A PIONEER DIAGNOSTIC CENTRE

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

NAME	: Mr. JAGTAR SINGH							
AGE/ GENDER	: 56 YRS/MALE		PATIENT ID	: 1813599				
COLLECTED BY	:		REG. NO./LAB NO.	: 122504010002				
REFERRED BY	:		REGISTRATION DATE	: 01/Apr/2025 08:11 AM				
BARCODE NO.	: 12507829		COLLECTION DATE	: 01/Apr/2025 08:26AM				
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITUTE		REPORTING DATE	: 01/Apr/2025 01:24PM				
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBALA C	REG. N REGIST COLLER COLLER NAMBALA CITY - HARYANA Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value Value VV V V V V V V V V	RYANA					
Test Name	V	alue	Unit	Biological Reference interval				
	SWASTHY	A WE	LLNESS PANEL: 1	2				
	COMPLE	TE BL	OOD COUNT (CBC)					
RED BLOOD CEL	LS (RBCS) COUNT AND INDICES							
HAEMOGLOBIN (H by CALORIMETRIC	B)	14.4	gm/dL	12.0 - 17.0				
RED BLOOD CELL by HYDRO DYNAMIC F	(RBC) COUNT FOCUSING, ELECTRICAL IMPEDENCE	4.33	Millions/	cmm 3.50 - 5.00				
PACKED CELL VO	LUME (PCV) NUTOMATED HEMATOLOGY ANALYZER	40.4	%	40.0 - 54.0				
	LAR VOLUME (MCV) NUTOMATED HEMATOLOGY ANALYZER	93.2	fL	80.0 - 100.0				
	LAR HAEMOGLOBIN (MCH) NUTOMATED HEMATOLOGY ANALYZER	33.3	pg	27.0 - 34.0				
	LAR HEMOGLOBIN CONC. (MCHC) NUTOMATED HEMATOLOGY ANALYZER	35.7	g/dL	32.0 - 36.0				
	BUTION WIDTH (RDW-CV)	15.6	%	11.00 - 16.00				
	BUTION WIDTH (RDW-SD)	56	fL	35.0 - 56.0				
MENTZERS INDEX by CALCULATED		21.52	RATIO	BETA THALASSEMIA TRAIT 13.0				
				IRON DEFICIENCY ANEMIA: >13.0				
GREEN & KING IN by CALCULATED	DEX	94.1	RATIO	BETA THALASSEMIA TRAIT: <= 74.1				
				IRON DEFICIENCY ANEMIA: >= 74.1				
WHITE BLOOD C	ELLS (WBCS)							
TOTAL LEUCOCY	TE COUNT (TLC) y by sf cube & microscopy	8170	/cmm	4000 - 11000				
DIFFERENTIAL L	EUCOCYTE COUNT (DLC)							
		60	%	50 - 70				

