

**PROCESSED AT :****Thyrocare**

260 - 261, Tribhuvan Complex,  
Ishwar Nagar,  
New Delhi - 110 065



Tests you can trust

**Corporate office : Thyrocare Technologies Limited, D-37/3, TTC MIDC, Turbhe, Navi Mumbai - 400 703**

☎ 022 - 3090 0000 / 6712 3400 📞 9870666333 ✉ wellness@thyrocare.com 🌐 www.thyrocare.com

**REPORT****NAME** : VIVEK KUMAR (32Y/M)**REF. BY** : HCL**TEST ASKED** : BLOOD SUGAR (F), COMPLETE URINE ANALYSIS, HbA1c**PATIENTID** : VK21292241**MOBILE NO** : 8750200587**DOB** : 06/22/1990**SAMPLE COLLECTED AT :**

HCL HEALTHCARE NOIDA SEZ CORPORATE  
CENTER,  
HCL HEALTHCARE, GATE 4, PLOT 3A,  
SECTOR-126, NOIDA, UP - 201301,  
INDIA-201301

**PAN ID** : HN1-20397

| TEST NAME      | TECHNOLOGY | VALUE | UNITS |
|----------------|------------|-------|-------|
| HbA1c - (HPLC) | H.P.L.C    | 5.6   | %     |

**Reference Range :****Reference Range: 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 114 mg/dl**Reference Range :**

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)** : 14 Mar 2023 08:00**Sample Received on (SRT)** : 14 Mar 2023 17:32**Report Released on (RRT)** : 14 Mar 2023 20:13**Sample Type** : EDTA**Labcode** : 1403101022/HCL04**Barcode** : AR240005

Dr Neha Prabhakar MD(Path)

Dr V Sandeep MD(Path)

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**REPORT**

**NAME** : VIVEK KUMAR (32Y/M)  
**REF. BY** : HCL  
**TEST ASKED** : BLOOD SUGAR (F), COMPLETE URINE ANALYSIS, HbA1c  
**PATIENTID** : VK21292241  
**MOBILE NO** : 8750200587  
**DOB** : 06/22/1990

**SAMPLE COLLECTED AT :**  
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HCL HEALTHCARE, GATE 4, PLOT 3A,  
SECTOR-126, NOIDA, UP - 201301,  
**PAN ID** : HN1-20397

| TEST NAME                                   | VALUE       | UNITS                                       | REFERENCE RANGE  |
|---|-------------|---|------------------|
| HEMOGLOBIN                                  | 13.8        | g/dL  | 13-17            |
| HEMATOCRIT(PCV)                             | 44.3        | %   | 40-50            |
| TOTAL RBC                                   | 4.9         | $\times 10^6/\mu\text{L}$                   | 4.5-5.5          |
| MEAN CORPUSCULAR VOLUME(MCV)                | 90.4        | fL  | 83-101           |
| MEAN CORPUSCULAR HEMOGLOBIN(MCH)            | 28.2        | pg  | 27-32            |
| <b>MEAN CORP.HEMO.CONC(MCHC)</b>            | <b>31.2</b> | <b>g/dL</b>                                 | <b>31.5-34.5</b> |
| RED CELL DISTRIBUTION WIDTH (RDW-CV)        | 13.5        | %   | 11.6-14          |
| TOTAL LEUCOCYTES COUNT (WBC)                | 5.12        | $\times 10^3/\mu\text{L}$                   | 4.0-10.0         |
| NEUTROPHILS                                 | 64          | %   | 40-80            |
| LYMPHOCYTE PERCENTAGE                       | 26.2        | %   | 20-40            |
| EOSINOPHILS                                 | 3.5         | %   | 0.0-6.0          |
| MONOCYTES                                   | 4.9         | %   | 0-10             |
| BASOPHILS                                   | 0.4         | %   | <2               |
| NEUTROPHILS - ABSOLUTE COUNT                | 3.28        | $\times 10^3/\mu\text{L}$                   | 2.0-7.0          |
| LYMPHOCYTES - ABSOLUTE COUNT                | 1.34        | $\times 10^3/\mu\text{L}$                   | 1.0-3.0          |
| EOSINOPHILS - ABSOLUTE COUNT                | 0.18        | $\times 10^3/\mu\text{L}$                   | 0-0.5            |
| MONOCYTES - ABSOLUTE COUNT                  | 0.25        | $\times 10^3/\mu\text{L}$                   | 0.2-1            |
| BASOPHILS - ABSOLUTE COUNT                  | 0.02        | $\times 10^3/\mu\text{L}$                   | 0-0.1            |
| <b>PLATELET COUNT</b>                       | <b>120</b>  | <b><math>\times 10^3/\mu\text{L}</math></b> | <b>150-400</b>   |
| IMMATURE GRANULOCYTES(IG)                   | 0.05        | $\times 10^3/\mu\text{L}$                   | 0-0.3            |
| <b>IMMATURE GRANULOCYTE PERCENTAGE(IG%)</b> | <b>1</b>    | <b>%</b>                                    | <b>0-0.5</b>     |
| NUCLEATED RED BLOOD CELLS                   | Nil         | $\times 10^3/\mu\text{L}$                   | <0.01            |
| <b>NUCLEATED RED BLOOD CELLS %</b>          | <b>Nil</b>  | <b>%</b>                                    | <b>&lt;0.01</b>  |

