



PKR JAIN HEALTHCARE INSTITUTE NASIRPUR, Hissar Road, AMBALA CITY- (Haryana)

A PIONEER DIAGNOSTIC CENTRE

■ 0171-2532620, 8222896961 **■** pkrjainhealthcare@gmail.com

NAME : Mrs. SAVITRI

AGE/ GENDER : 49 YRS/FEMALE **PATIENT ID** : 1817914

COLLECTED BY : 122504040015 REG. NO./LAB NO.

REFERRED BY **REGISTRATION DATE** : 04/Apr/2025 02:32 PM BARCODE NO. : 12507901 **COLLECTION DATE** : 04/Apr/2025 03:50PM CLIENT CODE. : P.K.R JAIN HEALTHCARE INSTITUTE REPORTING DATE : 04/Apr/2025 04:07PM

CLIENT ADDRESS : NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYANA

Value Unit Test Name **Biological Reference interval**

HAEMATOLOGY HAEMOGLOBIN (HB)

HAEMOGLOBIN (HB) gm/dL 12.0 - 16.0 8.5^{L}

by CALORIMETRIC

INTERPRETATION:-

Hemoglobin is the protein molecule in red blood cells that carries oxygen from the lungs to the bodys tissues and returns carbon dioxide from the tissues back to the lungs.

A low hemoglobin level is referred to as ANEMIA or low red blood count.

ANEMIA (DECRESED HAEMOGLOBIN):

- 1) Loss of blood (traumatic injury, surgery, bleeding, colon cancer or stomach ulcer)
- 2) Nutritional deficiency (iron, vitamin B12, folate)
- 3) Bone marrow problems (replacement of bone marrow by cancer)
- 4) Suppression by red blood cell synthesis by chemotherapy drugs
- 5) Kidney failure
- 6) Abnormal hemoglobin structure (sickle cell anemia or thalassemia). POLYCYTHEMIA (INCREASED HAEMOGLOBIN):

- 1) People in higher altitudes (Physiological)
- 2) Smoking (Secondary Polycythemia)
- 3) Dehydration produces a falsely rise in hemoglobin due to increased haemoconcentration
- 4) Advanced lung disease (for example, emphysema)
- 5) Certain tumors
- 6) A disorder of the bone marrow known as polycythemia rubra vera,
- 7) Abuse of the drug erythropoetin (Epogen) by athletes for blood doping purposes (increasing the amount of oxygen available to the body by chemically raising the production of red blood cells).

NOTE: TEST CONDUCTED ON EDTA WHOLE BLOOD



DR.VINAY CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY & MICROBIOLOGY)



440 Dated 17.5.2012 u/s 80 G OF INCOME TAX ACT. PAN NO. AAAAP1600. REPORT ATTRACTS THE CONDITIONS PRINTED OVERLEAF (P.T.O.)





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NASIRPUR, Hissar Road, AMBALA CITY- (Haryana)

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: 04/Apr/2025 04:31PM

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REPORTING DATE

GLYCOSYLATED HAEMOGLOBIN (HBA1C)

GLYCOSYLATED HAEMOGLOBIN (HbA1c): 8.2H 4.0 - 6.4

WHOLE BLOOD

CLIENT CODE.

by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY)

60.00 - 140.00 ESTIMATED AVERAGE PLASMA GLUCOSE mg/dL 188.64^H by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY)

INTERPRETATION:

AS PER AMERICAN DIABETES ASSOCIATION (ADA):					
REFERENCE GROUP	GLYCOSYLATED HEMOGLOGIB (HBAIC) in %				
Non diabetic Adults >= 18 years	<5.7				
At Risk (Prediabetes)	5.7 – 6.4				
Diagnosing Diabetes	>= 6.5				
Therapeutic goals for glycemic control	Age > 19 Years				
	Goals of Therapy:	< 7.0			
	Actions Suggested:	>8.0			
	Age < 19 Years				
	Goal of therapy:	<7.5			

COMMENTS:

- 1. Glycosylated hemoglobin (HbA1c) test is three monthly monitoring done to assess compliace with therapeutic regimen in diabetic patients.
- 2. Since Hb1c reflects long term fluctuations in blood glucose concentration, a diabetic patient who has recently under good control may still have high concentration of HbAlc. Converse is true for a diabetic previously under good control but now poorly controlled.
- 3.Target goals of < 7.0 % may be beneficial in patients with short duration of diabetes, long life expectancy and no significant cardiovascular disease. In patients with significant complications of diabetes, limited life expectancy or extensive co-morbid conditions, targetting a goal of < 7.0% may not be 4.High
- HbA1c (>9.0 -9.5 %) is strongly associated with risk of development and rapid progression of microvascular and nerve complications
- 5.Any condition that shorten RBC life span like acute blood loss, hemolytic anemia falsely lower HbA1c results.
- 6.HbA1c results from patients with HbSS,HbSC and HbD must be interpreted with caution, given the pathological processes including anemia,increased red cell turnover, and transfusion requirement that adversely impact HbA1c as a marker of long-term gycemic control.

7. Specimens from patients with polycythemia or post-splenctomy may exhibit increse in HbA1c values due to a somewhat longer life span of the red cells.



