



	Dr. Vinay Cł MD (Pathology & Chairman & Cor			(Pathology)
NAME	: Mrs. RADHA			
AGE/ GENDER	: 62 YRS/FEMALE		PATIENT ID	: 1563694
COLLECTED BY	:		REG. NO./LAB NO.	: 012407290003
REFERRED BY	:		REGISTRATION DATE	: 29/Jul/2024 06:04 AM
BARCODE NO.	:01514025		COLLECTION DATE	: 29/Jul/2024 08:36AM
CLIENT CODE.	: KOS DIAGNOSTIC LAB		REPORTING DATE	: 29/Jul/2024 08:52AM
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD,	, AMBALA CANTT		
Test Name		Value	Unit	Biological Reference interval
HAEMOGLOBIN (HB)		12.6	gm/dL	12.0 - 16.0
by CALORIMETRIC		12.0	gin/uL	12.0 - 10.0
tissues back to the lu A low hemoglobin lev ANEMIA (DECRESED I 1) Loss of blood (trau 2) Nutritional deficiel 3) Bone marrow prob 4) Suppression by red 5) Kidney failure 6) Abnormal hemogle POLYCYTHEMIA (INCR 1) People in higher a	ngs. rel is referred to as ANEMIA or lo HAEMOGLOBIN): Imatic injury, surgery, bleeding, ncy (iron, vitamin B12, folate) lems (replacement of bone mari blood cell synthesis by chemo obin structure (sickle cell anemi EASED HAEMOGLOBIN): Ititudes (Physiological)	ow red blood coun colon cancer or s row by cancer) therapy drugs	it. tomach ulcer)	odys tissues and returns carbon dioxide from t
4) Advanced lung dise5) Certain tumors6) A disorder of the b	ices a falsely rise in hemoglobin ease (for example, emphysema) one marrow known as polycythe	emia rubra vera,		e amount of oxygen available to the body by

KOS Diagnostic Lab (A Unit of KOS Healthcare)

NOTE: TEST CONDUCTED ON EDTA WHOLE BLOOD



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TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.

chemically raising the production of red blood cells).

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	Dr. Vinay Che MD (Pathology & Chairman & Cons	Microbiology)	Dr. Yugam MD CEO & Consultant	(Pathology)
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CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, A	AMBALA CANTT		
Test Name		Value	Unit	Biological Reference interval
	GĽ	YCOSYLATED HAEM	OGLOBIN (HBA1C)	
GLYCOSYLATED HAEM	OGLOBIN (HbA1c):	7.5 ^H	%	4.0 - 6.4
ESTIMATED AVERAGE		168.55 ^H	mg/dL	60.00 - 140.00
	AS PER AMERICAN DIAB	ETES ASSOCIATION (ADA)	:	
RE	FERENCE GROUP	GLYCOSYLATE	D HEMOGLOGIB (HBAIC) i	n %
Non diabetic Adults >= 18 years		<5.7		
	Risk (Prediabetes)	5.7 – 6.4		
Dia	gnosing Diabetes		>= 6.5	
			Age > 19 Years	
-		Goals of Therapy:	< 7.0	•
Therapeutic	goals for glycemic control	Actions Suggested:)
			Age < 19 Years	
		Goal of therapy:	<7.5	

COMMENTS:

