difference. In reference ranges and hence it is recommended to interpret result with assay specific reference ranges provided in the reports. To diagnose and monitor therapy doses, it is recommended to get tested every time at the same Laboratory.

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Sample Type : SERUM

Labcode : 0303040177/DS853 Dr.Samrita Samaddar MD (Pathor Sumanta Basak, DPB

Barcode : DT280269 Page: 6 of 13

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HOME COLLECTION:

B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, Mumbai, Maharashtra,

TEST NAME VALUE TECHNOLOGY UNITS EST. GLOMERULAR FILTRATION RATE (eGFR) **CALCULATED** mL/min/1.73 m2 128

Bio. Ref. Interval. :-

> = 90 : Normal 60 - 89 : Mild Decrease

45 - 59 : Mild to Moderate Decrease 30 - 44 : Moderate to Severe Decrease

15 - 29 : Severe Decrease

Clinical Significance

The normal serum creatinine reference interval does not necessarily reflect a normal GFR for a patient. Because mild and moderate kidney injury is poorly inferred from serum creatinine alone. Thus, it is recommended for clinical laboratories to routinely estimate glomerular filtration rate (eGFR), a "gold standard" measurement for assessment of renal function, and report the value when serum creatinine is measured for patients 18 and older, when appropriate and feasible. It cannot be measured easily in clinical practice, instead, GFR is estimated from equations using serum creatinine, age, race and sex. This provides easy to interpret information for the doctor and patient on the degree of renal impairment since it approximately equates to the percentage of kidney function remaining. Application of CKD-EPI equation together with the other diagnostic tools in renal medicine will further improve the detection and management of patients with CKD.

Reference

Levey AS, Stevens LA, Schmid CH, Zhang YL, Castro AF, 3rd, Feldman HI, et al. A new equation to estimate glomerular filtration rate. Ann Intern Med. 2009;150(9):604-12.

Please correlate with clinical conditions.

2021 CKD EPI Creatinine Equation Method:-

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: 03 Mar 2025 08:07

Sample Received on (SRT)

: 03 Mar 2025 10:19

Report Released on (RRT)

: 03 Mar 2025 14:37

Sample Type

. SERUM

Labcode

: 0303040177/DS853 Dr.Samrita Samaddar MD (Path)Dr Sumanta Basak, DPB

Barcode : DT280269

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Maharashtra, 400013

TEST NAME	TECHNOLOGY	VALUE	UNITS
HbA1c - (HPLC)			
	H.P.L.C	5.7	%

Bio. Ref. Interval. :

Bio. Ref. Interval.: As per ADA Guidelines

Below 5.7% : Normal 5.7% - 6.4% : Prediabetic >=6.5% : Diabetic

Guidance For Known Diabetics

Below 6.5%: Good Control 6.5% - 7% : Fair Control

7.0% - 8% : Unsatisfactory Control

: Poor Control

Method: Fully Automated H.P.L.C method

AVERAGE BLOOD GLUCOSE (ABG) **CALCULATED** 117 mg/dL

Bio. Ref. Interval. :

90 - 120 mg/dl : Good Control 121 - 150 mg/dl : Fair Control

151 - 180 mg/dl: Unsatisfactory Control

: Poor Control > 180 mg/dl

Method: Derived from HBA1c values

Please correlate with clinical conditions.

Sample Collected on (SCT) :03 Mar 2025 08:07 Sample Received on (SRT) : 03 Mar 2025 10:23 Report Released on (RRT) : 03 Mar 2025 12:55

Sample Type : EDTA Whole Blood

:0303040191/DS853

Barcode : DM600279

Labcode

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HOME COLLECTION:

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400013

VALUE TEST NAME TECHNOLOGY UNITS **ERYTHROCYTE SEDIMENTATION RATE (ESR) MODIFIED WESTERGREN** 38 mm / hr Bio. Ref. Interval. :-

Male: 0-15 Female: 0-20

Clinical Significance:

