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 13:57 Hrs.

OPD/IPD DOC NO OP40162332

PATIENT CODE 1000351368

REFERRING DOCTOR Dr. Mohit Kharbanda

ACCESSION NO DH-101/2025-26/0024303

AGE 56 Yrs 10 Mths 21 Dys SEX Female



Results relate only to the samples tested

TEST REPORT STATUS	RESULTS	BIOLOGICAL REFERENCE INTERVAL	UNITS
Glucose - Fasting			
Glucose - Fasting Specimen : Plasma Flouride Methodology : Hexokinase	108	70.0 - 100.0	mg/dL
Urea			
Urea Specimen : Serum Methodology : Urease GLDH, UV	14.2	15.0 - 45.0	mg/dL
Creatinine			
Creatinine Specimen : Serum Methodology : Jaffe	0.79	Adult Male : < 50 yrs : 0.84 - 1.25 > 50 yrs : 0.81 - 1.44 Adult Female : 0.66 - 1.09 Child : 0.5 - 1.2 Infant : 0.4 - 0.7 Neonate : 0.5 - 1.2	mg/dL
Electrolytes (Na, K, Cl)			
Sodium Specimen : Serum Methodology : Direct ISE / Indirect ISE	140.6	135.0 - 150.0	mmol/L
Potassium Specimen : Serum Methodology : Direct ISE / Indirect ISE	3.99	3.5 - 5.0	mmol/L
Chloride Specimen : Serum Methodology : Direct ISE / Indirect ISE	105.1	94.0 - 110.0	mmol/L

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 Dr. JAYATI GUPTA
 Ph.D (Bio.Chem)
 Senior Consultant Biochemist

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PATIENT NAME & ADDRESS

MRS. MEENAKSHI SANYAL

PATHOLOGY



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LFT (Liver Function Test)			
Total Bilirubin Specimen : Serum Methodology : Diazo Salt	0.62	Normal : 0.3 - 1.1 New Born : 1.2 - 12.0	mg/dL
Direct Bilirubin Specimen : Serum Methodology : Diazo Salt	0.13	Adult and Child : < 0.2	mg/dL
Indirect Bilirubin Methodology : Calculated Value	0.49		mg/dL
Total Protein Specimen : Serum Methodology : Biuret	7.55	6.7 - 8.3	gm/dl
Albumin Specimen : Serum Methodology : Bromocresol Green (BCG)	4.15	4.0 - 5.0	g/dL
Globulin Methodology : Calculated Value	3.40	1.8 - 3.6	g/dL
A/G Ratio Methodology : Calculated Value	1.22	1.0 - 2.0 : 1	ratio
Aspartate Aminotransferase (SGOT) (AST) Specimen : Serum Methodology : UV without P5P	33	13.0 - 33.0	U/L
Alanine Aminotransferase (SGPT) (ALT) Specimen : Serum Methodology : UV without P-5-P	36	6 - 27	U/L
Alkaline Phosphatase (ALP) Specimen : Serum Methodology : IFCC pNPP with AMP Buffer	98	30 - 120	U/L

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TEST REPORT STATUS	RESULTS	BIOLOGICAL REFERENCE INTERVAL	UNITS
Lipid Profile			
Cholesterol - Total Specimen : Serum Methodology : Cholesterol oxidase, POD	155	130.0 - 220.0	mg/dL
Cholesterol - HDL Specimen : Serum Methodology : Direct / Innovative Detergent	59.8	40.0 - 59.0	mg/dL
Cholesterol - LDL Methodology : Calculated Value	69.0	> 160.0 : High Risk 130.0 – 160.0 : Borderline High <= 130.0 : Desirable	mg/dL
Cholesterol - VLDL Methodology : Calculated Value	26.2	< 40.0	mg/dL
Triglyceride Specimen : Serum Methodology : GPO/POD	131	30.0 - 150.0	mg/dL
Cholesterol - Total/HDL ratio Methodology : Calculated Value	2.59	3.3 : 1/2 Average Risk 4.4 : Average Risk 7.1 : 2 x Average Risk 11.0 : 3 x Average Risk	ratio
Cholesterol - LDL/HDL ratio Methodology : Calculated Value	1.15		

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TEST REPORT STATUS	RESULTS	BIOLOGICAL REFERENCE INTERVAL	UNITS
HbA1c (Glycosylated Haemoglobin) Glycosylated Haemoglobin (HBA1C) Specimen : Whole Blood - EDTA Methodology : NGSP (Enzymatic Assay) * CLINICAL CORRELATION REQUESTED.	* 5.97	< 5.7 : Normal 5.7 - 6.4 : Prediabetes ≥ 6.5 : Higher Diabetics	%

