A PIONEER DIAGNOSTIC CENTRE

🕻 0171-2532620, 8222896961 🛛 🖾 pkrjainhealthcare@gmail.com

	: Mr. KRISHAN LAL					
AGE/ GENDER	: 76 YRS/MALE		PATIENT ID		: 1273398	
COLLECTED BY	:		REG. NO./LAI	3 NO.	: 122502260005	
REFERRED BY	:		REGISTRATI	ON DATE	: 26/Feb/2025 09:06 AM	
BARCODE NO.	: 12507233		COLLECTION	DATE	: 26/Feb/2025 09:18AM	
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INS	FITUTE	REPORTING		: 26/Feb/2025 04:49PM	
CLIENT ADDRESS					. 20/ FeD/ 2023 04.49FM	
LIENT ADDRESS	. NASIRI UR, IIISSAR ROAD, AN					
Test Name		Value		Unit	Biological Reference inte	erva
		HATA	ATOLOCY			
		HALV	ATOLOGY			
	GLYCO	DSYLATED H	AEMOGLOBI	N (HBA1C	C)	
WHOLE BLOOD	EMOGLOBIN (HbA1c):		AEMOGLOBI	N (HBA10 %	C) 4.0 - 6.4	
WHOLE BLOOD by hplc (high perfor ESTIMATED AVERAC		OSYLATED H	AEMOGLOBI	•		
WHOLE BLOOD by HPLC (HIGH PERFOR ESTIMATED AVERAC by HPLC (HIGH PERFOR	EMOGLOBIN (HbA1c): RMANCE LIQUID CHROMATOGRAPHY) GE PLASMA GLUCOSE RMANCE LIQUID CHROMATOGRAPHY)	DSYLATED H 7.9 ^H 180.03 ^H		%	4.0 - 6.4	
WHOLE BLOOD by HPLC (HIGH PERFOR ESTIMATED AVERA(by HPLC (HIGH PERFOR NTERPRETATION:	EMOGLOBIN (HbA1c): RMANCE LIQUID CHROMATOGRAPHY) GE PLASMA GLUCOSE	DSYLATED H 7.9 ^H 180.03 ^H DIABETES ASSOC		% mg/dL	4.0 - 6.4 60.00 - 140.00	
NHOLE BLOOD by HPLC (HIGH PERFOR ESTIMATED AVERA(by HPLC (HIGH PERFOR <u>NTERPRETATION:</u> R Non dia	EMOGLOBIN (HbA1c): RMANCE LIQUID CHROMATOGRAPHY) GE PLASMA GLUCOSE RMANCE LIQUID CHROMATOGRAPHY) AS PER AMERICAN REFERENCE GROUP Ibetic Adults >= 18 years	DSYLATED H 7.9 ^H 180.03 ^H DIABETES ASSOC	IATION (ADA): SLYCOSYLATED H	% mg/dL EMOGLOGIB <5.7	4.0 - 6.4 60.00 - 140.00	
WHOLE BLOOD by HPLC (HIGH PERFOR ESTIMATED AVERA(by HPLC (HIGH PERFOR NTERPRETATION: R R Non dia At	EMOGLOBIN (HbA1c): RMANCE LIQUID CHROMATOGRAPHY) GE PLASMA GLUCOSE RMANCE LIQUID CHROMATOGRAPHY) AS PER AMERICAN REFERENCE GROUP Ibetic Adults >= 18 years Risk (Prediabetes)	DSYLATED H 7.9 ^H 180.03 ^H DIABETES ASSOC	IATION (ADA): SLYCOSYLATED H	% mg/dL EMOGLOGIB <5.7 5.7 - 6.4	4.0 - 6.4 60.00 - 140.00	
WHOLE BLOOD by HPLC (HIGH PERFOR ESTIMATED AVERA(by HPLC (HIGH PERFOR NTERPRETATION: R R Non dia At	EMOGLOBIN (HbA1c): RMANCE LIQUID CHROMATOGRAPHY) GE PLASMA GLUCOSE RMANCE LIQUID CHROMATOGRAPHY) AS PER AMERICAN REFERENCE GROUP Ibetic Adults >= 18 years	DSYLATED H 7.9 ^H 180.03 ^H DIABETES ASSOC	IATION (ADA): SLYCOSYLATED H	% mg/dL < <u>EMOGLOGIB</u> < <u>5.7</u> 5.7 - 6.4 >= 6.5	4.0 - 6.4 60.00 - 140.00	
WHOLE BLOOD by HPLC (HIGH PERFOR ESTIMATED AVERA(by HPLC (HIGH PERFOR NTERPRETATION: R R Non dia At	EMOGLOBIN (HbA1c): RMANCE LIQUID CHROMATOGRAPHY) GE PLASMA GLUCOSE RMANCE LIQUID CHROMATOGRAPHY) AS PER AMERICAN REFERENCE GROUP Ibetic Adults >= 18 years Risk (Prediabetes)	DSYLATED H 7.9 ^H 180.03 ^H DIABETES ASSOC	IATION (ADA): SLYCOSYLATED H	% mg/dL EMOGLOGIB <5.7 5.7 - 6.4	4.0 - 6.4 60.00 - 140.00 (HBAIC) in %	
WHOLE BLOOD by HPLC (HIGH PERFOR ESTIMATED AVERA(by HPLC (HIGH PERFOR INTERPRETATION: R Non dia At Dia	EMOGLOBIN (HbA1c): RMANCE LIQUID CHROMATOGRAPHY) GE PLASMA GLUCOSE RMANCE LIQUID CHROMATOGRAPHY) AS PER AMERICAN REFERENCE GROUP Ibetic Adults >= 18 years Risk (Prediabetes) agnosing Diabetes	DSYLATED H 7.9 ^H 180.03 ^H DIABETES ASSOC	HATION (ADA): SLYCOSYLATED H Age s of Therapy:	% mg/dL < <u>EMOGLOGIB</u> < <u>5.7</u> 5.7 - 6.4 >= 6.5	4.0 - 6.4 60.00 - 140.00 (HBAIC) in %	
WHOLE BLOOD by HPLC (HIGH PERFOR ESTIMATED AVERA(by HPLC (HIGH PERFOR NTERPRETATION: R Non dia At Dia	EMOGLOBIN (HbA1c): RMANCE LIQUID CHROMATOGRAPHY) GE PLASMA GLUCOSE RMANCE LIQUID CHROMATOGRAPHY) AS PER AMERICAN REFERENCE GROUP Ibetic Adults >= 18 years Risk (Prediabetes)	DSYLATED H 7.9 ^H 180.03 ^H DIABETES ASSOC	SILYCOSYLATED H SILYCOSYLATED H Age s of Therapy: ns Suggested:	% mg/dL < <u>EMOGLOGIB</u> < <u>5.7</u> 5.7 - 6.4 >= 6.5	4.0 - 6.4 60.00 - 140.00 (HBAIC) in %	