TEST PERFORMED AT KOS DIAGNOSTIC LAB. AMBALA CANTT

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 appropriate. 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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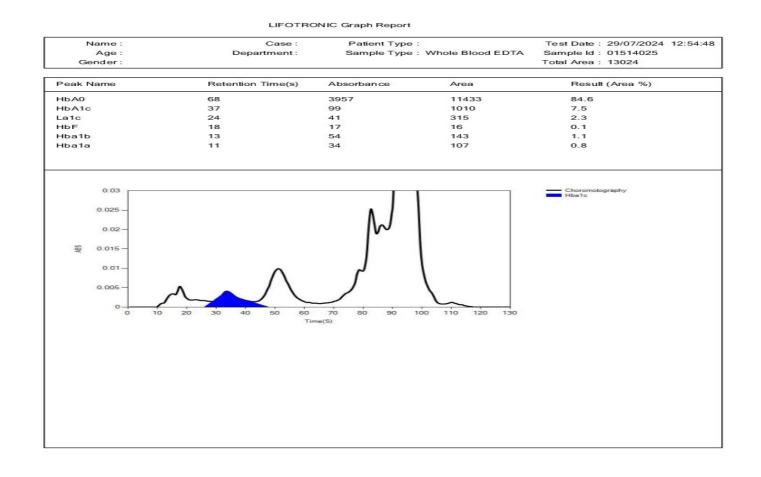








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Test Name	Valu	ue Unit	Biological Reference interval





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CLIENT CODE.	: KOS DIAGNOSTIC LAB	R	EPORTING DATE	: 29/Jul/2024 09:58AM
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD,	AMBALA CANTT		
Test Name		Value	Unit	Biological Reference interval
	CLIN	ICAL CHEMIST	RY/BIOCHEMISTR	Y
		GLUCOSE F	FASTING (F)	
GLUCOSE FASTING (by glucose oxidas	F): PLASMA se - peroxidase (god-pod)	113.78 ^H	mg/dL	NORMAL: < 100.0 PREDIABETIC: 100.0 - 125.0 DIABETIC: > 0R = 126.0
1. A fasting plasma g 2. A fasting plasma g test (after consumpti 3. A fasting plasma g	ion of 75 gms of glucose) is recor	considered normal. mg/dl is considered mmended for all suc is highly suggestive	as glucose intolerant or ch patients. of diabetic state. A repe	prediabetic. A fasting and post-prandial blood at post-prandial is strongly recommended for atory for diabetic state.





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CLIENT ADDRESS	: 6349/1, NICHOL	SON ROAD, AMBALA CANTT		
	_	Value	Unit	Biological Reference interval
Test Name				
Test Name		CHOLESTER	OL: SERUM	

NATIONAL LIPID ASSOCIATION RECOMMENDATIONS (NLA-2014)	CHOLESTEROL IN ADULTS (mg/dL)	CHOLESTEROL IN ADULTS (mg/dL)
DESIRABLE	< 200.0	< 170.0
BORDERLINE HIGH	200.0 – 239.0	171.0 - 199.0
HIGH	>= 240.0	>= 200.0

NOTE:

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.
 As per National Lipid association - 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.





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COLLECTED BY : REFERRED BY : BARCODE NO. : 01514025 CLIENT CODE. : KOS DIAC CLIENT ADDRESS : 6349/1, N Test Name TRIIODOTHYRONINE (T3): SERUN by CMIA (CHEMILUMINESCENT MICR	5 ENOSTIC LAB NICHOLSON ROAD, AMBALA CA	REG. NO./LAB NO. REGISTRATION DATE COLLECTION DATE REPORTING DATE	: 012407290003 : 29/Jul/2024 06:04 AM : 29/Jul/2024 08:36AM : 29/Jul/2024 10:18AM
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TRIIODOTHYRONINE (T3): SERUN by CMIA (CHEMILUMINESCENT MICR	Value	e Unit	Biological Reference interval
by CMIA (CHEMILUMINESCENT MICR			
by CMIA (CHEMILUMINESCENT MICR	ENI	DOCRINOLOGY	
by CMIA (CHEMILUMINESCENT MICR	THYROID F	UNCTION TEST: TOTAL	
		5 ng/mL	0.35 - 1.93
	OPARTICLE IMMUNOASSAY) 8.87	µgm/dL	4.87 - 12.60
by CMIA (CHEMILUMINESCENT MICR		μgin/uc	4.07 - 12.00
THYROID STIMULATING HORMC by CMIA (CHEMILUMINESCENT MICH IMMUNOASSAY)		23 ^H μlU/mL	0.35 - 5.50
3rd GENERATION, ULTRASENSITIVE			
INTERPRETATION:			om. The variation is of the order of 50%.Hence time o

