

Report for Shubham Tejani(21Y/M)

Tests asked Comprehensive Full Body Checkup With Vitamin D And B12 - New,  
Prot

Test date 26 Jun 2024

Report status Complete Report



**6** STEP

quality control to ensure 100% report accuracy



Qualified and trained technicians



Temperature-controlled containers to store samples



Strict quality checks on samples before processing



Regular monitoring of lab analyzers by experts



Assured machine inspection on a daily basis



Verified reports by qualified pathologists



25+ Years of Trust & Experience



NABL Accredited Labs



100+ Crore Samples Processed

**Name** : SHUBHAM TEJANI(21Y/M)  
**Ref. By** : SELF

**ADDRESS :**

B-13 ROOM NO 9 NEW SIDDHIVINAYAK CHS MG  
 COMPLEX SEC-14 VASHI NAVI MUMBAI

## Report Availability Summary

☒ Full Report Available

**Note** : This is summary page. Please refer to the table below for the details

Test	Report Status
<b>COMPREHENSIVE FULL BODY CHECKUP WITH VITAMIN D AND B12 - NEW</b>	<input checked="" type="checkbox"/> Available
25-OH VITAMIN D (TOTAL)	<input checked="" type="checkbox"/> Available
ALANINE TRANSAMINASE (SGPT)	<input checked="" type="checkbox"/> Available
ALBUMIN - SERUM	<input checked="" type="checkbox"/> Available
ALKALINE PHOSPHATASE	<input checked="" type="checkbox"/> Available
ASPARTATE AMINOTRANSFERASE (SGOT )	<input checked="" type="checkbox"/> Available
BILIRUBIN - TOTAL	<input checked="" type="checkbox"/> Available
BILIRUBIN -DIRECT	<input checked="" type="checkbox"/> Available
COMPLETE URINE ANALYSIS	<input checked="" type="checkbox"/> Available
FASTING BLOOD SUGAR(GLUCOSE)	<input checked="" type="checkbox"/> Available
GAMMA GLUTAMYL TRANSFERASE (GGT)	<input checked="" type="checkbox"/> Available
HbA1c	<input checked="" type="checkbox"/> Available
HEMOGRAM - 6 PART (DIFF)	<input checked="" type="checkbox"/> Available
IRON	<input checked="" type="checkbox"/> Available
KIDPRO	<input checked="" type="checkbox"/> Available
LIPID PROFILE	<input checked="" type="checkbox"/> Available
PROTEIN - TOTAL	<input checked="" type="checkbox"/> Available
TOTAL IRON BINDING CAPACITY (TIBC)	<input checked="" type="checkbox"/> Available
TOTAL THYROXINE (T4)	<input checked="" type="checkbox"/> Available

**Note** : Underlined values are Critical Values, Clinician's attention required.

**Clinically Tested by** : Thyrocare Technologies Ltd.

Name : SHUBHAM TEJANI(21Y/M)  
Ref. By : SELF

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Test	Report Status
TOTAL TRIIODOTHYRONINE (T3)	<input checked="" type="checkbox"/> Available
TSH - ULTRASENSITIVE	<input checked="" type="checkbox"/> Available
UNSAT.IRON-BINDING CAPACITY(UIBC)	<input checked="" type="checkbox"/> Available
VITAMIN B-12	<input checked="" type="checkbox"/> Available

**NAME** : SHUBHAM TEJANI(21Y/M)  
**REF. BY** : SELF  
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**HOME COLLECTION :**  
 B-13 ROOM NO 9 NEW SIDDHIVINAYAK CHS MG COMPLEX SEC-14 VASHI NAVI MUMBAI

TEST NAME	TECHNOLOGY	VALUE	UNITS
HbA1c - (HPLC)	H.P.L.C	4.5	%

**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  
 >8% : Poor Control

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

AVERAGE BLOOD GLUCOSE (ABG)	CALCULATED	<u>82</u>	mg/dL
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**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)** : 26 Jun 2024 09:54  
**Sample Received on (SRT)** : 26 Jun 2024 11:28  
**Report Released on (RRT)** : 26 Jun 2024 12:52  
**Sample Type** : EDTA Whole Blood  
**Labcode** : 2606071127/PE002  
**Barcode** : CM579843




Dr Sachin Patil MD(Path)

