

## REPORT

**NAME** : N.KANNAIMMAL (63Y/F)  
**REF. BY** : WELLNESS - DSA  
**TEST ASKED** : HBA,HEMOGRAM - 6 PART (DIFF)

**SAMPLE COLLECTED AT** :  
OLD NO 41 , NEW NO 85, C.N.K  
ROAD,TRIPLICANE,CHENNAI-600005 - 600005

| TEST NAME   | METHOD     | VALUE | UNITS |
|---|------------|-------|-------|
| <b>DIABETES SCREEN (BLOOD)</b>                              |            |       |       |
| <b>HbA1c</b>  | H.P.L.C    | 6.2   | %     |
| <b>Reference Range :</b>                                    |            |       |       |
| Below 6.0% - Normal Value                                   |            |       |       |
| 6.0% - 7.0% - Good Control                                  |            |       |       |
| 7.0% - 8.0% - Fair Control                                  |            |       |       |
| 8.0% - 10% - Unsatisfactory Control                         |            |       |       |
| Above 10% - Poor Control                                    |            |       |       |
| <b>Technology</b> : FULLY AUTOMATED H.P.L.C USING TOSOH G8. |            |       |       |
| <b>AVERAGE BLOOD GLUCOSE (ABG)</b>                          | CALCULATED | 131   | mg/dl |
| <b>Reference Range :</b>                                    |            |       |       |
| 90 - 120 mg/dl : Excellent Control                          |            |       |       |
| 121 - 150 mg/dl : Good Control                              |            |       |       |
| 151 - 180 mg/dl : Average Control                           |            |       |       |
| 181 - 210 mg/dl : Action Suggested                          |            |       |       |
| > 211 mg/dl : Panic Value                                   |            |       |       |

(Note: Average Blood Glucose value is calculated from HBA1c value and it indicates Average Blood Sugar level over past three months.)

**Technology** : Derived from HBA1c values

**Please correlate with clinical conditions.**

**Sample Collected on** : 05 Apr 2014 06:00  
**Sample Received on** : 06 Apr 2014 03:48  
**Report Released on** : 06 Apr 2014 08:15  
**Sample Type** : EDTA  
**Labcode** : 050434239/CHE02  
**Barcode** : 35679379/HOME

Dr. Suhas Sakhare MD

Dr. Caesar Sengupta MD

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| TEST NAME                                | VALUE | UNITS                       | REFERENCE RANGE                 |
|--|-------|-----------------------------|---------------------------------|
| TOTAL LEUCOCYTES COUNT                   | 7.13  | X 10 <sup>3</sup> / $\mu$ L | 4 - 10                          |
| NEUTROPHILS                              | 51.3  | %                           | 40-80                           |
| LYMPHOCYTE PERCENTAGE                    | 40.1  | %                           | M:20-40; F:20-40                |
| MONOCYTES                                | 3.9   | %                           | 0-10                            |
| EOSINOPHILS                              | 4.3   | %                           | 0-6                             |
| BASOPHILS                                | 0.1   | %                           | < 1-2                           |
| IMMATURE GRANULOCYTE PERCENTAGE(IG%)     | 0.3   | %                           | Male : 0-0.5 Female: 0-0.4      |
| NEUTROPHILS - ABSOLUTE COUNT             | 3.65  | X 10 <sup>3</sup> / $\mu$ L | 2.0 - 7.0                       |
| LYMPHOCYTES - ABSOLUTE COUNT             | 2.86  | X 10 <sup>3</sup> / $\mu$ L | 1.00 - 3.00                     |
| MONOCYTES - ABSOLUTE COUNT               | 0.28  | X 10 <sup>3</sup> / $\mu$ L | 0.20 - 1.00                     |
| BASOPHILS - ABSOLUTE COUNT               | 0.01  | X 10 <sup>3</sup> / $\mu$ L | 0.02 - 0.10                     |
| EOSINOPHILS - ABSOLUTE COUNT             | 0.31  | X 10 <sup>3</sup> / $\mu$ L | 0.02 - 0.50                     |
| IMMATURE GRANULOCYTES(IG)                | 0.02  | X 10 <sup>3</sup> / $\mu$ L | 0.03                            |
| TOTAL RBC                                | 4.35  | X 10 <sup>6</sup> / $\mu$ L | Male : 4.5-5.5 Female : 3.9-4.8 |
| NUCLEATED RED BLOOD CELLS                | Nil   | X 10 <sup>3</sup> / $\mu$ L | Nil in adults                   |
| NUCLEATED RED BLOOD CELLS %              | Nil   | %                           | Nil in adults                   |
| HEMOGLOBIN                               | 12.7  | g/dL                        | Male : 13-17 Female : 12-15     |
| HEMATOCRIT(PCV)                          | 41.5  | %                           | Male : 40-50 Female : 36-46     |
| MEAN CORPUSCULAR VOLUME(MCV)             | 95.4  | fL                          | 83-101                          |
| MEAN CORPUSCULAR HEMOGLOBIN(MCH)         | 29.2  | pg                          | 27-32                           |
| MEAN CORP.HEMO.CONC(MCHC)                | 30.6  | g/dL                        | 31.5-34.5                       |
| RED CELL DISTRIBUTION WIDTH - SD(RDW-SD) | 50    | fL                          | 39 - 46                         |
| RED CELL DISTRIBUTION WIDTH (RDW-CV)     | 14.3  | %                           | 11.6-14                         |
| PLATELET DISTRIBUTION WIDTH(PDW)         | 12.61 | fL                          | 9.6-15.2                        |
| MEAN PLATELET VOLUME(MPV)                | 8.7   | fL                          | 6.5-12.0                        |
| PLATELET COUNT                           | 347   | X 10 <sup>3</sup> / $\mu$ L | 150-400                         |
| PLATELET TO LARGE CELL RATIO(PLCR)       | 29.5  | %                           | 19.7 - 42.4                     |
| PLATELETCRIT(PCT)                        | 0.37  | %                           | 0.19 - 0.39                     |