Please Correlate with clinical conditions.

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

(This device performs hematology analyses according to the Hydrodynamic Focussing (DC method), Flow Cytometry Method (using a semiconductor laser), and SLS- hemoglobin method)

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**Report Released on (RRT)** : 14 Mar 2023 20:13  
**Sample Type** : EDTA  
**Labcode** : 1403101022/HCL04  
**Barcode** : AR240005

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**REPORT**

**NAME** : VIVEK KUMAR (32Y/M)  
**REF. BY** : HCL  
**TEST ASKED** : BLOOD SUGAR (F), COMPLETE URINE  
ANALYSIS, HbA1c  
**PATIENTID** : VK21292241  
**MOBILE NO** : 8750200587  
**DOB** : 06/22/1990

**SAMPLE COLLECTED AT :**  
HCL HEALTHCARE NOIDA SEZ CORPORATE  
CENTER,  
HCL HEALTHCARE, GATE 4, PLOT 3A, SECTOR-126,  
NOIDA, UP - 201301, INDIA-201301

**PAN ID** : HN1-20397

| TEST NAME                     | TECHNOLOGY | VALUE  | UNITS |
|-------------------------------|------------|--------|-------|
| FASTING BLOOD SUGAR (GLUCOSE) | PHOTOMETRY | 110.18 | mg/dL |

**Reference Range :-**

| 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)** : 14 Mar 2023 08:00  
**Sample Received on (SRT)** : 14 Mar 2023 17:35  
**Report Released on (RRT)** : 14 Mar 2023 21:38  
**Sample Type** : FLUORIDE  
**Labcode** : 1403101279/HCL04  
**Barcode** : AR198230

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**REPORT**

**NAME** : VIVEK KUMAR (32Y/M)  
**REF. BY** : HCL  
**TEST ASKED** : COMPLETE URINE ANALYSIS

**SAMPLE COLLECTED AT :**  
HCL HEALTHCARE NOIDA SEZ CORPORATE  
CENTER,  
HCL HEALTHCARE, GATE 4, PLOT 3A, SECTOR-126,  
NOIDA, UP - 201301, INDIA-201301