**NAME :** SHUBHAM TEJANI(21Y/M)  
**REF. BY :** SELF  
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**HOME COLLECTION :**  
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TEST NAME	METHODOLOGY	VALUE	UNITS	Bio. Ref. Interval.
TOTAL LEUCOCYTES COUNT (WBC)	HF & FC	5.51	X 10 <sup>3</sup> / µL	4.0 - 10.0
<b>NEUTROPHILS</b>	<b>Flow Cytometry</b>	<b><u>39.9</u></b>	<b>%</b>	<b>40-80</b>
<b>LYMPHOCYTE</b>	<b>Flow Cytometry</b>	<b><u>52.5</u></b>	<b>%</b>	<b>20-40</b>
<b>MONOCYTES</b>	<b>Flow Cytometry</b>	<b><u>1.8</u></b>	<b>%</b>	<b>2-10</b>
EOSINOPHILS	Flow Cytometry	5.1	%	1-6
BASOPHILS	Flow Cytometry	0.4	%	0-2
IMMATURE GRANULOCYTE PERCENTAGE(IG%)	Flow Cytometry	0.3	%	0-0.5
NEUTROPHILS - ABSOLUTE COUNT	Calculated	2.2	X 10 <sup>3</sup> / µL	2.0-7.0
LYMPHOCYTES - ABSOLUTE COUNT	Calculated	2.89	X 10 <sup>3</sup> / µL	1.0-3.0
<b>MONOCYTES - ABSOLUTE COUNT</b>	<b>Calculated</b>	<b><u>0.1</u></b>	<b>X 10<sup>3</sup> / µL</b>	<b>0.2 - 1.0</b>
BASOPHILS - ABSOLUTE COUNT	Calculated	0.02	X 10 <sup>3</sup> / µL	0.02 - 0.1
EOSINOPHILS - ABSOLUTE COUNT	Calculated	0.28	X 10 <sup>3</sup> / µL	0.02 - 0.5
IMMATURE GRANULOCYTES(IG)	Calculated	0.02	X 10 <sup>3</sup> / µL	0-0.3
<b>TOTAL RBC</b>	<b>HF &amp; EI</b>	<b><u>5.66</u></b>	<b>X 10<sup>6</sup>/µL</b>	<b>4.5-5.5</b>
NUCLEATED RED BLOOD CELLS	Calculated	0.01	X 10 <sup>3</sup> / µL	0.0-0.5
NUCLEATED RED BLOOD CELLS %	Flow Cytometry	0.01	%	0.0-5.0
HEMOGLOBIN	SLS-Hemoglobin Method	16.3	g/dL	13.0-17.0
HEMATOCRIT(PCV)	CPH Detection	47.2	%	40.0-50.0
MEAN CORPUSCULAR VOLUME(MCV)	Calculated	83.4	fL	83.0-101.0
MEAN CORPUSCULAR HEMOGLOBIN(MCH)	Calculated	28.8	pq	27.0-32.0
MEAN CORP.HEMO.CONC(MCHC)	Calculated	34.5	g/dL	31.5-34.5
<b>RED CELL DISTRIBUTION WIDTH - SD(RDW-SD)</b>	<b>Calculated</b>	<b><u>37</u></b>	<b>fL</b>	<b>39-46</b>
RED CELL DISTRIBUTION WIDTH (RDW-CV)	Calculated	12.2	%	11.6-14
<b>PLATELET DISTRIBUTION WIDTH(PDW)</b>	<b>Calculated</b>	<b><u>8.1</u></b>	<b>fL</b>	<b>9.6-15.2</b>
MEAN PLATELET VOLUME(MPV)	Calculated	8.3	fL	6.5-12
PLATELET COUNT	HF & EI	319	X 10 <sup>3</sup> / µL	150-410
<b>PLATELET TO LARGE CELL RATIO(PLCR)</b>	<b>Calculated</b>	<b><u>12.4</u></b>	<b>%</b>	<b>19.7-42.4</b>
PLATELETCRIT(PCT)	Calculated	0.27	%	0.19-0.39

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

**Please Correlate with clinical conditions.**

**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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D AND B12 - NEW,PROTEIN

**HOME COLLECTION :**  
B-13 ROOM NO 9 NEW SIDDHIVINAYAK CHS MG  
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TEST NAME	TECHNOLOGY	VALUE	UNITS
FASTING BLOOD SUGAR(GLUCOSE)	PHOTOMETRY	<u>101.09</u>	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

**Sample Collected on (SCT)** : 26 Jun 2024 09:54  
**Sample Received on (SRT)** : 26 Jun 2024 11:30  
**Report Released on (RRT)** : 26 Jun 2024 14:35  
**Sample Type** : FLUORIDE  
**Labcode** : 2606071305/PE002  
**Barcode** : CK513049



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TEST NAME	TECHNOLOGY	VALUE	UNITS
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**25-OH VITAMIN D (TOTAL)** **E.C.L.I.A** **6.46** **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. Vitamin D the underappreciated D-Lightful hormone that is important for Skeletal and cellular health Curr Opin Endocrinol Diabetes 2002;9(1)87-98.