ISH levels are subject to circadian 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 concentrations. TSH stimulates the production and secretion of the metabolically active hormones, thyroxine (T4) and trilodothyronine (T3). Failure at any level of regulation of the hypothalamic-pituitary-thyroid axis will result in either underproduction (hypothyroidism) or overproduction(hyperthyroidism) of T4 and/or T3.

CLINICAL CONDITION	T3	T4	TSH
Primary Hypothyroidism:	Reduced	Reduced	Increased (Significantly)
Subclinical Hypothyroidism:	Normal or Low Normal	Normal or Low Normal	High
Primary Hyperthyroidism:	Increased	Increased	Reduced (at times undetectable)
Subclinical Hyperthyroidism:	Normal or High Normal	Normal or High Normal	Reduced

LIMITATIONS:-

1. T3 and T4 circulates in reversibly bound form with Thyroid binding globulins (TBG), and to a lesser extent albumin and Thyroid binding Pre Albumin so conditions in which TBG and protein levels alter such as pregnancy, excess estrogens, androgens, anabolic steroids and glucocorticoids may falsely affect the T3 and T4 levels and may cause false thyroid values for thyroid function tests.

2. Normal levels of T4 can also be seen in Hyperthyroid patients with :T3 Thyrotoxicosis, Decreased binding capacity due to hypoproteinemia or ingestion of certain drugs (eg: phenytoin , salicylates).

3. Serum T4 levles in neonates and infants are higher than values in the normal adult , due to the increased concentration of TBG in neonate serum.

4. TSH may be normal in central hypothyroidism, recent rapid correction of hyperthyroidism or hypothroidism, pregnancy, phenytoin therapy.

TRIIODOTHYRONINE (T3)		THYROX	THYROXINE (T4)		ATING HORMONE (TSH)
Age	Refferance Range (ng/mL)	Age	Refferance Range (µg/dL)	Age	Reference Range (µIU/mL)
0-7 Days	0.20 - 2.65	0 - 7 Days	5.90 - 18.58	0 - 7 Days	2.43 - 24.3
7 Days - 3 Months	0.36 - 2.59	7 Days - 3 Months	6.39 - 17.66	7 Days - 3 Months	0.58 - 11.00
3 - 6 Months	0.51 - 2.52	3 - 6 Months	6.75 - 17.04	3 Days – 6 Months	0.70 - 8.40





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Test Name			Value	Unit	:	Biological Reference interva
6 - 12 Months	0.74 - 2.40	6 - 12 Months	7.10 - 16.16	6 – 12 Months	0.70 - 7.00	
1 - 10 Years	0.92 - 2.28	1 - 10 Years	6.00 - 13.80	1 – 10 Years	0.60 - 5.50	
11- 19 Years	0.35 - 1.93	11 - 19 Years	4.87- 13.20	11 – 19 Years	0.50 - 5.50	
> 20 years (Adults)	0.35 - 1.93	> 20 Years (Adults)	4.87 - 12.60	> 20 Years (Adults)	0.35-5.50	
	RECO	DMMENDATIONS OF TSH L	EVELS DURING PRE	GNANCY (µIU/mL)		
1st Trimester			0.10 - 2.50			
2nd Trimester			0.20 - 3.00			
	3rd Trimester		0.30 - 4.10			

INCREASED TSH LEVELS:

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

2.Hypothyroid patients receiving insufficient thyroid replacement therapy.

3.Hashimotos thyroiditis

4.DRUGS: Amphetamines, idonie containing agents & dopamine antagonist.

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

DECREASED TSH 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.

8.Pregnancy: 1st and 2nd Trimester

*** End Of Report ***





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