TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT

PKR JAIN HEALTHCARE INSTITUTE NASIRPUR, Hissar Road, AMBALA CITY- (Haryana) A PIONEER DIAGNOSTIC CENTRE

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

NAME	: Mr. ROSHAN LAL				
AGE/ GENDER	: 45 YRS/MALE		PATIENT ID	: 177433	34
COLLECTED BY	:		REG. NO./LAB NO.	: 1225(03010005
REFERRED BY	:		REGISTRATION DATE	:01/Ma	r/2025 10:11 AM
BARCODE NO.	: 12507285		COLLECTION DATE	:01/Ma	r/2025 10:36AM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITU	TE	REPORTING DATE	:01/Ma	r/2025 12:08PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBAL	A CITY - H	ARYANA		
Test Name		Value	Unit		Biological Reference interval
	SWASTI	HYA WI	ELLNESS PANEL: 1.	4	
	СОМР	PLETE B	LOOD COUNT (CBC)		
RED BLOOD CELLS	(RBCS) COUNT AND INDICES				
HAEMOGLOBIN (HI	B)	14.4	gm/dL		12.0 - 17.0
RED BLOOD CELL (RBC) COUNT OCUSING, ELECTRICAL IMPEDENCE	5.28 ^H	Millions	/cmm	3.50 - 5.00
PACKED CELL VOLU		41.5	%		40.0 - 54.0
MEAN CORPUSCULA		78.5 ^L	KR fl		80.0 - 100.0
	AR HAEMOGLOBIN (MCH) utomated hematology analyzer	27.2	pg		27.0 - 34.0
by CALCULATED BY A	AR HEMOGLOBIN CONC. (MCHC) UTOMATED HEMATOLOGY ANALYZER	34.7	g/dL		32.0 - 36.0
	UTION WIDTH (RDW-CV) utomated hematology analyzer	12.7	%		11.00 - 16.00
RED CELL DISTRIB	UTION WIDTH (RDW-SD) utomated hematology analyzer	39.2	fL		35.0 - 56.0
MENTZERS INDEX by CALCULATED		14.87	RATIO		BETA THALASSEMIA TRAIT: < 13.0 IRON DEFICIENCY ANEMIA:
					>13.0
GREEN & KING IND	DEX	18.83	RATIO		BETA THALASSEMIA TRAIT:<=
by CALCOLATED					65.0 IRON DEFICIENCY ANEMIA: > 65.0
WHITE BLOOD CEI	LLS (WBCS)				
,	BY SF CUBE & MICROSCOPY	11950 ^H	/cmm		4000 - 11000
	<u>UCOCYTE COUNT (DLC)</u>				
NEUTROPHILS	Y BY SF CUBE & MICROSCOPY	55	%		50 - 70
LYMPHOCYTES		37	%		20 - 40

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DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST



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Test Name		Value	Unit	Biological Reference interval
by FLOW CYTOMETR	Y BY SF CUBE & MICROSCOPY			
EOSINOPHILS	Y BY SF CUBE & MICROSCOPY	2	%	1 - 6
MONOCYTES	Y BY SF CUBE & MICROSCOPY	6	%	2 - 12
BASOPHILS	T BT SF COBE & MICROSCOPT	0	%	0 - 1
	Y BY SF CUBE & MICROSCOPY			
	OCYTES (WBC) COUNT			
ABSOLUTE NEUTR	OPHIL COUNT Y BY SF CUBE & MICROSCOPY	6573	/cmm	2000 - 7500
ABSOLUTE LYMPH	OCYTE COUNT	4422 ^L	/cmm	800 - 4900
by FLOW CYTOMETR ABSOLUTE EOSIN	Y BY SF CUBE & MICROSCOPY	239	KR /omm	40, 440
	Y BY SF CUBE & MICROSCOPY	239	/cmm	40 - 440
ABSOLUTE MONOC		717	/cmm	80 - 880
ABSOLUTE BASOP	Y BY SF CUBE & MICROSCOPY HIL COUNT	0	/cmm	0 - 110
by FLOW CYTOMETR	Y BY SF CUBE & MICROSCOPY		/ chilli	0 - 110
PLATELETS AND (OTHER PLATELET PREDICTIVE	MARKERS.		
PLATELET COUNT	(PLT) FOCUSING, ELECTRICAL IMPEDENCE	239000	/cmm	150000 - 450000
PLATELETCRIT (P		0.29	%	0.10 - 0.36
by HYDRO DYNAMÌC I	OCUSING, ELECTRICAL IMPEDENCE			
MEAN PLATELET V	OLUME (MPV)	12 ^H	fL	6.50 - 12.0
PLATELET LARGE	CELL COUNT (P-LCC) FOCUSING, ELECTRICAL IMPEDENCE	102000 ^H	I /cmm	30000 - 90000
PLATELET LARGE	CELL RATIO (P-LCR) FOCUSING, ELECTRICAL IMPEDENCE	42.6	%	11.0 - 45.0
PLATELET DISTRI	BUTION WIDTH (PDW) FOCUSING, ELECTRICAL IMPEDENCE	16.6	%	15.0 - 17.0
-	UCTED ON EDTA WHOLE BLOOD			