**Method :** Fully Automated Electrochemiluminescence Competitive Immunoassay

**VITAMIN B-12** **E.C.L.I.A** **> 100** **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

**Method :** Fully Automated Electrochemiluminescence Competitive Immunoassay

**Please correlate with clinical conditions.**

**Sample Collected on (SCT)** : 26 Jun 2024 09:54  
**Sample Received on (SRT)** : 26 Jun 2024 11:36  
**Report Released on (RRT)** : 26 Jun 2024 17:58  
**Sample Type** : SERUM  
**Labcode** : 2606071893/PE002  
**Barcode** : CF749448



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TEST NAME	TECHNOLOGY	VALUE	UNITS
IRON	PHOTOMETRY	159.7	µg/dL
<b>Bio. Ref. Interval. :</b> Male : 65 - 175 Female : 50 - 170 <b>Method :</b> Ferrozine method without deproteinization			
TOTAL IRON BINDING CAPACITY (TIBC)	PHOTOMETRY	355.9	µg/dL
<b>Bio. Ref. Interval. :</b> Male: 225 - 535 µg/dl Female: 215 - 535 µg/dl <b>Method :</b> Spectrophotometric Assay			
% TRANSFERRIN SATURATION	CALCULATED	44.87	%
<b>Bio. Ref. Interval. :</b> 13 - 45 <b>Method :</b> Derived from IRON and TIBC values			
UNSAT.IRON-BINDING CAPACITY(UIBC)	PHOTOMETRY	196.2	µg/dL
<b>Bio. Ref. Interval. :</b> 162 - 368 <b>Method :</b> SPECTROPHOTOMETRIC ASSAY			

**Please correlate with clinical conditions.**

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TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
TOTAL CHOLESTEROL	PHOTOMETRY	144	mg/dL	< 200
<b>HDL CHOLESTEROL - DIRECT</b>	<b>PHOTOMETRY</b>	<b><u>32</u></b>	<b>mg/dL</b>	<b>40-60</b>
<b>HDL / LDL RATIO</b>	<b>CALCULATED</b>	<b><u>0.32</u></b>	<b>Ratio</b>	<b>&gt; 0.40</b>
LDL CHOLESTEROL - DIRECT	PHOTOMETRY	98	mg/dL	< 100
TC/ HDL CHOLESTEROL RATIO	CALCULATED	4.5	Ratio	3 - 5
TRIG / HDL RATIO	CALCULATED	2.75	Ratio	< 3.12
TRIGLYCERIDES	PHOTOMETRY	88	mg/dL	< 150
LDL / HDL RATIO	CALCULATED	3.1	Ratio	1.5-3.5
NON-HDL CHOLESTEROL	CALCULATED	111.7	mg/dL	< 160
VLDL CHOLESTEROL	CALCULATED	17.5	mg/dL	5 - 40

