Thyrocare

Tests you can trust

Name : Bhakti Bhakti(34Y/F)

Date : 06 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#**

98% Reports released within **06 Hours** of sample reaching the lab+

9 out of 10 Doctors Trust that Thyrocare reports are **Accurate & Reliable***

1200+ Tests & Profiles

Temperature-Controlled Sample Logistics

Unique Barcode Tracking

Fully Automated Machines Inspected Daily

Abnormal Values Re-Checked Twice

Reports Verified By Expert MD Pathologists Stationed at Every Lab

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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

NABL From 2005

ISO 9001: 2015 – From 2015

CAP From 2007

PROCESSED AT :
Thyrocare
D-37/1,TTC MIDC,Turbhe,
Navi Mumbai-400 703

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Thyrocare Technologies Limited, D-37/3, TTC MIDC, Turbhe, Navi Mumbai - 400 703 9870666333 wellness@thyrocare.com

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NAME : BHAKTI BHAKTI(34Y/F)
REF. BY : SELF
TEST ASKED : MEDIWHEEL HEALTH CHECKUP BELOW 40

HOME COLLECTION :
H 306 Hawre Splendor Sector 20 home Opp Cidco garden Raigarh Maharashtra Cidco garden Raigarh Maharashtra

Report Availability Summary

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

13 Ready | **0** Ready with Cancellation | **0** Processing | **0** Cancelled in Lab

TEST DETAILS	REPORT STATUS
MEDIWHEEL 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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MC-2407

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Summary Report

Tests outside reference range

TEST NAME	OBSERVED VALUE	UNITS	Bio. Ref. Interval.
COMPLETE HEMOGRAM			
HEMOGLOBIN	11.6	g/dL	12.0-15.0
LYMPHOCYTE	43.5	%	20-40
LYMPHOCYTES - ABSOLUTE COUNT	3.03	X 10^3 / µL	1.0-3.0
MEAN CORP.HEMO.CONC(MCHC)	29.2	g/dL	31.5-34.5
MEAN CORPUSCULAR HEMOGLOBIN(MCH)	23.2	pq	27.0-32.0
MEAN CORPUSCULAR VOLUME(MCV)	79.6	fL	83.0-101.0
MONOCYTES - ABSOLUTE COUNT	0.17	X 10^3 / µL	0.2 - 1.0
PLATELET COUNT	440	X 10^3 / µL	150-410
PLATELETCRIT(PCT)	0.51	%	0.19-0.39
RED CELL DISTRIBUTION WIDTH (RDW-CV)	16	%	11.6-14.0
TOTAL RBC	4.99	X 10^6/µL	3.8-4.8
LIPID			
HDL CHOLESTEROL - DIRECT	25	mg/dL	40-60
TRIG / HDL RATIO	5.37	Ratio	< 3.12
RENAL			
BLOOD UREA NITROGEN (BUN)	5.09	mg/dL	7.94 - 20.07
BUN / SR.CREATININE RATIO	7.17	Ratio	9:1-23:1
UREA (CALCULATED)	10.89	mg/dL	Adult : 17-43
THYROID			
TSH - ULTRASENSITIVE	5.47	µIU/mL	0.54-5.30

Disclaimer: The above listed is the summary of the parameters with values outside the BRI. For detailed report values, parameter correlation and clinical interpretation, kindly refer to the same in subsequent pages.

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PATIENTID : BB25920114

TEST NAME	TECHNOLOGY	VALUE	UNITS
HbA1c - (HPLC)			
	H.P.L.C	5.2	%

Bio. Ref. Interval. :

Bio. Ref. Interval.: As per ADA Guidelines	Guidance For Known Diabetics
Below 5.7% : Normal 5.7% - 6.4% : Prediabetic >=6.5% : Diabetic	Below 6.5% : Good Control 6.5% - 7% : Fair Control 7.0% - 8% : Unsatisfactory Control >8% : Poor Control

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

AVERAGE BLOOD GLUCOSE (ABG)	CALCULATED	103	mg/dL

Bio. Ref. Interval. :

90 - 120 mg/dl : Good Control
121 - 150 mg/dl : Fair Control
151 - 180 mg/dl : Unsatisfactory Control
> 180 mg/dl : Poor Control

Method : Derived from HBA1c values

Please correlate with clinical conditions.