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Test Name		Value	Unit	Biological Reference interva				
by FLOW CYTOMETRY	BY SF CUBE & MICROSCOPY							
LYMPHOCYTES by FLOW CYTOMETRY	BY SF CUBE & MICROSCOPY	33	%	20 - 40				
EOSINOPHILS by FLOW CYTOMETRY	BY SF CUBE & MICROSCOPY	0 ^L	%	1 - 6				
MONOCYTES by FLOW CYTOMETRY	Y BY SF CUBE & MICROSCOPY	7	%	2 - 12				
BASOPHILS by FLOW CYTOMETRY	Y BY SF CUBE & MICROSCOPY	0	%	0 - 1				
ABSOLUTE LEUK	OCYTES (WBC) COUNT							
ABSOLUTE NEUTR	OPHIL COUNT BY SF CUBE & MICROSCOPY	4902	/cmm	2000 - 7500				
ABSOLUTE LYMPH by FLOW CYTOMETRY	OCYTE COUNT ' BY SF CUBE & MICROSCOPY	2696 ^L	/cmm	800 - 4900				
ABSOLUTE EOSINO	OPHIL COUNT ' BY SF CUBE & MICROSCOPY	0 ^L	/cmm	40 - 440				
ABSOLUTE MONO	CYTE COUNT ' BY SF CUBE & MICROSCOPY	572	/cmm	80 - 880				
ABSOLUTE BASOP	HIL COUNT ' BY SF CUBE & MICROSCOPY	0	/cmm	0 - 110				
PLATELETS AND (OTHER PLATELET PREDICTIV	E MARKERS.						
PLATELET COUNT by HYDRO DYNAMIC F	' (PLT) OCUSING, ELECTRICAL IMPEDENCE	264000	/cmm	150000 - 450000				
PLATELETCRIT (P	CT) OCUSING, ELECTRICAL IMPEDENCE	0.2	%	0.10 - 0.36				
MEAN PLATELET V by hydro dynamic f	VOLUME (MPV) OCUSING, ELECTRICAL IMPEDENCE	8	fL	6.50 - 12.0				
	CELL COUNT (P-LCC) OCUSING, ELECTRICAL IMPEDENCE	33000	/cmm	30000 - 90000				
	CELL RATIO (P-LCR) OCUSING, ELECTRICAL IMPEDENCE	12.6	%	11.0 - 45.0				
by HYDRO DYNAMIC F	BUTION WIDTH (PDW) ocusing, electrical impedence CTED ON EDTA WHOLE BLOOD	15.5	%	15.0 - 17.0				



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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMB	ALA CITY - HARYAN	IA	
Test Name		Value	Unit	Biological Reference interval
	ERYTHROO	CYTE SEDIMEN	NTATION RATE	(ESR)
	EDIMENTATION RATE (ESR) GATION BY CAPILLARY PHOTOMETRY	5	mm/1st h	r 0 - 20
1. ESR is a non-specif immune disease, but 2. An ESR can be affe as C-reactive protein	does not tell the health practitione cted by other conditions besides inf	r exactly where the flammation. For this	inflammation is in the reason, the ESR is typ	pically used in conjunction with other test suc
3. This test may also systemic lupus erythe CONDITION WITH LOV	ematosus	and response to the	erapy in both of the al	bove diseases as well as some others, such as

A low ESR can be seen with conditions that inhibit the normal sedimentation of red blood cells, such as a high red blood cell count

(polycythaemia), significantly high white blood cell count (leucocytosis), and some protein abnormalities. Some changes in red cell shape (such as sickle cells in sickle cell anaemia) also lower the ESR. NOTE:

1. ESR and C - reactive protein (C-RP) are both markers of inflammation.
2. Generally, ESR does not change as rapidly as does CRP, either at the start of inflammation or as it resolves.
3. CRP is not affected by as many other factors as is ESR, making it a better marker of inflammation.

If the ESR is elevated, it is typically a result of two types of proteins, globulins or fibrinogen.
Women tend to have a higher ESR, and menstruation and pregnancy can cause temporary elevations.

6. Drugs such as dextran, methyldopa, oral contraceptives, penicillamine procainamide, theophylline, and vitamin A can increase ESR, while aspirin, cortisone, and quinine may decrease it



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Test Name		Value	T I:4	Dislagical Deference interval
Test Name		Value	Unit	Biological Reference interval
	CLINI	CAL CHEMISTR	Y/BIOCHEMIS	TRY
		GLUCOSE FA	STING (F)	
GLUCOSE FASTIN by GLUCOSE OXIDAS	G (F): PLASMA E - PEROXIDASE (GOD-POD)	97.3	mg/dL	NORMAL: < 100.0 PREDIABETIC: 100.0 - 125.0 DIABETIC: > 0R = 126.0
INTERPRETATION				

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.