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CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTI		REPORTING DATE	:01/Mar/202504:50PM	
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AME	BALA CITY - HAI	RYANA		
Test Name		Value	Unit	Biological Referen	ce interval
	GLYCOS	SYLATED HA	EMOGLOBIN (HBA10		
	AEMOGLOBIN (HbA1c):	5	%	4.0 - 6.4	
WHOLE BLOOD	RMANCE LIQUID CHROMATOGRAPHY)				
	AGE PLASMA GLUCOSE	96.8	mg/dL	60.00 - 140.00	
	RMANCE LIQUID CHROMATOGRAPHY)	50.0	ilig/ uL	00.00 140.00	
INTERPRETATION:					
	AS PER AMERICAN D	IARETES ASSOCI			
	REFERENCE GROUP		YCOSYLATED HEMOGLOGIB	(HBAIC) in %	
Non di	abetic Adults >= 18 years		<5.7		
	t Risk (Prediabetes)		5.7 - 6.4		
D	Diagnosing Diabetes		>= 6.5		
			Age > 19 Years	< 7.0	
		Coals			
	tic goals for glycemic control	Goals	1		
	tic goals for glycemic control		s Suggested: Age < 19 Years	>8.0	

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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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AM	IBALA CITY - HARYA	NA	
Test Name		Value	Unit	Biological Reference interval
	ERYTHR	OCYTE SEDIME	NTATION RATE (ESR)
	DIMENTATION RATE (ESR) GATION BY CAPILLARY PHOTOMETRY	10 Y	mm/1st	hr 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 practitior ected by other conditions besides i	ner exactly where the inflammation. For th	e inflammation is in the is reason, the ESR is ty	pically used in conjunction with other test suc
systemic lupus eryth	ematosus W FSR	ty and response to th	lerapy in both of the a	above diseases as well as some others, such as
A low ESR can be see (polycythaemia), sign	en with conditions that inhibit the	unt (leucocytosis), a	on of red blood cells, s ind some protein abno	such as a high red blood cell count ormalities. Some changes in red cell shape (suc
1. ESR and C - reactiv 2. Generally, ESR doe 3. CRP is not affected 4. If the ESR is elevat 5. Women tend to ha 6. Drugs such as dexi	e protein (C-RP) are both markers es not change as rapidly as does Cl I by as many other factors as is ESR ed, it is typically a result of two ty ave a higher ESR, and menstruation tran, methyldopa, oral contracept nd quinine may decrease it	RP, either at the star R, making it a better (pes of proteins, glob and pregnancy can	marker of inflammation oulins or fibrinogen. cause temporary eleva	n.



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: NASIRPUR, HISSAR ROAD, A	MBALA CITY - HARYA	NA	
	Value	Unit	Biological Reference interval
CLINI	CAL CHEMISTR GLUCOSE FA	Y/BIOCHEMIST STING (F)	RY
-	: 45 YRS/MALE : : : 12507285 : P.K.R JAIN HEALTHCARE INS	: 45 YRS/MALE PAT : REC : REC : 12507285 COI : P.K.R JAIN HEALTHCARE INSTITUTE REI : NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYA	: 45 YRS/MALE PATIENT ID : REG. NO./LAB NO. : REGISTRATION DATE : 12507285 COLLECTION DATE : P.K.R JAIN HEALTHCARE INSTITUTE REPORTING DATE : NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYANA

2. A fasting plasma glucose level below 100 mg/dl is considered normal. 2. A fasting plasma glucose level between 100 - 125 mg/dl is considered as glucose intolerant or prediabetic. A fasting and post-prandial blood

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 CODE.	: P.K.R JAIN HEALTHCARE INS	STITUTE	REPORTING DATE	:01/Mar/2025 11:55AM
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 ON		244.09 ^H	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)	324.27 ^H	mg/dL	OPTIMAL: < 150.0 BORDERLINE HIGH: 150.0 - 199.0 HIGH: 200.0 - 499.0 VERY HIGH: > OR = 500.0
HDL CHOLESTERO by SELECTIVE INHIBIT	L (DIRECT): SERUM	45.1	mg/dL	LOW HDL: < 30.0 BORDERLINE HIGH HDL: 30.0 60.0 HIGH HDL: > OR = 60.0
LDL CHOLESTEROI by CALCULATED, SPE		134.14 ^H	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		198.99 ^H	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(64.85 ^H	mg/dL	0.00 - 45.00
TOTAL LIPIDS: SER	RUM	812.45 ^H	mg/dL	350.00 - 700.00
CHOLESTEROL/HE by CALCULATED, SPE	DL RATIO: SERUM	5.41 ^H	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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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.)**



Page 6 of 17

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

Test Name	Value	Unit	Biological Reference interval
LDL/HDL RATIO: SERUM by Calculated, SPECTROPHOTOMETRY	2.97	RATIO	LOW RISK: 0.50 - 3.0 MODERATE RISK: 3.10 - 6.0 HIGH RISK: > 6.0
TRIGLYCERIDES/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	7.19 ^H	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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Test Name		Value	Unit	Biological Reference interva
	LIVER	FUNCTIO	ON TEST (COMPLETE)	
BILIRUBIN TOTAL: by DIAZOTIZATION, SF	SERUM PECTROPHOTOMETRY	0.55	mg/dL	INFANT: 0.20 - 8.00 ADULT: 0.00 - 1.20
	C (CONJUGATED): SERUM	0.14	mg/dL	0.00 - 0.40
BILIRUBIN INDIRE by CALCULATED, SPE	CT (UNCONJUGATED): SERUM	0.41	mg/dL	0.10 - 1.00
SGOT/AST: SERUM by IFCC, WITHOUT PY	RIDOXAL PHOSPHATE	19.89	U/L	7.00 - 45.00
SGPT/ALT: SERUM	RIDOXAL PHOSPHATE	37.03	U/L	0.00 - 49.00
AST/ALT RATIO: SI by CALCULATED, SPE	ERUM	0. <mark>54</mark>	RATIO	0.00 - 46.00
ALKALINE PHOSPH by Para Nitrophen Propanol	IATASE: SERUM YL PHOSPHATASE BY AMINO METHYL	97.17	U/L	40.0 - 130.0
GAMMA GLUTAMY by SZASZ, SPECTROF	L TRANSFERASE (GGT): SERUM	36.93	U/L	0.00 - 55.0
TOTAL PROTEINS: by BIURET, SPECTRO		6.23	gm/dL	6.20 - 8.00
ALBUMIN: SERUM by BROMOCRESOL G	REEN	4.25	gm/dL	3.50 - 5.50
GLOBULIN: SERUM by CALCULATED, SPE		1.98 ^L	gm/dL	2.30 - 3.50
A : G RATIO: SERUN by CALCULATED, SPE		2.15 ^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.

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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5