**Please correlate with clinical conditions.**

**Method :**

CHOL - Cholesterol Oxidase, Esterase, Peroxidase  
 HCHO - Direct Enzymatic Colorimetric  
 HD/LD - Derived from HDL and LDL values.  
 LDL - Direct Measure  
 TC/H - Derived from serum Cholesterol and Hdl values  
 TRI/H - Derived from TRIG and HDL Values  
 TRIG - Enzymatic, End Point  
 LDL/ - Derived from serum 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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**HOME COLLECTION :**  
 B-13 ROOM NO 9 NEW SIDDHIVINAYAK CHS MG  
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TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
ALKALINE PHOSPHATASE	PHOTOMETRY	103.6	U/L	45-129
BILIRUBIN - TOTAL	PHOTOMETRY	1.07	mg/dL	0.3-1.2
BILIRUBIN -DIRECT	PHOTOMETRY	0.2	mg/dL	< 0.3
BILIRUBIN (INDIRECT)	CALCULATED	0.87	mg/dL	0-0.9
GAMMA GLUTAMYL TRANSFERASE (GGT)	PHOTOMETRY	13.9	U/L	< 55
SGOT / SGPT RATIO	CALCULATED	1.78	Ratio	< 2
ASPARTATE AMINOTRANSFERASE (SGOT )	PHOTOMETRY	18.9	U/L	< 35
ALANINE TRANSAMINASE (SGPT)	PHOTOMETRY	10.6	U/L	< 45
PROTEIN - TOTAL	PHOTOMETRY	7.01	gm/dL	5.7-8.2
ALBUMIN - SERUM	PHOTOMETRY	4.46	gm/dL	3.2-4.8
SERUM GLOBULIN	CALCULATED	2.55	gm/dL	2.5-3.4
SERUM ALB/GLOBULIN RATIO	CALCULATED	1.75	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  
 OT/PT - Derived from SGOT and SGPT values.  
 SGOT - IFCC\* Without Pyridoxal Phosphate Activation  
 SGPT - IFCC\* Without Pyridoxal Phosphate Activation  
 PROT - Biuret Method  
 SALB - Albumin Bcg<sup>1</sup>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.
<b>UREA (CALCULATED)</b>	<b>CALCULATED</b>	<b><u>14.98</u></b>	<b>mg/dL</b>	<b>Adult : 17-43</b>
<b>BLOOD UREA NITROGEN (BUN)</b>	<b>PHOTOMETRY</b>	<b><u>7</u></b>	<b>mg/dL</b>	<b>7.94 - 20.07</b>
UREA / SR.CREATININE RATIO	CALCULATED	20.52	Ratio	< 52
CREATININE - SERUM	PHOTOMETRY	0.73	mg/dL	0.72-1.18
BUN / SR.CREATININE RATIO	CALCULATED	9.59	Ratio	9:1-23:1
CALCIUM	PHOTOMETRY	9.2	mg/dL	8.8-10.6
<b>URIC ACID</b>	<b>PHOTOMETRY</b>	<b><u>7.7</u></b>	<b>mg/dL</b>	<b>4.2 - 7.3</b>

**Please correlate with clinical conditions.**

**Method :**

UREAC - Derived from BUN Value.  
 BUN - Kinetic UV Assay.  
 UR/CR - Derived from UREA and Sr.Creatinine values.  
 SCRE - Creatinine Enzymatic Method  
 B/CR - Derived from serum Bun and Creatinine values  
 CALC - Arsenazo III Method, End Point.  
 URIC - Uricase / Peroxidase Method

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TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
TOTAL TRIIODOTHYRONINE (T3)	E.C.L.I.A	106	ng/dL	80-200
TOTAL THYROXINE (T4)	E.C.L.I.A	10.4	µg/dL	4.8-12.7
TSH - ULTRASENSITIVE	E.C.L.I.A	2.92	µIU/mL	0.54-5.30

**Comments :** SUGGESTING THYRONORMALCY

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

**Method :**

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

**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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TEST NAME	TECHNOLOGY	VALUE	UNITS
EST. GLOMERULAR FILTRATION RATE (eGFR)	CALCULATED	133	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:-** CKD-EPI Creatinine Equation

**Sample Collected on (SCT)** : 26 Jun 2024 09:54  
**Sample Received on (SRT)** : 26 Jun 2024 11:36  
**Report Released on (RRT)** : 26 Jun 2024 17:58  
**Sample Type** : SERUM  
**Labcode** : 2606071893/PE002  
**Barcode** : CF749448



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**Note:- Underlined values are Critical Values, Clinician's attention required.**

**Clinically Tested by :Thyrocare Technologies Ltd - (NABL accredited)**

**NAME** : SHUBHAM TEJANI(21Y/M)  
**REF. BY** : SELF  
**TEST ASKED** : COMPREHENSIVE FULL BODY CHECKUP WITH VITAMIN D AND B12 - NEW,PROTEIN

**HOME COLLECTION :**  
 B-13 ROOM NO 9 NEW SIDDHIVINAYAK CHS MG  
 COMPLEX SEC-14 VASHI NAVI MUMBAI

TEST NAME	METHODOLOGY	VALUE	UNITS	Bio. Ref. Interval.
<b>Complete Urinogram</b>				
<b><u>Physical Examination</u></b>				
VOLUME	Visual Determination	3	mL	-
COLOUR	Visual Determination	PALE YELLOW	-	Pale Yellow
APPEARANCE	Visual Determination	CLEAR	-	Clear
SPECIFIC GRAVITY	pKa change	1.025	-	1.003-1.030
PH	pH indicator	5.5	-	5-8
<b><u>Chemical Examination</u></b>				
<b>URINARY PROTEIN</b>	<b>PEI</b>	<b>Trace (15-30 mg/dl)</b>	<b>mg/dL</b>	<b>Absent</b>
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
<b><u>Microscopic Examination</u></b>				
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	ABSENT	cells/HPF	0-5
CASTS	Microscopy	ABSENT	-	Absent
<b>CRYSTALS</b>	<b>Microscopy</b>	<b>CALCIUM OXALATE CRYSTALS</b>	-	<b>Absent</b>
BACTERIA	Microscopy	ABSENT	-	Absent
YEAST	Microscopy	ABSENT	-	Absent
PARASITE	Microscopy	ABSENT	-	Absent

