A PIONEER DIAGNOSTIC CENTRE

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

NAME	: Mr. JAGDEEP			
AGE/ GENDER	: 30 YRS/MALE		PATIENT ID	: 1708497
COLLECTED BY	:		REG. NO./LAB NO.	: 122412250009
REFERRED BY	:		<b>REGISTRATION DATE</b>	: 25/Dec/2024 02:02 PM
BARCODE NO.	: 12506290		<b>COLLECTION DATE</b>	: 25/Dec/2024 02:14PM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITU	TE	<b>REPORTING DATE</b>	: 25/Dec/2024 04:14PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBAL	A CITY - HA	RYANA	
Test Name		Value	Unit	Biological Reference interval
	SWASTI	HYA WE	LLNESS PANEL: 1.2	
	COMP	PLETE BL	OOD COUNT (CBC)	
RED BLOOD CELLS	(RBCS) COUNT AND INDICES			
HAEMOGLOBIN (HI	B)	16.6	gm/dL	12.0 - 17.0
RED BLOOD CELL (	RBC) COUNT ocusing, electrical impedence	4.72	Millions/	cmm 3.50 - 5.00
PACKED CELL VOLU	JME (PCV) utomated hematology analyzer	48.4	%	40.0 - 54.0
MEAN CORPUSCULA by CALCULATED BY A	AR VOLUME (MCV) utomated hematology analyzer	102.4 <sup>H</sup>	KR fl	80.0 - 100.0
	AR HAEMOGLOBIN (MCH) utomated hematology analyzer	35.2 <sup>H</sup>	pg	27.0 - 34.0
	AR HEMOGLOBIN CONC. (MCHC) UTOMATED HEMATOLOGY ANALYZER	34.4	g/dL	32.0 - 36.0
by CALCULATED BY A	UTION WIDTH (RDW-CV) utomated hematology analyzer	13.4	%	11.00 - 16.00
	UTION WIDTH (RDW-SD) utomated hematology analyzer	51.7	fL	35.0 - 56.0
MENTZERS INDEX by CALCULATED		21.69	RATIO	BETA THALASSEMIA TRAIT: < 13.0 IRON DEFICIENCY ANEMIA: >13.0
GREEN & KING IND by CALCULATED		29.1	RATIO	BETA THALASSEMIA TRAIT:< 65.0 IRON DEFICIENCY ANEMIA: > 65.0
WHITE BLOOD CEI				
	COUNT (TLC) / by sf cube & microscopy UCOCYTE COUNT (DLC)	8600	/cmm	4000 - 11000
NEUTROPHILS	Y BY SF CUBE & MICROSCOPY	59	%	50 - 70
LYMPHOCYTES		33	%	20 - 40

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Test Name		Value	Unit	Biological Reference interval
by FLOW CYTOMETRY	' BY SF CUBE & MICROSCOPY			
EOSINOPHILS	BY SF CUBE & MICROSCOPY	1	%	1 - 6
MONOCYTES		7	%	2 - 12
-	BY SF CUBE & MICROSCOPY	0	0/	0.1
BASOPHILS by FLOW CYTOMETRY	BY SF CUBE & MICROSCOPY	0	%	0 - 1
ABSOLUTE LEUKO	<u>CYTES (WBC) COUNT</u>			
ABSOLUTE NEUTRO		5074	/cmm	2000 - 7500
by FLOW CYTOMETRY ABSOLUTE LYMPH	Y BY SF CUBE & MICROSCOPY	2838 <sup>L</sup>	/cmm	800 - 4900
	Y BY SF CUBE & MICROSCOPY		K R	000 1000
ABSOLUTE EOSINO	PHIL COUNT ' by sf cube & microscopy	86	/cmm	40 - 440
ABSOLUTE MONOC		602	/cmm	80 - 880
	BY SF CUBE & MICROSCOPY	0	1	0 110
ABSOLUTE BASOPH by FLOW CYTOMETRY	11L COUN I ' BY SF CUBE & MICROSCOPY	0	/cmm	0 - 110
PLATELETS AND O	THER PLATELET PREDICTIVE	MARKERS.		
PLATELET COUNT (	(PLT) OCUSING, ELECTRICAL IMPEDENCE	176000	/cmm	150000 - 450000
PLATELETCRIT (PC	T) OCUSING, ELECTRICAL IMPEDENCE	0.22	%	0.10 - 0.36
MEAN PLATELET V	OLUME (MPV)	12 <sup>H</sup>	fL	6.50 - 12.0
-	OCUSING, ELECTRICAL IMPEDENCE CELL COUNT (P-LCC)	76000	/cmm	30000 - 90000
	JELL COUNT (P-LCC) OCUSING, ELECTRICAL IMPEDENCE	10000	/ cmm	20000 - 20000
	CELL RATIO (P-LCR) OCUSING, ELECTRICAL IMPEDENCE	43.1	%	11.0 - 45.0
by HYDRO DYNAMIC F	UTION WIDTH (PDW) OCUSING, ELECTRICAL IMPEDENCE CTED ON EDTA WHOLE BLOOD	16.7	%	15.0 - 17.0