Sample Collected on (SCT) :06 Mar 2025 07:10
Sample Received on (SRT) : 06 Mar 2025 12:42
Report Released on (RRT) : 06 Mar 2025 16:27
Sample Type : EDTA Whole Blood
Labcode : 0603000334/DS853
Barcode : DM213939

Dr Renuka MD(Path)

Dr Arshiya MD(Path)

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

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TEST NAME	TECHNOLOGY	VALUE	UNITS
ERYTHROCYTE SEDIMENTATION RATE (ESR)	MODIFIED WESTERGREN	20	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.
Method:- MODIFIED WESTERGREN

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TEST NAME	METHODOLOGY	VALUE	UNITS	Bio. Ref. Interval.
HEMOGLOBIN	**SLS-Hemoglobin Method**	**11.6**	**g/dL**	**12.0-15.0**
Hematocrit (PCV)	CPH Detection	39.7	%	36.0-46.0
Total RBC	**HF & EI**	**4.99**	**X 10^6/µL**	**3.8-4.8**
Mean Corpuscular Volume (MCV)	**Calculated**	**79.6**	**fL**	**83.0-101.0**
Mean Corpuscular Hemoglobin (MCH)	**Calculated**	**23.2**	**pq**	**27.0-32.0**
Mean Corp.Hemo. Conc (MCHC)	**Calculated**	**29.2**	**g/dL**	**31.5-34.5**
Red Cell Distribution Width - SD (RDW-SD)	Calculated	45.4	fL	39.0-46.0
Red Cell Distribution Width (RDW - CV)	**Calculated**	**16**	**%**	**11.6-14.0**
RED CELL DISTRIBUTION WIDTH INDEX (RDWI)	Calculated	255.2	-	*Refer Note below
MENTZER INDEX	Calculated	16	-	*Refer Note below
TOTAL LEUCOCYTE COUNT (WBC)	HF & FC	6.97	X 10^3 / µL	4.0 - 10.0
DIFFERENTIAL LEUCOCYTE COUNT				
Neutrophils Percentage	Flow Cytometry	48.1	%	40-80
Lymphocytes Percentage	**Flow Cytometry**	**43.5**	**%**	**20-40**
Monocytes Percentage	Flow Cytometry	2.4	%	2-10
Eosinophils Percentage	Flow Cytometry	4.7	%	1-6
Basophils Percentage	Flow Cytometry	1	%	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	3.35	X 10^3 / µL	2.0-7.0
Lymphocytes - Absolute Count	**Calculated**	**3.03**	**X 10^3 / µL**	**1.0-3.0**
Monocytes - Absolute Count	**Calculated**	**0.17**	**X 10^3 / µL**	**0.2 - 1.0**
Basophils - Absolute Count	Calculated	0.07	X 10^3 / µL	0.02 - 0.1
Eosinophils - Absolute Count	Calculated	0.33	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**	**440**	**X 10^3 / µL**	**150-410**
Mean Platelet Volume (MPV)	Calculated	11.6	fL	6.5-12
Platelet Distribution Width (PDW)	Calculated	15.2	fL	9.6-15.2
Platelet to Large Cell Ratio (PLCR)	Calculated	39.6	%	19.7-42.4
Plateletcrit (PCT)	**Calculated**	**0.51**	**%**	**0.19-0.39**

Remarks : Alert!!! RBCs:Mild anisopoikilocytosis. Predominantly normocytic normochromic with microcytes & ovalocytes. Platelets:Appear adequate in smear.

***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.**
Method : Fully automated bidirectional analyser (6 Part Differential SYSMEX XN-1000)
(Reference : *FC- flowcytometry, *HF- hydrodynamic focussing, *EI- Electric Impedence, *Hb- hemoglobin, *CPH- Cumulative pulse height**)**

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Dr Arshiya MD(Path)

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TEST ASKED : MEDIWHEEL HEALTH CHECKUP BELOW 40

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TEST NAME	TECHNOLOGY	VALUE	UNITS
25-OH VITAMIN D (TOTAL)	E.C.L.I.A	38.5	ng/mL

Bio. Ref. Interval. :-

Deficiency : <=20 ng/ml || Insufficiency : 21-29 ng/ml
Sufficiency : >= 30 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):9.20%, Inter assay (%CV):8.50%
Kit Validation Reference : Holick M. Vtamin D the underappreciated D-Lightful hormone that is important for Skeletal and cellular health Curr Opin Endocrinol Diabetes 2002:9(1)87-98.

Please correlate with clinical conditions.
Method:- Fully Automated Electrochemiluminescence Compititive Immunoassay

Sample Collected on (SCT) : 06 Mar 2025 07:10
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Labcode : 0603000346/DS853
Barcode : DC137346

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TEST NAME	TECHNOLOGY	VALUE	UNITS
VITAMIN B-12	E.C.L.I.A	365	pg/mL

Bio. Ref. Interval. :-

Normal: 197-771 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):2.6%, Inter assay (%CV):2.3 %

Kit Validation Reference : Thomas L.Clinical laborator Diagnostics : Use and Assessment of Clinical laboratory Results 1st Edition,TH Books-Verl-Ges,1998:424-431