**PATIENTID** : VK21292241  
**MOBILE NO** : 8750200587  
**DOB** : 06/22/1990

**PAN ID** : HN1-20397

| TEST NAME                      | OBSERVATION | UNITS     | REFERENCE RANGE |
|--------------------------------|-------------|-----------|-----------------|
| <b>Complete Urinogram</b>      |             |           |                 |
| <b>Physical Examination</b>    |             |           |                 |
| VOLUME                         | 3           | mL        | -               |
| COLOUR                         | PALE YELLOW | -         | Pale Yellow     |
| APPEARANCE                     | CLEAR       | -         | Clear           |
| SPECIFIC GRAVITY               | < 1.003     | -         | 1.003-1.030     |
| PH                             | 6.5         | -         | 5 - 8           |
| <b>Chemical Examination</b>    |             |           |                 |
| URINARY PROTEIN                | ABSENT      | mg/dl     | Absent          |
| URINARY GLUCOSE                | ABSENT      | mg/dl     | Absent          |
| URINE KETONE                   | ABSENT      | mg/dl     | Absent          |
| URINARY BILIRUBIN              | ABSENT      | mg/dl     | Absent          |
| UROBILINOGEN                   | Normal      | mg/dl     | <=0.2           |
| BILE SALT                      | ABSENT      | -         | Absent          |
| BILE PIGMENT                   | ABSENT      | -         | Absent          |
| URINE BLOOD                    | ABSENT      | -         | Absent          |
| NITRITE                        | ABSENT      | -         | Absent          |
| MICROALBUMIN                   | 10          | mg/l      | < 30            |
| <b>Microscopic Examination</b> |             |           |                 |
| MUCUS                          | PRESENT     | -         | Absent          |
| RED BLOOD CELLS                | ABSENT      | Cells/HPF | 0-5             |
| URINARY LEUCOCYTES (PUS CELLS) | ABSENT      | Cells/HPF | 0-5             |
| EPITHELIAL CELLS               | 2           | Cells/HPF | 0-5             |
| CASTS                          | ABSENT      | -         | Absent          |
| CRYSTALS                       | ABSENT      | -         | Absent          |
| BACTERIA                       | ABSENT      | -         | Absent          |
| YEAST                          | ABSENT      | -         | Absent          |
| PARASITE                       | ABSENT      | -         | Absent          |

**Method :** Fully Automated Matrix AVE Urinalysis Dipstick Method, Microscopy

**Sample Collected on (SCT)** : 14 Mar 2023 08:00  
**Sample Received on (SRT)** : 14 Mar 2023 19:03  
**Report Released on (RRT)** : 14 Mar 2023 20:06  
**Sample Type** : URINE  
**Labcode** : 1403107246/HCL04  
**Barcode** : AR710628

Dr Neha Prabhakar MD(Path)

Dr V Sandeep MD(Path)



**PROCESSED AT :**

**Thyrocare,**  
Plot No.428,Phase-IV,  
Udyog Vihar,  
Gurgaon,Haryana - 122 015



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**REPORT**

**NAME** : VIVEK KUMAR (32Y/M)  
**REF. BY** : HCL  
**TEST ASKED** : BLOOD SUGAR (F),COMPLETE URINE ANALYSIS,HbA1c  
**PATIENTID** : VK21292241  
**MOBILE NO** : 8750200587  
**DOB** : 06/22/1990

**SAMPLE COLLECTED AT :**  
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**PAN ID** : HN1-20397

| TEST NAME                       | TECHNOLOGY | VALUE | UNITS |
|---------------------------------|------------|-------|-------|
| PROSTATE SPECIFIC ANTIGEN (PSA) | C.L.I.A    | 0.26  | ng/ml |

**Reference Range :-**

Normal : < 4.00 ng/ml  
Border line : 4.01 to 10.00 ng/ml

**Clinical Significance:**

Elevated levels of PSA are associated with prostate cancer, but may also be seen with prostatitis (Inflammation of the prostate) and benign prostatic hyperplasia (BPH). PSA test done along with free PSA provides additional information. Studies have suggested that the percentage of free PSA in total PSA is lower in patients with prostate cancer than those with benign prostate hyperplasia.