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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, A	MBALA CITY - HA	RYANA	
Test Name		Value	Unit	Biological Reference interval
		LIPID PRO	OFILE : BASIC	
CHOLESTEROL TO by CHOLESTEROL OX		272.38 ^H	mg/dL	OPTIMAL: < 200.0 BORDERLINE HIGH: 200.0 - 239.0 HIGH CHOLESTEROL: > OR = 240.0
TRIGLYCERIDES: 5 by GLYCEROL PHOSF	SERUM PHATE OXIDASE (ENZYMATIC)	182.68 ^H	mg/dL	OPTIMAL: < 150.0 BORDERLINE HIGH: 150.0 - 199.0 HIGH: 200.0 - 499.0 VERY HIGH: > OR = 500.0
HDL CHOLESTER(by SELECTIVE INHIBIT	DL (DIRECT): SERUM 70N	116.18 ^H	mg/dL	LOW HDL: < 30.0 BORDERLINE HIGH HDL: 30.0 60.0
LDL CHOLESTERC		119.66	mg/dL	HIGH HDL: > OR = 60.0 OPTIMAL: < 100.0 ABOVE OPTIMAL: 100.0 - 129 BORDERLINE HIGH: 130.0 - 159.0 HIGH: 160.0 - 189.0
NON HDL CHOLES by CALCULATED, SPE		156.2 ^H	mg/dL	VERY HIGH: > OR = 190.0 OPTIMAL: < 130.0 ABOVE OPTIMAL: 130.0 - 159 BORDERLINE HIGH: 160.0 - 189.0 HIGH: 190.0 - 219.0
VLDL CHOLESTER		36.54	mg/dL	VERY HIGH: > OR = 220.0 0.00 - 45.00
TOTAL LIPIDS: SE by CALCULATED, SPE	RUM	727.44 ^H	mg/dL	350.00 - 700.00
CHOLESTEROL/HI		2.34	RATIO	LOW RISK: 3.30 - 4.40 AVERAGE RISK: 4.50 - 7.0

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Test Name	Value	Unit	Biological Reference interval
			MODERATE RISK: 7.10 - 11.0

LDL/HDL RATIO: SERUM	1.03	RATIO	HIGH RISK: > 11.0 LOW RISK: 0.50 - 3.0
by CALCULATED, SPECTROPHOTOMETRY	1.05	KAHO	MODERATE RISK: 3.10 - 6.0 HIGH RISK: > 6.0
TRIGLYCERIDES/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	1.57 ^L	RATIO	3.00 - 5.00

INTERPRETATION:

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 HDL

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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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBALA	A CITY - HARY	YANA	•				
Test Name		Value	Unit	Biological Reference interval				
	LIVER FU	JNCTION	TEST (COMPLETE)				
BILIRUBIN TOTAL by DIAZOTIZATION, SF	: SERUM PECTROPHOTOMETRY	0.61	mg/dL	INFANT: 0.20 - 8.00 ADULT: 0.00 - 1.20				
	T (CONJUGATED): SERUM	0.22	mg/dL	0.00 - 0.40				
BILIRUBIN INDIRE	ECT (UNCONJUGATED): SERUM	0.39	mg/dL	0.10 - 1.00				
SGOT/AST: SERUN by IFCC, WITHOUT PY	I RIDOXAL PHOSPHATE	38.27	U/L	7.00 - 45.00				
SGPT/ALT: SERUM by IFCC, WITHOUT PY	I RIDOXAL PHOSPHATE	22.46		0.00 - 49.00				
AST/ALT RATIO: S		1.7	RATIO	0.00 - 46.00				
ALKALINE PHOSPI by PARA NITROPHEN [®] PROPANOL	HATASE: SERUM YL PHOSPHATASE BY AMINO METHYL	87.97	U/L	40.0 - 130.0				
GAMMA GLUTAM by SZASZ, SPECTROF	YL TRANSFERASE (GGT): SERUM Phtometry	85.87 ^H	U/L	0.00 - 55.0				
TOTAL PROTEINS by BIURET, SPECTRO		5.67 ^L	gm/dL	6.20 - 8.00				
ALBUMIN: SERUM by BROMOCRESOL G		3.95	gm/dL	3.50 - 5.50				
GLOBULIN: SERUN by CALCULATED, SPE		1.72 ^L	gm/dL	2.30 - 3.50				
A : G RATIO: SERU by CALCULATED, SPE		2.3 ^H	RATIO	1.00 - 2.00				

INTERPRETATION

NOTE: • To be correlated in individuals having SGOT and SGPT values higher than Normal Referance Range. USE: • Differential diagnosis of diseases of hepatobiliary system and pancreas.