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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CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITUTE		RTING DATE	: 26/Feb/2025 01:35PM	
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, A	MBALA CITY - HARYAN	A		
Test Name		Value	Unit	Biological Reference interval	
	CLINI	CAL CHEMISTRY	/BIOCHEMIST	RY	
		GLUCOSE FAS			
		die cosi mo			
GLUCOSE FASTING by GLUCOSE OXIDAS	G (F): PLASMA E - PEROXIDASE (GOD-POD)	113.05 ^H	mg/dL	NORMAL: < 100.0 PREDIABETIC: 100.0 - 125.0 DIABETIC: > 0R = 126.0	

A fasting plasma glucose level below 100 mg/dl is considered normal.
A fasting plasma glucose level between 100 - 125 mg/dl is considered as glucose intolerant or prediabetic. A fasting and post-prandial blood test (after consumption of 75 gms of glucose) is recommended for all such patients.
A fasting plasma glucose level of above 125 mg/dl is highly suggestive of diabetic state. A repeat post-prandial is strongly recommended for all such patients. A fasting plasma glucose level in excess of 125 mg/dl on both occasions is confirmatory for diabetic state.





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CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITUTE : NASIRPUR, HISSAR ROAD, AMBALA CITY -	REPORTING DATE	: 26/Feb/2025 01:32PM
CLIENT ADDRESS			
Test Name	Value	Unit	Biological Reference interval
by CMIA (CHEMILUMIN 3rd GENERATION, ULT	THYROID STIMU ATING HORMONE (TSH): SERUM 5.65 ^H JESCENT MICROPARTICLE IMMUNOASSAY) 5.65 ^H	DCRINOLOGY ILATING HORMONE (TS µIU/mL	5H) 0.35 - 5.50
by CMIA (CHEMILUMIN 3rd GENERATION, ULT	THYROID STIMU ATING HORMONE (TSH): SERUM MESCENT MICROPARTICLE IMMUNOASSAY) RASENSITIVE	LATING HORMONE (TS μIU/mL	0.35 - 5.50
by CMIA (CHEMILUMIN 3rd GENERATION, ULT	THYROID STIMU TING HORMONE (TSH): SERUM S.65 ^H IESCENT MICROPARTICLE IMMUNOASSAY) RASENSITIVE	LATING HORMONE (TS μIU/mL REFFERENCE RANGE	0.35 - 5.50 (µlU/mL)
by CMIA (CHEMILUMIN 3rd GENERATION, ULT	THYROID STIMU TING HORMONE (TSH): SERUM IESCENT MICROPARTICLE IMMUNOASSAY) RASENSITIVE AGE 0 – 5 DAYS	LATING HORMONE (TS μIU/mL REFFERENCE RANGE (0.70 – 15.20	0.35 - 5.50 (µlU/mL)