**Specification:**

Precision: Intra assay (%CV): 4.38%, Inter assay (%CV): 4.67%; Sensitivity: 0.01 ng/ml

**Kit validation references:**

Wang MC, Valenzuela LA, Murphy GP, and Chu TM. Purification of a human prostate-specific antigen. Invest. Urol. 1979; 17: 159

**Please correlate with clinical conditions.**

**Method:-** TWO SITE SANDWICH IMMUNOASSAY

**Sample Collected on (SCT)** : 14 Mar 2023 08:00  
**Sample Received on (SRT)** : 15 Mar 2023 03:17  
**Report Released on (RRT)** : 15 Mar 2023 06:32  
**Sample Type** : SERUM  
**Labcode** : 1403134268/HCL04  
**Barcode** : AR559203

Dr Saakshi Mittal MD(Path)

Dr Preet Kaur MD(Path)

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**PAN ID** : HN1-20397

| TEST NAME                       | TECHNOLOGY        | VALUE       | UNITS        | NORMAL RANGE     |
|---------------------------------|-------------------|-------------|--------------|------------------|
| TOTAL CHOLESTEROL               | PHOTOMETRY        | 191         | mg/dl        | < 200            |
| HDL CHOLESTEROL - DIRECT        | PHOTOMETRY        | 41          | mg/dl        | 40-60            |
| <b>HDL / LDL RATIO</b>          | <b>CALCULATED</b> | <b>0.32</b> | <b>Ratio</b> | <b>&gt; 0.40</b> |
| <b>LDL CHOLESTEROL - DIRECT</b> | <b>PHOTOMETRY</b> | <b>130</b>  | <b>mg/dl</b> | <b>&lt; 100</b>  |
| <b>TRIG / HDL RATIO</b>         | <b>CALCULATED</b> | <b>3.51</b> | <b>Ratio</b> | <b>&lt; 3.12</b> |
| TRIGLYCERIDES                   | PHOTOMETRY        | 146         | mg/dl        | < 150            |
| TC/ HDL CHOLESTEROL RATIO       | CALCULATED        | 4.6         | Ratio        | 3 - 5            |
| LDL / HDL RATIO                 | CALCULATED        | 3.1         | Ratio        | 1.5-3.5          |
| VLDL CHOLESTEROL                | CALCULATED        | 29.1        | 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  
TRI/H - DERIVED FROM TRIG AND HDL VALUES  
TRIG - ENZYMATIC, END POINT  
TC/H - DERIVED FROM SERUM CHOLESTEROL AND HDL VALUES  
LDL/ - DERIVED FROM SERUM HDL AND LDL 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)** : 15 Mar 2023 03:17  
**Report Released on (RRT)** : 15 Mar 2023 06:32  
**Sample Type** : SERUM  
**Labcode** : 1403134268/HCL04  
**Barcode** : AR559203

*Saakshi*

Dr Saakshi Mittal MD(Path)

*Preet Kaur*

Dr Preet Kaur MD(Path)

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**PATIENTID** : VK21292241  
**MOBILE NO** : 8750200587  
**DOB** : 06/22/1990

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NOIDA, UP - 201301, INDIA-201301

**PAN ID** : HN1-20397

| TEST NAME                          | TECHNOLOGY        | VALUE        | UNITS        | NORMAL RANGE   |
|------------------------------------|-------------------|--------------|--------------|----------------|
| ALKALINE PHOSPHATASE               | PHOTOMETRY        | 79.11        | U/L          | 45-129         |
| BILIRUBIN - TOTAL                  | PHOTOMETRY        | 0.65         | mg/dl        | 0.3-1.2        |
| BILIRUBIN -DIRECT                  | PHOTOMETRY        | 0.19         | mg/dl        | < 0.3          |
| BILIRUBIN (INDIRECT)               | CALCULATED        | 0.46         | mg/dl        | 0-0.9          |
| SGOT / SGPT RATIO                  | CALCULATED        | 0.56         | Ratio        | < 2            |
| ASPARTATE AMINOTRANSFERASE (SGOT ) | PHOTOMETRY        | 31.54        | U/l          | < 35           |
| <b>ALANINE TRANSAMINASE (SGPT)</b> | <b>PHOTOMETRY</b> | <b>56.48</b> | <b>U/l</b>   | <b>&lt; 45</b> |
| PROTEIN - TOTAL                    | PHOTOMETRY        | 7.49         | gm/dl        | 5.7-8.2        |
| <b>ALBUMIN - SERUM</b>             | <b>PHOTOMETRY</b> | <b>5.13</b>  | <b>gm/dl</b> | <b>3.2-4.8</b> |
| <b>SERUM GLOBULIN</b>              | <b>CALCULATED</b> | <b>2.36</b>  | <b>gm/dL</b> | <b>2.5-3.4</b> |
| <b>SERUM ALB/GLOBULIN RATIO</b>    | <b>CALCULATED</b> | <b>2.17</b>  | <b>Ratio</b> | <b>0.9 - 2</b> |