: Mr. JAGTAR SINGH

INCREASED:

DRUG HEPATOTOXICITY	>2
ALCOHOLIC HEPATITIS	> 2 (Highly Suggestive)
CIRRHOSIS	1.4 - 2.0
INTRAHEPATIC CHOLESTATIS	> 1.5





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Test Name	Value	Unit	Biological Reference interval
HEPATOCELLULAR CARCINOMA & CHRONIC HEPATITIS		> 1.3 (Slightly Increased)	
DECREASED:			

1. Acute Hepatitis due to virus, drugs, toxins (with AST increased 3 to 10 times upper limit of normal)

2. Extra Hepatic cholestatis: 0.8 (normal or slightly decreased).

PRO	G	NO	ST	ПC	S	IGN	lif	IC	:A	١Ν	ICE	•

NORMAL	< 0.65
GOOD PROGNOSTIC SIGN	0.3 - 0.6
POOR PROGNOSTIC SIGN	1.2 - 1.6



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Test Name		Value	Unit	Biological Reference interval	
	KIDNEY	FUNCTI	ON TEST (COMPLET	F)	
UREA: SERUM		17.81		10.00 - 50.00	
	ATE DEHYDROGENASE (GLDH)	17.81	mg/dL	10.00 - 50.00	
CREATININE: SERU	JM	0.94	mg/dL	0.40 - 1.40	
BLOOD UREA NITH by CALCULATED, SPE	ROGEN (BUN): SERUM CTROPHOTOMETRY	8.32	mg/dL	7.0 - 25.0	
BLOOD UREA NITH RATIO: SERUM by CALCULATED, SPEN	ROGEN (BUN)/CREATININE	8.85 ^L	RATIO	10.0 - 20.0	
UREA/CREATININE by CALCULATED, SPE	E RATIO: SERUM	18.95	RATIO		
URIC ACID: SERUN by URICASE - OXIDASI		4.71	mg/dL	3.60 - 7.70	
CALCIUM: SERUM by ARSENAZO III, SPEC	CTROPHOTOMETRY	9.08	mg/dL	8.50 - 10.60	
PHOSPHOROUS: SE by PHOSPHOMOLYBD	ERUM ATE, SPECTROPHOTOMETRY	2.75	mg/dL	2.30 - 4.70	
<u>ELECTROLYTES</u>					
SODIUM: SERUM by ISE (ION SELECTIVE	E ELECTRODE)	145.8	mmol/L	135.0 - 150.0	
POTASSIUM: SERU by ISE (ION SELECTIVE		3.67	mmol/L	3.50 - 5.00	
CHLORIDE: SERUN	E ELECTRODE)	109.35	mmol/L	90.0 - 110.0	
	MERULAR FILTERATION RAT				
ESTIMATED GLON (eGFR): SERUM by calculated INTERPRETATION:	IERULAR FILTERATION RATE	95.1			
To differentiate betwe	een pre- and post renal azotemia.				

INCREASED RATIO (>20:1) WITH NORMAL CREATININE:

1. Prerenal azotemia (BUN rises without increase in creatinine) e.g. heart failure, salt depletion, dehydration, blood loss) due to decreased glomerular filtration rate.



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Test Name	Value	Unit	Biological Reference interval

- 4. High protein intake.
- 5. Impaired renal function plus
- 6. Excess protein intake or production or tissue breakdown (e.g. infection, GI bleeding, thyrotoxicosis, Cushing's syndrome, high protein diet,
- burns, surgery, cachexia, high fever).
- 7. Urine reabsorption (e.g. ureter colostomy)
- 8. Reduced muscle mass (subnormal creatinine production)
- 9. Certain drugs (e.g. tetracycline, glucocorticoids)

INCREASED RATIO (>20:1) WITH ELEVATED CREATININE LEVELS:

1. Postrenal azotemia (BUN rises disproportionately more than creatinine) (e.g. obstructive uropathy).

2. Prerenal azotemia superimposed on renal disease.

DECREASED RATIO (<10:1) WITH DECREASED BUN :

- 1. Acute tubular necrosis.
- 2. Low protein diet and starvation.
- 3. Severe liver disease.
- 4. Other causes of decreased urea synthesis.
- 5. Repeated dialysis (urea rather than creatinine diffuses out of extracellular fluid).
- 6. Inherited hyperammonemias (urea is virtually absent in blood).
- 7. SIADH (syndrome of inappropiate antidiuretic harmone) due to tubular secretion of urea.
- 8. Pregnancy.