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CLIENT ADDRESS					
Test Name		Value	Unit	Biological Reference interval	
	ERYTHE	ROCYTE SEDIME	NTATION RATE (H	ESR)	
ERYTHROCYTE SE	DIMENTATION RATE (ESR)	5	mm/1st ]	•	
by RED CELL AGGRE	GATION BY CAPILLARY PHOTOMET				
immune disease, but 2. An ESR can be affe as C-reactive protein	does not tell the health practitic ected by other conditions besides be used to monitor disease activ ematosus	oner exactly where the inflammation. For th	e inflammation is in the is reason, the ESR is typ	on associated with infection, cancer and auto body or what is causing it. bically used in conjunction with other test suc bove diseases as well as some others, such as	
A low ESR can be see (polycythaemia), sigi	en with conditions that inhibit the	ount (leucocytosis), a	on of red blood cells, sund some protein abnor	uch as a high red blood cell count malities. Some changes in red cell shape (su	
1. ESR and C - reactiv					

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



TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT



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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, A	MBALA CITY - HARYAN	JA		
Test Name		Value	Unit	Biological Reference ir	terva
Test Name		Value	Unit	Diological Meterence II	
	CLINI	CAL CHEMISTRY	/BIOCHEMIST	RY	
		GLUCOSE FAS	TING (F)		

2. 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. 3. 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 ADDRESS	: NASIRPUR, HISSAR ROAD, AI	MBALA CITY - HA	RYANA	
Test Name		Value	Unit	Biological Reference interval
		LIPID PR	OFILE : BASIC	
CHOLESTEROL TO by CHOLESTEROL O		233.51 <sup>H</sup>	mg/dL	OPTIMAL: < 200.0 BORDERLINE HIGH: 200.0 - 239.0 HIGH CHOLESTEROL: > OR = 240.0
TRIGLYCERIDES: S by GLYCEROL PHOSF	ERUM PHATE OXIDASE (ENZYMATIC)	132.3	mg/dL	OPTIMAL: < 150.0 BORDERLINE HIGH: 150.0 - 199.0 HIGH: 200.0 - 499.0
HDL CHOLESTERO by SELECTIVE INHIBIT	L (DIRECT): SERUM Ton	52.41	mg/dL	VERY HIGH: > OR = 500.0 LOW HDL: < 30.0 BORDERLINE HIGH HDL: 30.0 60.0 HIGH HDL: > OR = 60.0
LDL CHOLESTERO by CALCULATED, SPE		154.64 <sup>H</sup>	mg/dL	OPTIMAL: < 100.0 ABOVE OPTIMAL: 100.0 - 129 BORDERLINE HIGH: 130.0 - 159.0 HIGH: 160.0 - 189.0 VERY HIGH: > OR = 190.0
NON HDL CHOLES' by calculated, spe		181.1 <sup>H</sup>	mg/dL	OPTIMAL: < 130.0 ABOVE OPTIMAL: 130.0 - 159 BORDERLINE HIGH: 160.0 - 189.0 HIGH: 190.0 - 219.0 VERY HIGH: > OR = 220.0
VLDL CHOLESTER		26.46	mg/dL	0.00 - 45.00
TOTAL LIPIDS: SEF by CALCULATED, SPE	RUM	599.32	mg/dL	350.00 - 700.00
CHOLESTEROL/HI		4.46 <sup>H</sup>	RATIO	LOW RISK: 3.30 - 4.40 AVERAGE RISK: 4.50 - 7.0 MODERATE RISK: 7.10 - 11.0 HIGH RISK: > 11.0