by CMIA (CHEMILUMIN 3rd GENERATION, ULT	THYROID STIMU TING HORMONE (TSH): SERUM S.65 ^H IESCENT MICROPARTICLE IMMUNOASSAY) RASENSITIVE	LATING HORMONE (TS μIU/mL REFFERENCE RANGE	0.35 - 5.50 (µlU/mL)
by CMIA (CHEMILUMIN 3rd GENERATION, ULT	THYROID STIMU TING HORMONE (TSH): SERUM S.65 ^H ISECENT MICROPARTICLE IMMUNOASSAY) RASENSITIVE AGE 0 – 5 DAYS 6 6 Days – 2 Months O	LATING HORMONE (TS μlU/mL <u>REFFERENCE RANGE</u> 0.70 – 15.20 0.70 – 11.00	0.35 - 5.50 (µlU/mL)
by CMIA (CHEMILUMIN 3rd GENERATION, ULT	THYROID STIMU TING HORMONE (TSH): SERUM S.65 ^H ISECENT MICROPARTICLE IMMUNOASSAY) RASENSITIVE AGE 0-5 DAYS 0 6 Days - 2 Months 0 3 - 11 Months 0 1 - 5 Years 0 6 - 10 Years 0	LATING HORMONE (TS μlU/mL REFFERENCE RANGE 0.70 – 15.20 0.70 – 11.00 0.70 – 8.40	0.35 - 5.50 (µlU/mL)
by CMIA (CHEMILUMIN 3rd GENERATION, ULT	THYROID STIMU TING HORMONE (TSH): SERUM S.65 ^H IESCENT MICROPARTICLE IMMUNOASSAY) RASENSITIVE AGE 0 - 5 DAYS 0 6 Days - 2 Months 0 3 - 11 Months 0 1 - 5 Years 0 6 - 10 Years 0 11 - 15 0	LATING HORMONE (TS μIU/mL REFFERENCE RANGE (0.70 – 15.20 0.70 – 11.00 0.70 – 8.40 0.70 – 7.00 0.60 – 5.50 0.50 – 5.50	0.35 - 5.50 (µlU/mL)
	THYROID STIMU TING HORMONE (TSH): SERUM S.65 ^H IESCENT MICROPARTICLE IMMUNOASSAY) RASENSITIVE AGE 0 - 5 DAYS 0 6 Days - 2 Months 0 3 - 11 Months 0 1 - 5 Years 0 6 - 10 Years 0 11 - 15 0 > 20 Years (Adults) 0	LATING HORMONE (TS μIU/mL REFFERENCE RANGE (0.70 – 15.20 0.70 – 15.20 0.70 – 11.00 0.70 – 8.40 0.70 – 7.00 0.60 – 5.50 0.50 – 5.50 0.27 – 5.50	0.35 - 5.50 (µlU/mL)
by CMIA (CHEMILUMIN 3rd GENERATION, ULT	THYROID STIMU TING HORMONE (TSH): SERUM 15.65 ^H IESCENT MICROPARTICLE IMMUNOASSAY) RASENSITIVE AGE 0 - 5 DAYS 0 6 Days - 2 Months 0 3 - 11 Months 0 1 - 5 Years 0 6 - 10 Years 0 11 - 15 0 > 20 Years (Adults) 0	LATING HORMONE (TS μIU/mL REFFERENCE RANGE (0.70 – 15.20 0.70 – 11.00 0.70 – 8.40 0.70 – 7.00 0.60 – 5.50 0.50 – 5.50 0.27 – 5.50 Y	0.35 - 5.50 (µlU/mL)
by CMIA (CHEMILUMIN 3rd GENERATION, ULT	THYROID STIMU TING HORMONE (TSH): SERUM S.65 ^H IESCENT MICROPARTICLE IMMUNOASSAY) RASENSITIVE AGE 0 - 5 DAYS 0 6 Days - 2 Months 0 3 - 11 Months 0 1 - 5 Years 0 6 - 10 Years 0 11 - 15 0 > 20 Years (Adults) 0	LATING HORMONE (TS μIU/mL REFFERENCE RANGE (0.70 – 15.20 0.70 – 15.20 0.70 – 11.00 0.70 – 8.40 0.70 – 7.00 0.60 – 5.50 0.50 – 5.50 0.27 – 5.50	0.35 - 5.50 (µlU/mL)