Please correlate with clinical conditions.
Method:- Fully Automated Electrochemiluminescence Compitive Immunoassay

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PATIENTID : BB25920114

TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
TOTAL CHOLESTEROL	PHOTOMETRY	104	mg/dL	< 200
HDL CHOLESTEROL - DIRECT	**PHOTOMETRY**	**25**	**mg/dL**	**40-60**
LDL CHOLESTEROL - DIRECT	PHOTOMETRY	57	mg/dL	< 100
TRIGLYCERIDES	PHOTOMETRY	134	mg/dL	< 150
TC/ HDL CHOLESTEROL RATIO	CALCULATED	4.2	Ratio	3 - 5
TRIG / HDL RATIO	**CALCULATED**	**5.37**	**Ratio**	**< 3.12**
LDL / HDL RATIO	CALCULATED	2.3	Ratio	1.5-3.5
HDL / LDL RATIO	CALCULATED	0.44	Ratio	> 0.40
NON-HDL CHOLESTEROL	CALCULATED	79.1	mg/dL	< 160
VLDL CHOLESTEROL	CALCULATED	26.84	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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TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
ALKALINE PHOSPHATASE	PHOTOMETRY	72.5	U/L	45-129
BILIRUBIN - TOTAL	PHOTOMETRY	0.46	mg/dL	0.3-1.2
BILIRUBIN -DIRECT	PHOTOMETRY	0.09	mg/dL	< 0.3
BILIRUBIN (INDIRECT)	CALCULATED	0.37	mg/dL	0-0.9
GAMMA GLUTAMYL TRANSFERASE (GGT)	PHOTOMETRY	14.6	U/L	< 38
ASPARTATE AMINOTRANSFERASE (SGOT)	PHOTOMETRY	25.8	U/L	< 31
ALANINE TRANSAMINASE (SGPT)	PHOTOMETRY	21.6	U/L	< 34
SGOT / SGPT RATIO	CALCULATED	1.19	Ratio	< 2
PROTEIN - TOTAL	PHOTOMETRY	7.13	gm/dL	5.7-8.2
ALBUMIN - SERUM	PHOTOMETRY	4.29	gm/dL	3.2-4.8
SERUM GLOBULIN	CALCULATED	2.84	gm/dL	2.5-3.4
SERUM ALB/GLOBULIN RATIO	CALCULATED	1.51	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 Bcg[1]method (Colorimetric Assay Endpoint)
SEGB - DERIVED FROM SERUM ALBUMIN AND PROTEIN VALUES
A/GR - Derived from serum Albumin and Protein values

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TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
CALCIUM	PHOTOMETRY	9	mg/dL	8.8-10.6
URIC ACID	PHOTOMETRY	4.9	mg/dL	3.2 - 6.1
BLOOD UREA NITROGEN (BUN)	**PHOTOMETRY**	**5.09**	**mg/dL**	**7.94 - 20.07**
UREA (CALCULATED)	**CALCULATED**	**10.89**	**mg/dL**	**Adult : 17-43**
CREATININE - SERUM	PHOTOMETRY	0.71	mg/dL	0.55-1.02
UREA / SR.CREATININE RATIO	CALCULATED	15.34	Ratio	< 52
BUN / SR.CREATININE RATIO	**CALCULATED**	**7.17**	**Ratio**	**9:1-23:1**
PHOSPHOROUS	PHOTOMETRY	3.87	mg/dL	2.4 - 5.1
SODIUM	I.S.E - INDIRECT	137.3	mmol/L	136 - 145
POTASSIUM	I.S.E - INDIRECT	5.05	mmol/L	3.5 - 5.1
CHLORIDE	I.S.E - INDIRECT	104.9	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

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PATIENTID : BB25920114

TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
TOTAL TRIIODOTHYRONINE (T3)	E.C.L.I.A	102	ng/dL	80-200
TOTAL THYROXINE (T4)	E.C.L.I.A	8.12	µg/dL	4.8-12.7
TSH - ULTRASENSITIVE	**E.C.L.I.A**	**5.47**	**µIU/mL**	**0.54-5.30**

Comments : IF NOT ON DRUGS SUGGESTED FT3 & FT4 ESTIMATION

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

Method :

T3,T4 - Fully Automated Electrochemiluminescence Compititive Immunoassay
USTSH - Fully Automated Electrochemiluminescence Sandwich Immunoassay

Pregnancy reference ranges for TSH/USTSH :
Trimester || T3 (ng/dl) || T4 (µg/dl) || TSH/USTSH (µIU/ml)
1st || 83.9-196.6 || 4.4-11.5 || 0.1-2.5
2nd || 86.1-217.4 || 4.9-12.2 || 0.2-3.0
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. 2019; 6(4)