**Please Correlate with clinical conditions.**

**Technology** : 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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**REF. BY** : WELLNESS - DSA  
**TEST ASKED** : AAROGYAM 1.2, AMYL, LASE, SOD, CHL

**SAMPLE COLLECTED AT** :  
OLD NO 41 , NEW NO 85, C.N.K  
ROAD, TRIPLICANE, CHENNAI-600005 - 600005

| TEST NAME | METHOD     | VALUE | UNITS |
|-----------|------------|-------|-------|
| AMYLASE   | PHOTOMETRY | 132   | U/L   |

### Reference Range :-

Adults : 30-118 U/L

### Interpretation:

Lipemic Sera (Hypertriglyceridemia) may contain inhibitors, Which falsely depress results. About 20% of patients with Acute Pancreatitis have abnormal lipids. Normal serum amylase may occur in Pancreatitis, Especially relapsing and chronic pancreatitis. Moderate increases may be reported in normal pregnancy.

### Clinical Significance:

Causes of high Serum Amylase include Acute Pancreatitis, Pancreatic Pseudocyst, Pancreatic Ascites, Pancreatic Abscess, Neoplasm in or adjacent to Pancreas, Trauma to Pancreas, and common Duct Stones. Nonpancreatic Causes include inflammatory salivary lesions (Eg, Mumps), Perforated Peptic Ulcer, Intestinal Obstruction, Biliary Tract Disease, Peritonitis, Acute Appendicitis, Diabetic Ketoacidosis, and Extrapaneatic Carcinomas. Amylase levels more than 25-fold the upper limit of normal are often found when metastatic tumors produce Ectopic Amylase.

### Specifications:

Precision: Within run %CV has been recorded 1% and between run %CV of 1.5%. Analytical sensitivity (Lower Detection Limit) 3 U/l

### Kit Validation References:

Tietz Nw, Huang WY, Rauh DF ET Al. Laboratory tests in the differential diagnosis of Hyperamylasemia. Clin Chem 1986;32: 301-307

### Please correlate with clinical conditions.

**Technology :-** ENZYMATIC PHOTOMETRIC TEST.

**Sample Collected on (SCT)** : 05 Apr 2014 06:00  
**Sample Received on (SRT)** : 06 Apr 2014 03:32  
**Report Released on (RRT)** : 06 Apr 2014 17:59  
**Sample Type** : SERUM  
**Labcode** : 050432941/CHE02  
**Barcode** : 35966136/HOME

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|---|--------|-------|--------|
| CHLORIDE  | I.S.E  | 110.4 | mmol/l |
| <b>Reference Range :</b><br>Adults : 98 - 106 mmol/l<br><b>Technology :</b> Ion Selective Electrode in Olympus AU2700 |        |       |        |