DECREASED RATIO (<10:1) WITH INCREASED CREATININE:

- 1. Phenacimide therapy (accelerates conversion of creatine to creatinine).
- 2. Rhabdomyolysis (releases muscle creatinine).
- 3. Muscular patients who develop renal failure.

INAPPROPIATE RATIO:

1. Diabetic ketoacidosis (acetoacetate causes false increase in creatinine with certain methodologies, resulting in normal ratio when dehydration should produce an increased BUN/creatinine ratio).

2. Cephalosporin therapy (interferes with creatinine measurement).

CKD STAGE	DESCRIPTION	GFR (mL/min/1.73m2)	ASSOCIATED FINDINGS
G1	Normal kidney function	>90	No proteinuria
G2	Kidney damage with normal or high GFR	>90	Presence of Protein , Albumin or cast in urine
G3a	Mild decrease in GFR	60 -89	
G3b	Moderate decrease in GFR	30-59	
G4	Severe decrease in GFR	15-29	
G5	Kidney failure	<15	





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A PIONEER DIAGNOSTIC CENTRE

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NAME	: Mr. JAGTAR SINGH		
AGE/ GENDER	: 56 YRS/MALE	PATIENT ID	: 1813599
COLLECTED BY	:	REG. NO./LAB NO.	: 122504010002
REFERRED BY	:	REGISTRATION DATE	: 01/Apr/2025 08:11 AM
BARCODE NO.	: 12507829	COLLECTION DATE	:01/Apr/202508:26AM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITUTE	REPORTING DATE	: 01/Apr/2025 04:41PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBALA CITY - I	HARYANA	

Test Name	Value	Unit	Biological Reference interval

COMMENTS:

Estimated Glomerular filtration rate (eGFR) is the sum of filtration rates in all functioning nephrons and so an estimation of the GFR provides a measure of functioning nephrons of the kidney.
eGFR calculated using the 2009 CKD-EPI creatinine equation and GFR category reported as per KDIGO guideline 2012

3. In patients, with eGFR creatinine between 45-59 ml/min/1.73 m2 (G3) and without any marker of Kidney damage, It is recommended to measure eGFR with Cystatin C for confirmation of CKD

4. eGFR category G1 OR G2 does not fullfill the criteria for CKD, in the absence of evidence of Kidney Damage 5. In a suspected case of Acute Kidney Injury (AKI), measurement of eGFR should be done after 48-96 hours of any Intervention or procedure 6. eGFR calculated by Serum Creatinine may be less accurate due to certain factors like Race, Muscle Mass, Diet, Certain Drugs. In such cases, eGFR should be calculated using Serum Cystatin C 7. A decrease in eGFR implies either progressive renal disease, or a reversible process causing decreased nephron function (eg, severe dehydration).

ADVICE:

KDIGO guideline, 2012 recommends Chronic Kidney Disease (CKD) should be classified based on cause, eGFR category and Albuminuria (ACR) category. GFR & ACR category combined together reflect risk of progression and helps Clinician to identify the individual who are progressing at more rapid rate than anticipated



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BARCODE NO.	: 12507829	COL	LECTION DATE	: 01/Apr/2025 08:26AM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITUT	ГЕ Rep	ORTING DATE	:01/Apr/2025 01:24PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBAL	A CITY - HARYAN	NA	
Test Name		Value	Unit	Biological Reference interval
		ENDOCRIN	OLOGY	
	THYRO	ID FUNCTIO	N TEST: TOTAL	
TRIIODOTHYRON by CMIA (CHEMILUMIN	INE (T3): SERUM	1.322	ng/mL	0.35 - 1.93
THYROXINE (T4): by CMIA (CHEMILUMIN	SERUM NESCENT MICROPARTICLE IMMUNOASSAY)	8.65	µgm/dL	4.87 - 12.60
	ATING HORMONE (TSH): SERUM	0.821	µIU/mL	0.35 - 5.50

TSH 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 triiodothyronine (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 (e.g.: phenytoin , salicylates).