NOTE:-TSH levels are subjected to circardian variation, reaching peak levels between 2-4 a.m and at a minimum between 6-10 pm. The variation is of the order of 50 %. Hence time of the day has influence on the measured serum TSH concentration.

USE:- TSH controls biosynthesis and release of thyroid harmones T4 & T3. It is a sensitive measure of thyroid function, especially useful in early or subclinical hypothyroidism, before the patient develops any clinical findings or goitre or any other thyroid function abnormality. **INCREASED LEVELS**:

1. Primary or untreated hypothyroidism, may vary from 3 times to more than 100 times normal depending on degree of hypofunction.

2.Hypothyroid patients receiving insufficient thyroid replacement therapy.

3. Hashimotos thyroiditis.

4.DRUGS: Amphetamines, Iodine containing agents and dopamine antagonist.

5.Neonatal period, increase in 1st 2-3 days of life due to post-natal surge.

DECREASED LEVELS:

1. Toxic multi-nodular goitre & Thyroiditis.

2. Over replacement of thyroid harmone in treatment of hypothyroidism.

3. Autonomously functioning Thyroid adenoma

4. Secondary pituatary or hypothalmic hypothyroidism

5. Acute psychiatric illness

6.Severe dehydration.

7.DRUGS: Glucocorticoids, Dopamine, Levodopa, T4 replacement therapy, Anti-thyroid drugs for thyrotoxicosis.



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NOT VALID FOR MEDICO LEGAL PURPOSE

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Test Name	Value	Unit	Biological Reference interval

8.Pregnancy: 1st and 2nd Trimester

LIMITATIONS:

1.TSH may be normal in central hypothyroidism, recent rapid correction of hyperthyroidism or hypothyroidism, pregnancy, phenytoin therapy. 2.Autoimmune disorders may produce spurious results.





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