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  
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\*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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*Saakshi*

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**REPORT****NAME** : VIVEK KUMAR (32Y/M)**REF. BY** : HCL**TEST ASKED** : BLOOD SUGAR (F),COMPLETE URINE ANALYSIS,HbA1c**SAMPLE COLLECTED AT :**

HCL HEALTHCARE NOIDA SEZ CORPORATE CENTER,

HCL HEALTHCARE,GATE 4, PLOT 3A, SECTOR-126,

NOIDA, UP - 201301, INDIA-201301

**PATIENTID** : VK21292241**MOBILE NO** : 8750200587**DOB** : 06/22/1990**PAN ID** : HN1-20397

| TEST NAME                         | TECHNOLOGY | VALUE | UNITS  | REFERENCE RANGE |
|-----------------------------------|------------|-------|--------|-----------------|
| THYROID STIMULATING HORMONE (TSH) | C.L.I.A    | 3.96  | μIU/ml | 0.3-5.5         |
| FREE TRIIODOTHYRONINE (FT3)       | C.L.I.A    | 3.35  | pg/ml  | 1.7-4.2         |
| FREE THYROXINE (FT4)              | C.L.I.A    | 1.29  | ng/dl  | 0.7-1.8         |

**Comments :** SUGGESTING THYRONORMALCY**Please correlate with clinical conditions.****Method :**

TSH - Sandwich Chemi Luminescent Immuno Assay

FT3 - Competitive Chemi Luminescent Immuno Assay

FT4 - Competitive Chemi Luminescent Immuno Assay

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**REF. BY** : HCL  
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**PATIENTID** : VK21292241  
**MOBILE NO** : 8750200587  
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NOIDA, UP - 201301, INDIA-201301

**PAN ID** : HN1-20397

| TEST NAME                  | TECHNOLOGY | VALUE | UNITS | NORMAL RANGE  |
|----------------------------|------------|-------|-------|---------------|
| CALCIUM                    | PHOTOMETRY | 10    | mg/dl | 8.8-10.6      |
| URIC ACID                  | PHOTOMETRY | 8.3   | mg/dl | 4.2 - 7.3     |
| UREA (CALCULATED)          | CALCULATED | 20.97 | mg/dL | Adult : 17-43 |
| BLOOD UREA NITROGEN (BUN)  | PHOTOMETRY | 9.8   | mg/dL | 7.04-20.07    |
| UREA / SR.CREATININE RATIO | CALCULATED | 36.79 | Ratio | < 52          |
| CREATININE - SERUM         | PHOTOMETRY | 0.57  | mg/dl | 0.72-1.18     |
| BUN / SR.CREATININE RATIO  | CALCULATED | 17.19 | Ratio | 9:1-23:1      |

**Please correlate with clinical conditions.**

**Method :**

CALC - ARSENazo III METHOD, END POINT.  
URIC - URICASE / PEROXIDASE 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