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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Report Released on (RRT) : 06 Mar 2025 16:22
Sample Type : SERUM
Labcode : 0603000346/DS853
Barcode : DC137346

Dr Renuka MD(Path)

Dr Arshiya MD(Path)

NAME : BHAKTI BHAKTI(34Y/F)
REF. BY : SELF
TEST ASKED : MEDIWHEEL HEALTH CHECKUP BELOW 40

HOME COLLECTION :
H 306 Hawre Splendor Sector 20 home Opp Cidco garden Raigarh Maharashtra Cidco garden Raigarh Maharashtra

TEST NAME	TECHNOLOGY	VALUE	UNITS
EST. GLOMERULAR FILTRATION RATE (eGFR)	CALCULATED	114	mL/min/1.73 m2

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.
Method:- 2021 CKD EPI Creatinine Equation

Sample Collected on (SCT) : 06 Mar 2025 07:10
Sample Received on (SRT) : 06 Mar 2025 12:43
Report Released on (RRT) : 06 Mar 2025 16:22
Sample Type : SERUM
Labcode : 0603000346/DS853
Barcode : DC137346

Dr Renuka MD(Path)

Dr Arshiya MD(Path)

NAME : BHAKTI BHAKTI(34Y/F)
REF. BY : SELF
TEST ASKED : MEDIWHEEL HEALTH CHECKUP BELOW 40

HOME COLLECTION :
H 306 Hawre Splendor Sector 20 home Opp Cidco garden Raigarh Maharashtra Cidco garden Raigarh Maharashtra

PATIENTID : BB25920114

TEST NAME	METHODOLOGY	VALUE	UNITS	Bio. Ref. Interval.
Complete Urinogram				
Physical Examination				
VOLUME	Visual Determination	3	mL	-
COLOUR	Visual Determination	PALE YELLOW	-	Pale Yellow
APPEARANCE	Visual Determination	CLEAR	-	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	ABSENT	-	Absent
NITRITE	Diazo coupling	ABSENT	-	Absent
LEUCOCYTE ESTERASE	Esterase reaction	ABSENT	-	Absent
Microscopic Examination				
MUCUS	Microscopy	ABSENT	-	Absent
RED BLOOD CELLS	Microscopy	ABSENT	cells/HPF	0-5
URINARY LEUCOCYTES (PUS CELLS)	Microscopy	ABSENT	cells/HPF	0-5
EPITHELIAL CELLS	Microscopy	5	cells/HPF	0-5
CASTS	Microscopy	ABSENT	-	Absent
CRYSTALS	Microscopy	ABSENT	-	Absent
BACTERIA	Microscopy	ABSENT	-	Absent
YEAST	Microscopy	ABSENT	-	Absent
PARASITE	Microscopy	ABSENT	-	Absent

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

Sample Collected on (SCT) : 06 Mar 2025 07:10
Sample Received on (SRT) : 06 Mar 2025 12:32
Report Released on (RRT) : 06 Mar 2025 14:00
Sample Type : URINE
Labcode : 0603073136/DS853
Barcode : DK566717

Dr Renuka MD(Path)

Dr Arshiya MD(Path)

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Navi Mumbai-400 703

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NAME : BHAKTI BHAKTI(34Y/F)
REF. BY : SELF
TEST ASKED : MEDIWHEEL HEALTH CHECKUP BELOW 40

HOME COLLECTION :
H 306 Hawre Splendor Sector 20 home Opp Cidco garden Raigarh Maharashtra Cidco garden Raigarh Maharashtra

TEST NAME	TECHNOLOGY	VALUE	UNITS
FASTING BLOOD SUGAR(GLUCOSE)	PHOTOMETRY	90.14	mg/dL

Bio. Ref. Interval. :-

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

Note :
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.
Method:- GOD-PAP METHOD

~~ End of report ~~

Sample Collected on (SCT) : 06 Mar 2025 07:10
Sample Received on (SRT) : 06 Mar 2025 12:23
Report Released on (RRT) : 06 Mar 2025 13:27
Sample Type : FLUORIDE PLASMA
Labcode : 0603072387/DS853
Barcode : DJ138071

Dr Renuka MD(Path)

Dr Arshiya MD(Path)

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

CONDITIONS OF REPORTING

- The reported results are for information and interpretation of the referring doctor only.
- It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for the same patient.
- Should the results indicate an unexpected abnormality, the same should be reconfirmed.
- Only such medical professionals who understand reporting units, reference ranges and limitations of technologies should interpret results.
- This report is not valid for medico-legal purpose.
- 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.
- Thyrocare Discovery video link :- https://youtu.be/nbdYeRgYyQc

EXPLANATIONS

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

SUGGESTIONS

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

*+T&C Apply, #As on 5th December 2024, *As per a survey on doctors' perception of laboratory diagnostics (IJARIIT,2023)*