- An erythrocyte sedimentation rate (ESR) is a blood test that can rise if you have inflammation in your body. Its also used as a marker to monitor prognosis of an existing inflammatory/infective condition.
- Inflammation is your immune systems response to injury, infection, and many types of conditions, including immune system disorders, certain cancers and blood disorders.
- A high ESR test result may be from a condition that causes inflammation, such as: Arteritis, Arthritis, Systemic vasculitis, Polymyalgia rheumatica, Inflammatory bowel disease, Kidney disease, Infections like Tuberculosis etc, Rheumatoid arthritis and other autoimmune diseases, Heart disease, Certain cancers and many other Conditions.
- A low ESR test result may be caused by conditions such as: A blood disorder, such as: Polycythemia, Sickle cell disease (SCD), Leukocytosis, Heart failure, Certain kidney and liver problems etc.
- Certain physiological conditions also affect ESR results, these include: Pregnancy, menstrual cycle, ageing, obesity, drinking alcohol regularly, and exercise, Certain medicines and supplements also can affect ESR results.
- Hence Its always suggested to interpret ESR results in conjunction with Clinical History and other findings.

References:

https://medlineplus.gov/lab-tests/erythrocyte-sedimentation-rate-esr/

Please correlate with clinical conditions. MODIFIED WESTERGREN Method:-

Sample Collected on (SCT)

: 03 Mar 2025 08:07

Sample Received on (SRT)

: 03 Mar 2025 10:23

Report Released on (RRT)

: 03 Mar 2025 12:55

Sample Type

. EDTA Whole Blood

Labcode

: 0303040191/DS853 Dr.Samrita Samaddar MD (Path)Dr Sumanta Basak, DPB

Barcode : DM600279

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: MEDIWHEEL HEALTH CHECKUP BELOW 40 **TEST ASKED**

HOME COLLECTION:

B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, Mumbai, Maharashtra,

TEST NAME	METHODOLOGY	VALUE	UNITS	Bio. Ref. Interval
HEMOGLOBIN	SLS-Hemoglobin Method	12.2	g/dL	12.0-15.0
Hematocrit (PCV)	CPH Detection	38.6	%	36.0-46.0
Total RBC	HF & EI	4.44	X 10^6/μL	3.8-4.8
Mean Corpuscular Volume (MCV)	Calculated	86.9	fL	83.0-101.0
Mean Corpuscular Hemoglobin (MCH)	Calculated	27.5	pq	27.0-32.0
Mean Corp.Hemo. Conc (MCHC)	Calculated	31.6	g/dL	31.5-34.5
Red Cell Distribution Width - SD (RDW-SD)	Calculated	42	fL	39.0-46.0
Red Cell Distribution Width (RDW - CV)	Calculated	13.5	%	11.6-14.0
RED CELL DISTRIBUTION WIDTH INDEX (RDWI)	Calculated	264.2	-	*Refer Note below
MENTZER INDEX	Calculated	19.6	-	*Refer Note below
TOTAL LEUCOCYTE COUNT (WBC)	HF & FC	6.65	X 10 ³ / μL	4.0 - 10.0
DIFFERENTIAL LEUCOCYTE COUNT				
Neutrophils Percentage	Flow Cytometry	63.4	%	40-80
Lymphocytes Percentage	Flow Cytometry	32.5	%	20-40
Monocytes Percentage	Flow Cytometry	2	%	2-10
Eosinophils Percentage	Flow Cytometry	1.5	%	1-6
Basophils Percentage	Flow Cytometry	0.3	%	0-2
Immature Granulocyte Percentage (IG%)	Flow Cytometry	0.3	%	0.0-0.4
Nucleated Red Blood Cells %	Flow Cytometry	0.01	%	0.0-5.0
ABSOLUTE LEUCOCYTE COUNT				
Neutrophils - Absolute Count	Calculated	4.22	$X~10^3$ / μL	2.0-7.0
Lymphocytes - Absolute Count	Calculated	2.16	X 10 ³ / μL	1.0-3.0
Monocytes - Absolute Count	Calculated	0.13	X 10³ / μL	0.2 - 1.0
Basophils - Absolute Count	Calculated	0.02	$X~10^3$ / μL	0.02 - 0.1
Eosinophils - Absolute Count	Calculated	0.1	$X~10^3$ / μL	0.02 - 0.5
Immature Granulocytes (IG)	Calculated	0.02	$X~10^3$ / μL	0.0-0.3
Nucleated Red Blood Cells	Calculated	0.01	X 10 ³ / μL	0.0-0.5
PLATELET COUNT	HF & EI	218	$X~10^3$ / μL	150-410
Mean Platelet Volume (MPV)	Calculated	13	fL	6.5-12
Platelet Distribution Width (PDW)	Calculated	17.5	fL	9.6-15.2
Platelet to Large Cell Ratio (PLCR)	Calculated	50	%	19.7-42.4
Plateletcrit (PCT)	Calculated	0.28	%	0.19-0.39