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TEST REPORT STATUS	RESULTS	BIOLOGICAL REFERENCE INTERVAL	UNITS
Apolipoprotein A-I Apolipoprotein A-I Specimen : Serum Methodology : Spectrophotometry	161.2	122 - 161	mg/dL
Apolipoprotein B Apolipoprotein B Specimen : Serum Methodology : Spectrophotometry	71.2	69 - 105	mg/dL

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TEST REPORT STATUS	RESULTS	BIOLOGICAL REFERENCE INTERVAL	UNITS
TSH (Thyroid Stimulating Hormone) Thyroid Stimulating Hormone (TSH) Specimen : Serum Methodology : Electrochemiluminescence	1.78	0.27 - 4.20	μIU/mL

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TEST REPORT STATUS	RESULTS	BIOLOGICAL REFERENCE INTERVAL	UNITS
CBC (Complete Blood Count)			
Haemoglobin (Hb) Specimen : Whole Blood - EDTA Methodology : Colorimetric Non-cyanide Haemoglobin method	12.2	12.0 - 15.0	gm/dl
RBC Count Specimen : Whole Blood - EDTA Methodology :	4.08	3.8 - 4.8	million/cmm
Packed Cell Volume (Hematocrit) (PCV) Specimen : Whole Blood - EDTA Methodology : Pulse height detection	37.5	36.0 - 46.0	%
Mean Cell Volume (MCV) Specimen : Whole Blood - EDTA Methodology : Calculated Value	92.0	83.0 - 101.0	fL
Mean Cell Haemoglobin (MCH) Specimen : Whole Blood - EDTA Methodology : Calculated Value	29.9	27 - 32	pg
Mean Cell Haemoglobin Concentration (MCHC) Specimen : Whole Blood - EDTA Methodology : Calculated Value	32.5	31.5 - 34.5	g/dL
Platelet Count Specimen : Whole Blood - EDTA Methodology : Electrical Impedance & Neubauer's chamber	1.95	1.5 - 4.0	lakh/cmm
Total Count			
WBC Count Specimen : Whole Blood - EDTA Methodology : Electrical Impedance & Manual	5.4	4.0 - 11.0	thou/cmm
Differential Count (Elect. Impedance & Microscopy)			
Neutrophil	57	40 - 80	%
Lymphocyte	38	20 - 40	%

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Consultant Pathologist

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TEST REPORT STATUS	RESULTS	BIOLOGICAL REFERENCE INTERVAL	UNITS
CBC (Complete Blood Count)			
Monocyte	02	2 - 10	%
Eosinophil	03	1 - 6	%
Basophil	00	0 - 2	%
Peripheral Blood Smear (Microscopy)			
RBC	Normocytic Normochromic		
WBC	No immature cell seen		
Platelet	Adequate		
Specimen : Whole Blood - EDTA			
Erythrocyte Sedimentation Rate (ESR)	33	<= 20	mm / hr
Specimen : Whole Blood - EDTA			
Methodology : Westergren			

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TEST REPORT STATUS	RESULTS	BIOLOGICAL REFERENCE INTERVAL	UNITS
Apolipoprotein A-I			
The decreased level is associated with increased risk of coronary heart disease (CHD).			
Elevated in:			
i.Familial hyperalphalipoproteinemia			
ii.Pregnancy			
iii.Estrogen therapy			
iv.Alcohol consumption			
v.Exercise			
Reduced in :			
i.Tangier disease			
ii."Fish-eye" disease			
iii.Familial hypoalphalipoproteinemia			
iv.Familial lecithin-cholesterol acyltransferase deficiency			
v.Type I and type V hyperlipoproteinemia			
vi.Diabetes mellitus			
vii.Cholestasis			
viii.Hemodialysis			
ix.Infection			
x.Drugs (e.g. Diuretics, beta-blockers etc.)			
Apolipoprotein B			
Elevated levels are associated with increased risk of CHD.			
Elevated in:			
i.Hyperlipoproteinemia (types IIa, IIb, IV, V)			
ii.Familial hyperapobetalipoproteinemia			
iii.Nephrotic syndrome			
iv.Pregnancy			
v.Biliary obstruction			
vi.Hemodialysis			
vii.Cigarette smoking			
viii.Drugs (e.g. Diuretics, beta-blockers, cyclosporine, glucocorticoids)			
Reduced in			
i.Hypo- and Abetalipoproteinemia			
ii.Type I hyperlipoproteinemia (hyperchylomicronemia)			
iii.Liver disease			
iv.Exercise			
v.Infections			

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TEST REPORT STATUS	RESULTS	BIOLOGICAL REFERENCE INTERVAL	UNITS
Apolipoprotein B ii.Type I hyperlipoproteinemia (hyperchylomicronemia) iii.Liver disease iv.Exercise v.Infections vi.Drugs (e.g. Cholesterol-lowering drugs, estrogens)	<div>----- End of Report -----</div>		

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