**Sample Collected on (SCT)** : 14 Mar 2023 08:00  
**Sample Received on (SRT)** : 15 Mar 2023 03:17  
**Report Released on (RRT)** : 15 Mar 2023 06:32  
**Sample Type** : SERUM  
**Labcode** : 1403134268/HCL04  
**Barcode** : AR559203

*Saakshi*

Dr Saakshi Mittal MD(Path)

*Preet Kaur*

Dr Preet Kaur MD(Path)

**PROCESSED AT :**

**Thyrocare,**  
Plot No.428,Phase-IV,  
Udyog Vihar,  
Gurgaon,Haryana - 122 015



Tests you can trust

Corporate office : Thyrocare Technologies Limited, D-37/3, TTC MIDC, Turbhe, Navi Mumbai - 400 703

☎ 022 - 3090 0000 / 6712 3400 ☎ 9870666333 ✉ wellness@thyrocare.com 🌐 www.thyrocare.com

**REPORT**

**NAME** : VIVEK KUMAR (32Y/M)  
**REF. BY** : HCL  
**TEST ASKED** : BLOOD SUGAR (F), COMPLETE URINE  
ANALYSIS, HbA1c  
**PATIENTID** : VK21292241  
**MOBILE NO** : 8750200587  
**DOB** : 06/22/1990

**SAMPLE COLLECTED AT :**  
HCL HEALTHCARE NOIDA SEZ CORPORATE  
CENTER,  
HCL HEALTHCARE, GATE 4, PLOT 3A, SECTOR-126,  
NOIDA, UP - 201301, INDIA-201301

**PAN ID** : HN1-20397

| TEST NAME                              | TECHNOLOGY | VALUE | UNITS                      |
|--|------------|-------|----------------------------|
| EST. GLOMERULAR FILTRATION RATE (eGFR) | CALCULATED | 136   | mL/min/1.73 m <sup>2</sup> |

**Reference Range :-**

> = 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

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**REPORT****NAME** : VIVEK KUMAR (32Y/M)**REF. BY** : HCL**TEST ASKED** : BLOOD SUGAR (F),COMPLETE URINE ANALYSIS,HbA1c**PATIENTID** : VK21292241**MOBILE NO** : 8750200587**DOB** : 06/22/1990**SAMPLE COLLECTED AT :**HCL HEALTHCARE NOIDA SEZ CORPORATE  
CENTER,  
HCL HEALTHCARE,GATE 4, PLOT 3A,  
SECTOR-126, NOIDA, UP - 201301,  
INDIA-201301**PAN ID** : HN1-20397

| TEST NAME  | TECHNOLOGY | VALUE | UNITS |
|------------|------------|-------|-------|
| VITAMIN D2 | LC-MS/MS   | 0.63  | ng/mL |

**Method :** LIQUID CHROMATOGRAPHY TANDEM MASS SPECTROMETRY

|            |          |       |       |
|------------|----------|-------|-------|
| VITAMIN D3 | LC-MS/MS | 11.42 | ng/mL |
|------------|----------|-------|-------|

**Method :** LIQUID CHROMATOGRAPHY TANDEM MASS SPECTROMETRY

|                 |          |       |       |
|-----------------|----------|-------|-------|
| VITAMIN D TOTAL | LC-MS/MS | 12.05 | ng/mL |
|-----------------|----------|-------|-------|

**Reference Range :**

Deficiency : &lt;20 ng/mL

Insufficiency : 20-30 ng/mL

Sufficiency : 30-100 ng/mL

Toxicity : &gt;100 ng/mL

**Method :** LIQUID CHROMATOGRAPHY TANDEM MASS SPECTROMETRY**Please correlate with clinical conditions.**

~~ End of report ~~

**Sample Collected on (SCT)** :14 Mar 2023 08:00**Sample Received on (SRT)** : 15 Mar 2023 03:17**Report Released on (RRT)** : 15 Mar 2023 06:32**Sample Type** : SERUM**Labcode** : 1403134268/HCL04**Barcode** : AR559203*Saakshi*

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*Preet Kaur*

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#### 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 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

- ❖ 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.