**Please correlate with clinical conditions.**

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|--|------------|-------|-------|
| <b>IRON</b><br><b>Reference Range :</b><br>Male : 70 - 180<br>Female : 60 - 180<br><b>Technology :</b> FERROZINE METHOD WITHOUT DEPROTEINIZATION                       | PHOTOMETRY | 99.9  | µg/dl |
| <b>TOTAL IRON BINDING CAPACITY (TIBC)</b><br><b>Reference Range :</b><br>Male: 225 - 535 µg/dl Female: 215 - 535 µg/dl<br><b>Technology :</b> SPECTROPHOTOMETRIC ASSAY | PHOTOMETRY | 344.2 | µg/dl |
| <b>% TRANSFERRIN SATURATION</b><br><b>Reference Range :</b><br>13 - 45<br><b>Technology :</b> DERIVED FROM IRON AND TIBC VALUES  | CALCULATED | 29.02 | %     |

Please correlate with clinical conditions.

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| TEST NAME                 | METHOD     | VALUE | UNITS  | NORMAL RANGE                       |
|---------------------------|------------|-------|--------|------------------------------------|
| BLOOD UREA NITROGEN (BUN) | PHOTOMETRY | 15.1  | mg/dl  | 7.9 - 20                           |
| CREATININE - SERUM        | PHOTOMETRY | 0.66  | mg%    | Male: 0.6 - 1.1 Female: 0.5 - 0.8  |
| URIC ACID                 | PHOTOMETRY | 7.6   | mg/dl  | Male : 3.5 - 7.2 Female: 2.6 - 6.0 |
| CALCIUM                   | PHOTOMETRY | 9.87  | mg/dl  | 8.8 - 10.6                         |
| BUN / SR.CREATININE RATIO | CALCULATED | 22.88 | Ratio  | 9:1 - 23:1                         |
| SODIUM                    | I.S.E      | 145   | mmol/l | 136 - 146                          |

Please correlate with clinical conditions.

### Technology :

BUN - KINETIC UV ASSAY.

SCRE - CREATININE ENZYMATIC METHOD

URIC - ENZYMATIC COLORIMETRIC TEST

CALC - ARSENAZO III METHOD, END POINT.

B/CR - DERIVED FROM SERUM BUN AND CREATININE VALUES

SOD - Ion Selective Electrode in Olympus AU2700

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| TEST NAME | METHOD     | VALUE | UNITS |
|-----------|------------|-------|-------|
| LIPASE    | PHOTOMETRY | 63    | U/l   |

### Reference Range :-

Adults : 5.6 - 51.3 U/l

### Interpretation:

For diagnostic purposes, the results should always be assessed in conjunction with the patient's medical history, clinical examination and other findings like serum amylase. Serum Lipase is usually normal in patients with elevated serum amylase, having peptic ulcer, salivary adenitis, inflammatory bowel disease, intestinal obstruction, and macroamylasemia. Lipemic sera may interfere with results.

### Clinical Significance:

High serum Lipase is a specific marker for pancreatitis; after acute pancreatitis the Lipase activity increases within 4-8 hours, reaches a peak after 24 hours and decreases after 8 to 14 days. However, there is no correlation between the Lipase activity determined in serum and the extent of damage to the pancreas.

### Specifications:

Precision: Within run %CV has been recorded 1.16 % and between run %CV of 0.65 %. Analytical sensitivity (lower detection limit) 3 U/l

### Kit Validation References:

Tietz Nw Et Al. Lipase In Serum - The Elusive Enzyme: An Overview. Clin Chem 1993; 39:746-756.

**Please correlate with clinical conditions.**

**Technology :-** ENZYMATIC COLOUR TEST

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| TEST NAME                 | METHOD     | VALUE | UNITS | NORMAL RANGE |
|---------------------------|------------|-------|-------|--------------|
| TOTAL CHOLESTEROL         | PHOTOMETRY | 253   | mg%   | 125 - 200    |
| HDL CHOLESTEROL - DIRECT  | PHOTOMETRY | 47    | mg%   | 35-80        |
| LDL CHOLESTEROL - DIRECT  | PHOTOMETRY | 154   | mg%   | 85 - 130     |
| TRIGLYCERIDES             | PHOTOMETRY | 344   | mg%   | 25 - 200     |
| TC/ HDL CHOLESTEROL RATIO | CALCULATED | 5.4   | Ratio | 3.0 - 5.0    |
| LDL / HDL RATIO           | CALCULATED | 3.3   | Ratio | 1.5 - 3.5    |
| VLDL CHOLESTEROL          | CALCULATED | 68.8  | mg%   | 5 - 40       |

Please correlate with clinical conditions.