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, AMI	BALA CITY - H	IARYANA	
Test Name		Value	Unit	Biological Reference interva
	KIDNI	EY FUNCTI	ON TEST (COMPLETE)
•	ATE DEHYDROGENASE (GLDH)	17.61	mg/dL	10.00 - 50.00
CREATININE: SERU by ENZYMATIC, SPECT		0.73	mg/dL	0.40 - 1.40
by CALCULATED, SPEC		8.23	mg/dL	7.0 - 25.0
BLOOD UREA NITR RATIO: SERUM by calculated, spec	OGEN (BUN)/CREATININE CTROPHOTOMETRY	11.27	RATIO	10.0 - 20.0
JREA/CREATININE	E RATIO: SERUM	<mark>24.12</mark>	RATIO	
JRIC ACID: SERUM	E PEROXIDASE	3.65	mg/dL	3.60 - 7.70
CALCIUM: SERUM by ARSENAZO III, SPEC		9.84	mg/dL	8.50 - 10.60
PHOSPHOROUS: SE by phosphomolybd, ELECTROLYTES	RUM ate, spectrophotometry	3.41	mg/dL	2.30 - 4.70
SODIUM: SERUM by ISE (ION SELECTIVE	E ELECTRODE)	139	mmol/L	135.0 - 150.0
OTASSIUM: SERUN	Л	4.64	mmol/L	3.50 - 5.00
CHLORIDE: SERUM by ISE (ION SELECTIVE	E ELECTRODE)	104.25	mmol/L	90.0 - 110.0
	ERULAR FILTERATION RATE ERULAR FILTERATION RATE	114.3		

by CALCULATED

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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NOT VALID FOR MEDICO LEGAL PURPOSE



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REFERRED BY	:	R	EGISTRATION DAT	TE : 01/Mar/2025 10:1	1 AM
BARCODE NO.	: 12507285		OLLECTION DATE	: 01/Mar/2025 10:3	
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTIT		EPORTING DATE	: 01/Mar/2025 04:3	
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMB			. 01/ 141/ 2020 01.0	
Test Name		Value	Unit	Biologica	l Reference interval
9. Certain drugs (e.g. INCREASED RATIO (>2 1. Postrenal azotemia 2. Prerenal azotemia DECREASED RATIO (< 1. Acute tubular necr 2. Low protein diet ar 3. Severe liver diseas 4. Other causes of de 5. Repeated dialysis 6. Inherited hyperam	nd starvation.	EVELS: The than creatinine than creatinine sout of extracell in blood).	ular fluid).	ıropathy).	
1. Phenacimide thera 2. Rhabdomyolysis (r 3. Muscular patients INAPPROPIATE RATIO 1. Diabetic ketoacido should produce an in 2. Cephalosporin ther	osis (acetoacetate causes false incre creased BUN/creatinine ratio). rapy (interferes with creatinine mea	ease in creatinine		odologies,resulting in norma	al ratio when dehydrat
ESTIMATED GLOMERU CKD STAGE	JLAR FILTERATION RATE: DESCRIPTION		/min/1.73m2)	ASSOCIATED FINDINGS	1
G1	Normal kidney function		>90	No proteinuria	1
G2	Kidney damage with normal or high GFR		>90	Presence of Protein , Albumin or cast in urine	1
				i	-

0.	i torinar namoj ranotion	.,,,	no protoniuna
G2	Kidney damage with	>90	Presence of Protein ,
	normal or high GFR		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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NAME	: Mr. ROSHAN LAL		
AGE/ GENDER	: 45 YRS/MALE	PATIENT ID	: 1774334
COLLECTED BY	:	REG. NO./LAB NO.	: 122503010005
REFERRED BY	:	REGISTRATION DATE	: 01/Mar/2025 10:11 AM
BARCODE NO.	: 12507285	COLLECTION DATE	:01/Mar/2025 10:36AM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITUTE	REPORTING DATE	: 01/Mar/2025 04:38PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBALA CITY - H	IARYANA	

Test Name	Value	Unit	Biological Reference interval

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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NAME	: Mr. ROSHAN LAL		

	IRON	PROFILE	
IRON: SERUM by Ferrozine, spectrophotometry	97.9	μg/dL	59.0 - 158.0
UNSATURATED IRON BINDING CAPA :SERUM by Ferrozine, spectrophotometery	ACITY (UIBC) 93.8 ^L	μg/dL	150.0 - 336.0
TOTAL IRON BINDING CAPACITY (TI :SERUM by SPECTROPHOTOMETERY	BC) 191.7^L	µg/dL	230 - 430
%TRANSFERRIN SATURATION: SERI by Calculated, spectrophotometery	01.07	%	15.0 - 50.0
TRANSFERRIN: SERUM by SPECTROPHOTOMETERY (FERENE)	136.11 ^L	mg/dL	200.0 - 350.0
	NEMIA OF CHRONIC DISEASE	IRON DEFICIENCY ANEMIA	THALASSEMIA α/β TRAIT

VARIABLES	ANEMIA OF CHRONIC DISEASE	IRON DEFICIENCY ANEMIA	THALASSEMIA α/β TRAIT
SERUM IRON: Normal to Reduce		Reduced	Normal
TOTAL IRON BINDING CAPACITY:	Decreased	Increased	Normal
% TRANSFERRIN SATURATION:	Decreased	Decreased < 12-15 %	Normal
SERUM FERRITIN:	Normal to Increased	Decreased	Normal or Increased
IDON.			