**Remarks** : Alert!!! Calcium oxalate crystals seen/ HPF.

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

~~ End of report ~~

**Sample Collected on (SCT)** : 26 Jun 2024 09:54  
**Sample Received on (SRT)** : 26 Jun 2024 11:31  
**Report Released on (RRT)** : 26 Jun 2024 16:45  
**Sample Type** : URINE  
**Labcode** : 2606071499/PE002  
**Barcode** : BM069042



  
 Dr Sachin Patil MD(Path)

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Note:- Underlined values are Critical Values, Clinician's attention required.

Clinically Tested by :Thyrocare Technologies Ltd

#### CONDITIONS OF REPORTING

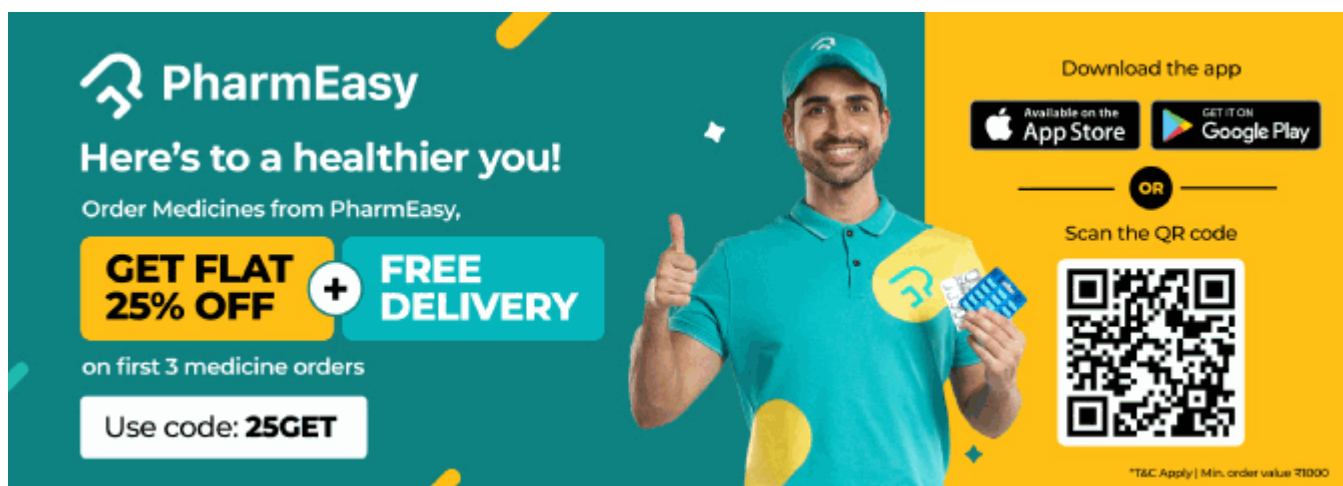
- ✓ The reported results are for information and interpretation of the referring doctor only.
- ✓ It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- ✓ Results of tests may vary from laboratory to laboratory and also in some parameters from time to time 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.
- ✓ Docon Technologies Private Limited, Thyrocare Technologies Limited and its employees/representatives do not assume 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.

#### EXPLANATIONS

- ✓ **Name** - The name is as declared by the client and recorded by the personnel who collected the specimen.
- ✓ **Ref.By** - 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).
- ✓ **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.
- ✓ For suggestions, complaints or feedback, write to us at [grievance-office@docon.co.in](mailto:grievance-office@docon.co.in) or call us on 7022000900.



The advertisement banner for PharmEasy features a smiling male delivery person in a blue uniform and cap, giving a thumbs up while holding a medicine box. The background is split into teal and yellow sections. On the teal side, the PharmEasy logo is at the top, followed by the slogan 'Here's to a healthier you!' and 'Order Medicines from PharmEasy,'. A large yellow button highlights 'GET FLAT 25% OFF + FREE DELIVERY' on first 3 medicine orders, with a code box below showing 'Use code: 25GET'. On the yellow side, it prompts to 'Download the app' with 'Available on the App Store' and 'GET IT ON Google Play' buttons, separated by an 'OR' circle. Below this is a 'Scan the QR code' section with a QR code. A small disclaimer at the bottom right states '\*T&C Apply | Min. order value ₹1000'.