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DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY)



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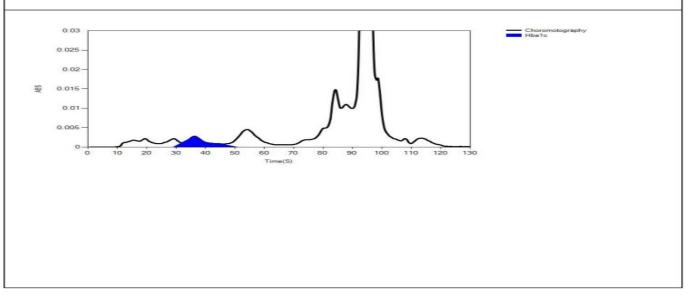
CLIENT ADDRESS : NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYANA

Value Unit Test Name **Biological Reference interval**

LIFOTRONIC Graph Report

Name :	Case:	Patient Type :	Test Date: 04/04/2025 16:27:12
Age:	Department:	Sample Type: Whole Blood EDTA	Sample ld: 12507901
Gender:			Total Area: 5352

Peak Name	Retention Time(s)	Absorbance	Area	Result (Area %)
нь ао	69	1461	4516	80.1
HbA1c	40	45	460	8.2
La1c	26	28	182	3.2
HbF	21	22	32	0.6
Hba1b	14	22	84	1.5
Hba1a	11	18	78	1.4





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NAME : Mrs. SAVITRI

by GLYCEROL PHOSPHATE OXIDASE (ENZYMATIC)

by CALCULATED, SPECTROPHOTOMETRY

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Value Unit Test Name **Biological Reference interval**

REPORTING DATE

CLINICAL CHEMISTRY/BIOCHEMISTRY

LIPID PROFILE: BASIC

CHOLESTEROL TOTAL: SERUM 118.99 OPTIMAL: < 200.0 mg/dL by CHOLESTEROL OXIDASE PAP

BORDERLINE HIGH: 200.0 -

239.0 HIGH CHOLESTEROL: > OR =

: 04/Apr/2025 05:17PM

240.0 TRIGLYCERIDES: SERUM 100.94 OPTIMAL: < 150.0 mg/dL

BORDERLINE HIGH: 150.0 -

199.0

HIGH: 200.0 - 499.0

VERY HIGH: > OR = 500.0

HDL CHOLESTEROL (DIRECT): SERUM 43.81 mg/dL LOW HDL: < 30.0

by SELECTIVE INHIBITION BORDERLINE HIGH HDL: 30.0 -

60.0

HIGH HDL: > OR = 60.0LDL CHOLESTEROL: SERUM 54.99 mg/dL OPTIMAL: < 100.0

by CALCULATED, SPECTROPHOTOMETRY ABOVE OPTIMAL: 100.0 - 129.0

BORDERLINE HIGH: 130.0 -

159.0

HIGH: 160.0 - 189.0

VERY HIGH: > OR = 190.0

NON HDL CHOLESTEROL: SERUM 75.18 OPTIMAL: < 130.0 mg/dL

ABOVE OPTIMAL: 130.0 - 159.0

BORDERLINE HIGH: 160.0 -

189.0

HIGH: 190.0 - 219.0

VERY HIGH: > OR = 220.0

VLDL CHOLESTEROL: SERUM 0.00 - 45.0020.19 mg/dL

by CALCULATED, SPECTROPHOTOMETRY TOTAL LIPIDS: SERUM 350.00 - 700.00 mg/dL 338.92^L

by CALCULATED, SPECTROPHOTOMETRY

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Test Name	Value	Unit	Biological Reference interval
CHOLESTEROL/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	2.72	RATIO	LOW RISK: 3.30 - 4.40 AVERAGE RISK: 4.50 - 7.0 MODERATE RISK: 7.10 - 11.0 HIGH RISK: > 11.0
LDL/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	1.26	RATIO	LOW RISK: 0.50 - 3.0 MODERATE RISK: 3.10 - 6.0 HIGH RISK: > 6.0
TRIGLYCERIDES/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	2.3 ^L	RATIO	3.00 - 5.00

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1.Measurements in the same patient can show physiological& analytical variations. Three serial samples 1 week apart are recommended for Total Cholesterol, Triglycerides, HDL & LDL Cholesterol.

2. As per NLA-2014 guidelines, all adults above the age of 20 years should be screened for lipid status. Selective screening of children above the age of 2 years with a family history of premature cardiovascular disease or those with at least one parent with high total cholesterol is recommended

3. Low HDL levels are associated with increased risk for Atherosclerotic Cardiovascular disease (ASCVD) due to insufficient HDL being available to participate in reverse cholesterol transport, the process by which cholesterol is eliminated from peripheral tissues.

4. NLA-2014 identifies Non HDL Cholesterol (an indicator of all atherogeniclipoproteins such as LDL, VLDL, IDL, Lpa, Chylomicron remnants) along

with LDL-cholesterol as co- primary target for cholesterol lowering therapy. Note that major risk factors can modify treatment goals for LDL &Non

5. Additional testing for Apolipoprotein B, hsCRP,Lp(a) & LP-PLA2 should be considered among patients with moderate risk for ASCVD for risk refinement



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Value Unit Test Name **Biological Reference interval**

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UREA

mg/dL **UREA: SERUM** 30.1 10.00 - 50.00 by UREASE - GLUTAMATE DEHYDROGENASE (GLDH)



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Value Unit Test Name **Biological Reference interval**

CREATININE

CREATININE: SERUM mg/dL 0.85 0.40 - 1.20

* End Of Report ***



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