Remarks: Alert!!! Predominantly normocytic normochromic with ovalocytes. Platelets: Appear adequate in smear.

Method: Fully automated bidirectional analyser (6 Part Differential SYSMEX XN-1000)

: DM600279

(Reference: *FC- flowcytometry, *HF- hydrodynamic focussing, *EI- Electric Impedence, *Hb- hemoglobin, *CPH- Cumulative pulse height)

:03 Mar 2025 08:07 Sample Collected on (SCT) : 03 Mar 2025 10:23 Sample Received on (SRT) : 03 Mar 2025 12:55 Report Released on (RRT)

Sample Type : EDTA Whole Blood

Barcode

Labcode : 0303040191/DS853

Dr.Samrita Samaddar MD (Path)

Dr Sumanta Basak, DPB

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^{*}Note - Mentzer index (MI), RDW-CV and RDWI are hematological indices to differentiate between Iron Deficiency Anemia (IDA) and Beta Thalassemia Trait (BTT). MI >13, RDWI >220 and RDW-CV >14 more likely to be IDA. MI <13, RDWI <220, and RDW-CV <14 more likely to be BTT. Suggested Clinical correlation. BTT to be confirmed with HB electrophoresis if clinically indicated.

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NAME : RAJLAXMI RAJLAXMI(26Y/F)

REF. BY : SELF

: MEDIWHEEL HEALTH CHECKUP BELOW 40 **TEST ASKED**

HOME COLLECTION:

B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, Mumbai, Maharashtra,

TEST NAME	TECHNOLOGY	VALUE	UNITS
FASTING BLOOD SUGAR(GLUCOSE)	PHOTOMETRY	90	mg/dL

Bio. Ref. Interval. :-

As per ADA Guideline: Fasting Plasma Glucose (FPG)				
Normal 70 to 100 mg/dl				
Prediabetes	Prediabetes 100 mg/dl to 125 mg/dl			
Diabetes 126 mg/dl or higher				

The assay could be affected mildly and may result in anomalous values if serum samples have heterophilic antibodies, hemolyzed, icteric or lipemic. The concentration of Glucose in a given specimen may vary due to differences in assay methods, calibration and reagent specificity. For diagnostic purposes results should always be assessed in conjunction with patients medical history, clinical findings and other findings.

Please correlate with clinical conditions.

GOD-PAP METHOD Method:-

Sample Collected on (SCT)

: 03 Mar 2025 08:07

Sample Received on (SRT)

: 03 Mar 2025 10:20

Report Released on (RRT)

: 03 Mar 2025 12:14

Sample Type

Labcode

. FLUORIDE PLASMA

Barcode : DT166257

: 0303063352/DS853 Dr.Samrita Samaddar MD (Path)Dr Sumanta Basak, DPB

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 ■ wellness@thyrocare.com

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NAME : RAJLAXMI RAJLAXMI(26Y/F)

REF. BY : SELF

TEST ASKED : MEDIWHEEL HEALTH CHECKUP BELOW 40 **HOME COLLECTION:**

B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, home, B3/509 Bhawani Mata CHS Lower Parel Deeepak Talkies, Mumbai, Maharashtra,