IRON:

1.Serum iron studies is recommended for differential diagnosis of microcytic hypochromic anemia.i.e iron deficiency anemia, zinc deficiency anemia, anemia of chronic disease and thalassemia syndromes.

It is essential to isolate iron deficiency anemia from Beta thalassemia syndromes because during iron replacement which is therapeutic for iron deficiency anemia, is severely contra-indicated in Thalassemia.
TOTAL IRON BINDING CAPACITY (TIBC):

1.It is a direct measure of protein transferrin which transports iron from the gut to storage sites in the bone marrow. % TRANSFERRIN SATURATION:

1. Occurs in idiopathic hemochromatosis and transfusional hemosiderosis where no unsaturated iron binding capacity is available for iron mobilization. Similar condition is seen in congenital deficiency of transferrin.



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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBAL	A CITY - HARYAN	A	
Test Name		Value	Unit	Biological Reference interval
Test Name		Value ENDOCRIN		Biological Reference interval
Test Name	THYRO	ENDOCRIN		Biological Reference interval
TRIIODOTHYRONII		ENDOCRING DID FUNCTION 1.416	DLOGY	0.35 - 1.93
TRIIODOTHYRONII by CMIA (CHEMILUMIN THYROXINE (T4): S	NE (T3): SERUM ESCENT MICROPARTICLE IMMUNOASSAY)	ENDOCRIN DID FUNCTION 1.416 7.33	DLOGY N TEST: TOTAL	U
TRIIODOTHYRONII by cmia (chemilumin THYROXINE (T4): S by cmia (chemilumin THYROID STIMULA	NE (T3): SERUM ESCENT MICROPARTICLE IMMUNOASSAY) ERUM ESCENT MICROPARTICLE IMMUNOASSAY) TING HORMONE (TSH): SERUM ESCENT MICROPARTICLE IMMUNOASSAY)	ENDOCRING DID FUNCTION 1.416 7.33 3.057	DLOGY N TEST: TOTAL ng/mL	0.35 - 1.93

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.

TRIIODOTHYRONINE (T3)		THYROXINE (T4)		THYROID STIMULATING 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	
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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Test Name		Value Unit		t	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	
	RECOM	MENDATIONS OF TSH LI	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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: Mr. ROSHAN LAL

NAME

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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYANA						
Test Name		Value	Unit	Biological Reference interva			
		CLINICAL PATHO	DLOGY				
	URINE ROU	UTINE & MICROSCO	PIC EXAMINA	ATION			
PHYSICAL EXAMIN	VATION						
QUANTITY RECIEV	ED tance spectrophotometry	30	ml				
COLOUR by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	PALE YELLOW		PALE YELLOW			
TRANSPARANCY by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	HAZY		CLEAR			
SPECIFIC GRAVITY by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	1.01 PKR		1.002 - 1.030			
CHEMICAL EXAMI	<u>NATION</u>						
REACTION by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	ACIDIC					
PROTEIN by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	NEGATIVE (-ve)		NEGATIVE (-ve)			
SUGAR by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	NEGATIVE (-ve)		NEGATIVE (-ve)			
pH by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	5.5		5.0 - 7.5			
BILIRUBIN by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY	NEGATIVE (-ve)		NEGATIVE (-ve)			
NITRITE by DIP STICK/REFLEC	TANCE SPECTROPHOTOMETRY.	NEGATIVE (-ve)		NEGATIVE (-ve)			
•	TANCE SPECTROPHOTOMETRY	NOT DETECTED	EU/dL	0.2 - 1.0			
	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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NOT VALID FOR MEDICO LEGAL PURPOSE



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Tost Namo	Valua	Unit	Biological Reference interval	

Test Name	Value	Unit	Biological Reference interval
by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT			
PUS CELLS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	4-5	/HPF	0 - 5
EPITHELIAL CELLS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	3-4	/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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