3. Serum T4 levels 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 hypothyroidism, pregnancy, phenytoin therapy.

TRIIODOTH	(RONINE (T3)	THYROXINE (T4)		THYROID STIMULATING HORMONE (TS	
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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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBALA CITY - H	IARYANA	

	.10 - 16.16 .00 - 13.80	6 – 12 Months 1 – 10 Years	0.70 - 7.00	
1 - 10 Years 0.92 - 2.28 1 - 10 Years 6.0	.00 - 13.80	1 – 10 Years	0.40 5.50	
			0.60 - 5.50	
11- 19 Years 0.35 - 1.93 11 - 19 Years 4.8	.87- 13.20	11 – 19 Years	0.50 - 5.50	
> 20 years (Adults) 0.35 - 1.93 > 20 Years (Adults) 4.8	.87 - 12.60	> 20 Years (Adults)	0.35- 5.50	
RECOMMENDATIONS OF TSH LEVELS	S DURING PREGN	ANCY (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, iodine 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 goiter & Thyroiditis.

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

3. Autonomously functioning Thyroid adenoma

4. Secondary pituitary or hypothalamic 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



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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AM	IBALA CITY - HARY	ANA	
Test Name		Value	Unit	Biological Reference interv
		CLINICAL PA	THOLOGY	
	URINE ROU	TINE & MICRO	DSCOPIC EXAMI	NATION
PHYSICAL EXAMI	INATION NATION			
QUANTITY RECIEN by DIP STICK/REFLEC	/ED TANCE SPECTROPHOTOMETRY	10	ml	
COLOUR by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	PALE YELLO	W	PALE YELLOW
TRANSPARANCY by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	CLEAR		CLEAR
SPECIFIC GRAVITY by DIP STICK/REFLEC	Y TANCE SPECTROPHOTOMETRY	>=1.030		1.002 - 1.030
CHEMICAL EXAM	<u>IINATION</u>			
REACTION by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	ACIDIC		
PROTEIN by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	Negative		NEGATIVE (-ve)
SUGAR by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	Negative		NEGATIVE (-ve)
pH	TANCE SPECTROPHOTOMETRY	5.5		5.0 - 7.5
BILIRUBIN by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	Negative		NEGATIVE (-ve)
NITRITE	TANCE SPECTROPHOTOMETRY.	Negative		NEGATIVE (-ve)
UROBILINOGEN	TANCE SPECTROPHOTOMETRY	Normal	EU/dL	0.2 - 1.0
KETONE BODIES	TANCE SPECTROPHOTOMETRY	Negative		NEGATIVE (-ve)
BLOOD by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	Negative		NEGATIVE (-ve)
ASCORBIC ACID	TANCE SPECTROPHOTOMETRY	NEGATIVE ((-ve)	NEGATIVE (-ve)



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NEGATIVE (-ve)

NEGATIVE (-ve)

NEGATIVE (-ve)

ABSENT

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Test Name		Value	Unit	Biological Reference interval
RED BLOOD CELLS (RBCs) by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT		NEGATIVE (-ve)	/HPF	0 - 3
PUS CELLS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT		2-4	/HPF	0 - 5
EPITHELIAL CELLS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT		1-3	/HPF	ABSENT
CRYSTALS		NEGATIVE (-ve)		NEGATIVE (-ve)

by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT CASTS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT BACTERIA

by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT OTHERS

by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT TRICHOMONAS VAGINALIS (PROTOZOA)

by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT

* * * End Of Report *

ABSENT

NEGATIVE (-ve)

NEGATIVE (-ve)

NEGATIVE (-ve)





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