400013

TEST NAME	METHODOLOGY	VALUE	UNITS	Bio. Ref. Interv
Complete Urinogram				
Physical Examination				
VOLUME	Visual Determination	3	mL	-
COLOUR	Visual Determination	PALE YELLOW	-	Pale Yellow
APPEARANCE	Visual Determination	SLIGHT CLOUDY	-	Clear
SPECIFIC GRAVITY	pKa change	1.01	-	1.003-1.030
PH	pH indicator	5.5	-	5-8
Chemical Examination				
URINARY PROTEIN	PEI	ABSENT	mg/dL	Absent
URINARY GLUCOSE	GOD-POD	ABSENT	mg/dL	Absent
URINE KETONE	Nitroprusside	ABSENT	mg/dL	Absent
URINARY BILIRUBIN	Diazo coupling	ABSENT	mg/dL	Absent
UROBILINOGEN	Diazo coupling	Normal	mg/dL	<=0.2
BILE SALT	Hays sulphur	ABSENT	-	Absent
BILE PIGMENT	Ehrlich reaction	ABSENT	-	Absent
URINE BLOOD	Peroxidase reaction	PRESENT	-	Absent
NITRITE	Diazo coupling	ABSENT	-	Absent
LEUCOCYTE ESTERASE	Esterase reaction	ABSENT	-	Absent
Microscopic Examination				
MUCUS	Microscopy	ABSENT	-	Absent
RED BLOOD CELLS	Microscopy	7	cells/HPF	0-5
URINARY LEUCOCYTES (PUS CELLS)	Microscopy	2	cells/HPF	0-5
EPITHELIAL CELLS	Microscopy	6	cells/HPF	0-5
CASTS	Microscopy	ABSENT	-	Absent
CRYSTALS	Microscopy	ABSENT	-	Absent
BACTERIA	Microscopy	PRESENT	-	Absent
YEAST	Microscopy	ABSENT	-	Absent
PARASITE	Microscopy	ABSENT	-	Absent

(Reference: *PEI - Protein error of indicator, *GOD-POD - Glucose oxidase-peroxidase)

~~ End of report ~~

Sample Collected on (SCT) Sample Received on (SRT) Report Released on (RRT) : 03 Mar 2025 08:07 : 03 Mar 2025 10:08

: 03 Mar 2025 11:31

Sample Type

Labcode **Barcode**

: URINE

: 0303062886/DS853

Dr.Samrita Samaddar MD (Path) Dr Sumanta Basak, DPB

: DE473552

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Scan QR code to verify authenticity of reported results; active for 30 days from release time.

CONDITIONS OF REPORTING

- v The reported results are for information and interpretation of the referring doctor only.
- v It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- v Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for the same patient.
- v Should the results indicate an unexpected abnormality, the same should be reconfirmed.
- v Only such medical professionals who understand reporting units, reference ranges and limitations of technologies should interpret results.
- v This report is not valid for medico-legal purpose.
- v Neither Thyrocare, nor its employees/representatives assume: (a) any liability, responsibility for any loss or damage that may be incurred by any person as a result of presuming the meaning or contents of the report, (b) any claims of any nature whatsoever arising from or relating to the performance of the requested tests as well as any claim for indirect, incidental or consequential damages. The total liability, in any case, of Thyrocare shall not exceed the total amount of invoice for the services provided and paid for.
- v Thyrocare Discovery video link :- https://voutu.be/nbdYeRqYvOc

EXPLANATIONS

- v Majority of the specimen processed in the laboratory are collected by Pathologists and Hospitals we call them as "Clients".
- v Name The name is as declared by the client and recored by the personnel who collected the specimen.
- v Ref.Dr The name of the doctor who has recommended testing as declared by the client.
- v **Labcode** This is the accession number in our laboratory and it helps us in archiving and retrieving the data.
- v **Barcode** This is the specimen identity number and it states that the results are for the specimen bearing the barcode (irrespective of the name).
- v **SCP** Specimen Collection Point This is the location where the blood or specimen was collected as declared by the client.
- v SCT Specimen Collection Time The time when specimen was collected as declared by the client.
- v **SRT** Specimen Receiving Time This time when the specimen reached our laboratory.
- v **RRT** Report Releasing Time The time when our pathologist has released the values for Reporting.
- v Reference Range Means the range of values in which 95% of the normal population would fall.

SUGGESTIONS

- v Values out of reference range requires reconfirmation before starting any medical treatment.
- v Retesting is needed if you suspect any quality shortcomings.
- v Testing or retesting should be done in accredited laboratories.
- v For suggestions, complaints, clinical support or feedback, write to us at **customersupport@thyrocare.com** or call us on **022-3090 0000**



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