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

Test Name	Value	Unit	<b>Biological Reference interval</b>
LDL/HDL RATIO: SERUM by calculated, spectrophotometry	2.95	RATIO	LOW RISK: 0.50 - 3.0 MODERATE RISK: 3.10 - 6.0 HIGH RISK: > 6.0
TRIGLYCERIDES/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	2.52 <sup>L</sup>	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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: Mr. JAGDEEP

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Test Name		Value	Unit	Biological Reference interval
	LIVER	FUNCTION	N TEST (COMPLETE)	
BILIRUBIN TOTAL by diazotization, si	: SERUM PECTROPHOTOMETRY	2.05 <sup>H</sup>	mg/dL	INFANT: 0.20 - 8.00 ADULT: 0.00 - 1.20
	C (CONJUGATED): SERUM	0.77 <sup>H</sup>	mg/dL	0.00 - 0.40
BILIRUBIN INDIRE by CALCULATED, SPE	CT (UNCONJUGATED): SERUM	1.28 <sup>H</sup>	mg/dL	0.10 - 1.00
SGOT/AST: SERUM by IFCC, WITHOUT PY	RIDOXAL PHOSPHATE	135.79 <sup>H</sup>	U/L	7.00 - 45.00
SGPT/ALT: SERUM by IFCC, WITHOUT PY	RIDOXAL PHOSPHATE	141.51 <sup>H</sup>	KR U/L	0.00 - 49.00
AST/ALT RATIO: S by CALCULATED, SPE		0.96	RATIO	0.00 - 46.00
ALKALINE PHOSPI by para nitrophen propanol	HATASE: SERUM YL PHOSPHATASE BY AMINO METHYL	116.81	U/L	40.0 - 130.0
GAMMA GLUTAMY by szasz, spectrof	L TRANSFERASE (GGT): SERUM PHTOMETRY	505.63 <sup>H</sup>	U/L	0.00 - 55.0
TOTAL PROTEINS: by biuret, spectro		6.82	gm/dL	6.20 - 8.00
ALBUMIN: SERUM by bromocresol g		4.34	gm/dL	3.50 - 5.50
GLOBULIN: SERUM by CALCULATED, SPE		2.48	gm/dL	2.30 - 3.50
A : G RATIO: SERUN by CALCULATED, SPE		1.75	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.

INCREASED:

DRUG HEPATOTOXICITY	> 2
ALCOHOLIC HEPATITIS	> 2 (Highly Suggestive)
CIRRHOSIS	1.4 - 2.0
INTRAHEPATIC CHOLESTATIS	> 1.5
HEPATOCELLULAR CARCINOMA & CHRONIC HEPATITIS	> 1.3 (Slightly Increased)





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#### **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).