### Technology :

CHOL - CHOD POD METHOD

HCHO - ENZYME SELECTIVE PROTECTION METHOD

LDL - HOMOGENOUS ENZYMATIC COLORIMETRIC ASSAY

TRIG - ENZYMATIC COLORIMETRIC METHOD (GPO) [HIGHLY INFLUENCED BY LEVEL OF FASTING]

TC/H - DERIVED FROM SERUM CHOLESTEROL AND HDL VALUES

LDL/ - DERIVED FROM SERUM HDL AND LDL VALUES

VLDL - DERIVED FROM SERUM TRIGLYCERIDE VALUES

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|------------------------------------|------------|-------|-------|---------------------------|
| ALKALINE PHOSPHATASE               | PHOTOMETRY | 107   | U/l   | M:53 to 128 - F:42 to 98  |
| BILIRUBIN -DIRECT                  | PHOTOMETRY | 0.14  | mg/dl | 0 - 0.20                  |
| BILIRUBIN - TOTAL                  | PHOTOMETRY | 0.44  | mg/dl | 0.30 - 1.20               |
| BILIRUBIN (INDIRECT)               | CALCULATED | 0.3   | mg/dl | 0 - 0.9                   |
| GAMMA GLUTAMYL TRANSFERASE (GGT)   | PHOTOMETRY | 45    | U/l   | M: 0 to 55 - F :0 to 38   |
| ASPARTATE AMINOTRANSFERASE (SGOT ) | PHOTOMETRY | 23    | U/l   | M: 0 to 37 - F: 0 to 31   |
| ALANINE TRANSAMINASE (SGPT)        | PHOTOMETRY | 28    | U/l   | M: 13 to 40 - F: 10 to 28 |
| PROTEIN - TOTAL                    | PHOTOMETRY | 8     | gm/dl | 6.6 - 8.3                 |
| ALBUMIN - SERUM                    | PHOTOMETRY | 4.8   | gm/dl | 3.5 - 5.2                 |
| SERUM GLOBULIN                     | PHOTOMETRY | 3.2   | gm/dL | 2.30-3.50                 |
| SERUM ALBUMIN/GLOBULIN RATIO       | CALCULATED | 1.5   | Ratio | 0.9 - 2.0                 |

**Please correlate with clinical conditions.**

### Technology :

ALKP - ALP IFCC\* LIQUID (COLORIMETRIC ASSAY)  
BILD - DIAZO METHOD OF PEARLMAN & LEE, ENDPOINT.  
BILT - DIAZO METHOD OF PEARLMAN & LEE, ENDPOINT.  
BILI - DERIVED FROM SERUM TOTAL AND DIRECT BILIRUBIN VALUES  
GGT - IFCC STANDARDISED SZASZ METHOD  
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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|-----------------------------------|---------|-------|--------|-----------------|
| TOTAL TRIIODOTHYRONINE (T3)       | C.L.I.A | 131   | ng/dl  | 60 - 200        |
| TOTAL THYROXINE (T4)              | C.L.I.A | 9     | µg/dl  | 4.5 - 12.0      |
| THYROID STIMULATING HORMONE (TSH) | C.L.I.A | 3.96  | µIU/ml | 0.30 - 5.5      |

**Comments** : SUGGESTING THYRONORMALCY

**Please correlate with clinical conditions.**

**Technology :**

T3 - COMPETITIVE CHEMI LUMINESCENT IMMUNO ASSAY

T4 - COMPETITIVE CHEMI LUMINESCENT IMMUNO ASSAY

TSH - ULTRA SENSITIVE SANDWICH CHEMI LUMINESCENT IMMUNO ASSAY

Pregnancy reference ranges for TSH

1st Trimester : 0.30 - 4.50

2nd Trimester : 0.50 - 4.60

3rd Trimester : 0.80 - 5.20

Reference:

National Health and Nutrition examination survey, J Clin Endocrinol Metab. 2002; 87;489

~~ End of report ~~

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