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



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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AME	BALA CITY - H	IARYANA		
Test Name		Value	Unit	Biological Reference interval	
	KIDNE	Y FUNCTI	ION TEST (COMPLETE)	)	
UREA: SERUM by UREASE - GLUTAM	IATE DEHYDROGENASE (GLDH)	16.61	mg/dL	10.00 - 50.00	
CREATININE: SERU by ENZYMATIC, SPEC		0.91	mg/dL	0.40 - 1.40	
BLOOD UREA NITR by CALCULATED, SPE	COGEN (BUN): SERUM	7.76	mg/dL	7.0 - 25.0	
	ROGEN (BUN)/CREATININE	8.53 <sup>L</sup>	RATIO	10.0 - 20.0	
UREA/CREATININI by CALCULATED, SPE	E RATIO: SERUM	1 <mark>8.25</mark>	RATIO		
URIC ACID: SERUM	[	9.06 <sup>H</sup>	mg/dL	3.60 - 7.70	
CALCIUM: SERUM by ARSENAZO III, SPE		9.06	mg/dL	8.50 - 10.60	
PHOSPHOROUS: SE by PHOSPHOMOLYBE		2.66	mg/dL	2.30 - 4.70	
ELECTROLYTES SODIUM: SERUM by ISE (ION SELECTIV		138.4	mmol/L	135.0 - 150.0	
POTASSIUM: SERUI by ISE (ION SELECTIV	M	5	mmol/L	3.50 - 5.00	
CHLORIDE: SERUM by ISE (ION SELECTIV	Ĩ	103.8	mmol/L	90.0 - 110.0	
	IERULAR FILTERATION RATE				
ESTIMATED GLOM (eGFR): SERUM by CALCULATED INTERPRETATION:	ERULAR FILTERATION RATE	116.3			

**INTERPRETATION:** 

To differentiate between 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.

2. Catabolic states with increased tissue breakdown.

3. GI haemorrhage.



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Test Name		Value Unit	Biological Reference interval
8. Reduced muscle m 9. Certain drugs (e.g. INCREASED RATIO (>2 1. Postrenal azotemia DECREASED RATIO (<1 1. Acute tubular necr 2. Low protein diet ar 3. Severe liver disease 4. Other causes of de 5. Repeated dialysis ( 6. Inherited hyperam 7. SIADH (syndrome c 8. Pregnancy. DECREASED RATIO (<1 1. Phenacimide thera 2. Rhabdomyolysis (r 3. Muscular patients INAPPROPIATE RATIO 1. Diabetic ketoacido should produce an in 2. Cephalosporin ther	nd starvation. e. creased urea synthesis. furea rather than creatinine diffuses ou monemias (urea is virtually absent in b of inappropiate antidiuretic harmone) di <b>10:1) WITH INCREASED CREATININE:</b> py (accelerates conversion of creatine t eleases muscle creatinine). who develop renal failure. :	an creatinine) (e.g. obstructive t of extracellular fluid). lood). ue to tubular secretion of urea. to creatinine). in creatinine with certain meth	uropathy). odologies,resulting in normal ratio when dehydratic ASSOCIATED FINDINGS
G1	Normal kidney function	SFR ( mL/min/1./3m2 ) >90	No proteinuria
G1 G2	Kidney damage with	>90	Presence of Protein ,
	normal or high GFR	~ / 0	_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	



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Kidney failure

<15

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G5





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Test Name	Value	Unit	<b>Biological Reference interval</b>

COMMENTS:

1. 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. 2. 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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**PKR JAIN HEALTHCARE INSTITUTE** 

NASIRPUR, Hissar Road, AMBALA CITY- (Haryana)

A PIONEER DIAGNOSTIC CENTRE

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NAME	: Mr. JAGDEEP			
AGE/ GENDER	: 30 YRS/MALE	РАТ	TENT ID	: 1708497
COLLECTED BY	:	REG	. NO./LAB NO.	: 122412250009
REFERRED BY	:	REG	ISTRATION DATE	: 25/Dec/2024 02:02 PM
BARCODE NO.	: 12506290	COL	LECTION DATE	: 25/Dec/2024 02:14PM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITU	TE <b>Rep</b>	ORTING DATE	: 25/Dec/2024 04:14PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBAL	A CITY - HARYAI	NA	
Test Name		Value	Unit	Biological Reference interva
		value	Unit	Diviogical weier ence miter va
		ENDOCRIN	OLOGY	
	THYRO		OLOGY N TEST: TOTAL	_
TRIIODOTHYRONII				0.35 - 1.93
TRIIODOTHYRONII by CMIA (CHEMILUMIN THYROXINE (T4): S	NE (T3): SERUM ESCENT MICROPARTICLE IMMUNOASSAY)	DID FUNCTIO	N TEST: TOTAL	0.35 - 1.93 4.87 - 12.60
TRIIODOTHYRONII by cmia (chemilumin THYROXINE (T4): S by cmia (chemilumin THYROID STIMULA	NE (T3): SERUM iescent microparticle immunoassay) SERUM	DID FUNCTIO 1.31	<b>N TEST: TOTAL</b> ng/mL	
TRIIODOTHYRONII by cmia (chemilumin THYROXINE (T4): S by cmia (chemilumin THYROID STIMULA	NE (T3): SERUM ESCENT MICROPARTICLE IMMUNOASSAY) SERUM ESCENT MICROPARTICLE IMMUNOASSAY) TTING HORMONE (TSH): SERUM ESCENT MICROPARTICLE IMMUNOASSAY)	9.19	<b>N TEST: TOTAL</b> ng/mL μgm/dL	4.87 - 12.60

overproduction(hyperthyroidism) of T4 and/or T3.

CLINICAL CONDITION	Т3	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	YRONINE (T3)	THYROXINE (T4)		THYROID STIMULATING HORMONE (1		
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	
6 - 12 Months	0.74 - 2.40	6 - 12 Months	7.10 - 16.16	6 – 12 Months	0.70 - 7.00	





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est Name			Value	Value Unit		Biological Reference interval
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	
	RECON	/IMENDATIONS OF TSH LE	EVELS DURING PREC	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, 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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Test Name		Value	Unit	Biological Reference interva
		CLINICAL PATHO	LOGY	
	URINE ROU	TINE & MICROSCOP	IC EXAMINA	ATION
PHYSICAL EXAMI	NATION			
QUANTITY RECIEV		25	ml	
COLOUR	TANCE SPECTROPHOTOMETRY	AMBER YELLOW		PALE YELLOW
TRANSPARANCY by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	CLEAR		CLEAR
SPECIFIC GRAVITY by DIP STICK/REFLEC CHEMICAL EXAMI	TANCE SPECTROPHOTOMETRY	1.02 PKR		1.002 - 1.030
REACTION	TANCE SPECTROPHOTOMETRY	ACIDIC		
PROTEIN by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	NEGATIVE (-ve)		NEGATIVE (-ve)
	TANCE SPECTROPHOTOMETRY	NEGATIVE (-ve)		NEGATIVE (-ve)
pH by DIP STICK/REELEC	TANCE SPECTROPHOTOMETRY	6		5.0 - 7.5
BILIRUBIN	TANCE SPECTROPHOTOMETRY	NEGATIVE (-ve)		NEGATIVE (-ve)
,	TANCE SPECTROPHOTOMETRY.	NEGATIVE (-ve)		NEGATIVE (-ve)
UROBILINOGEN by DIP STICK/REFLEC	CTANCE SPECTROPHOTOMETRY	<b>POSITIVE (+ve)</b>	EU/dL	0.2 - 1.0
KETONE BODIES	TANCE SPECTROPHOTOMETRY	NEGATIVE (-ve)		NEGATIVE (-ve)
,	TANCE SPECTROPHOTOMETRY	NEGATIVE (-ve)		NEGATIVE (-ve)
ASCORBIC ACID by DIP STICK/REFLEC MICROSCOPIC EXA	TANCE SPECTROPHOTOMETRY	NEGATIVE (-ve)		NEGATIVE (-ve)
RED BLOOD CELLS		NEGATIVE (-ve)	/HPF	0 - 3





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NAME

: Mr. JAGDEEP

**NOT VALID FOR MEDICO LEGAL PURPOSE** 



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Test Name		Value	Unit	<b>Biological Reference interval</b>

by MICKOSCOFT ON CENTRIFOGED ORINART SEDIMENT				
EPITHELIAL CELLS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	1-2	/HPF	ABSENT	
CRYSTALS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	NEGATIVE (-ve)		NEGATIVE (-ve)	
CASTS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	NEGATIVE (-ve)		NEGATIVE (-ve)	
BACTERIA by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	NEGATIVE (-ve)		NEGATIVE (-ve)	
OTHERS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	NEGATIVE (-ve)		NEGATIVE (-ve)	
TRICHOMONAS VAGINALIS (PROTOZOA) by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	ABSENT		ABSENT	

End Of Report



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