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. NARMAIL SINGH						
AGE/ GENDER	: 69 YRS/MALE	PAT	FIENT ID	: 1816215			
COLLECTED BY	:			: 122504030007 : 03/Apr/2025 09:19 AM			
REFERRED BY	:						
BARCODE NO.	: 12507873	COL	LECTION DATE	: 03/Apr/2025 09:48AM			
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITUTE	REI	PORTING DATE	: 03/Apr/2025 01:23PM			
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBALA (CITY - HARYA	NA				
Test Name	V	alue	Unit	Biological Reference interval			
	SWASTHY	A WELL	NESS PANEL: 1	.0			
	COMPLE	TE BLOO	D COUNT (CBC)				
RED BLOOD CEL	LS (RBCS) COUNT AND INDICES						
HAEMOGLOBIN (H	IB)	9.5 ^L	gm/dL	12.0 - 17.0			
RED BLOOD CELL	, (RBC) COUNT FOCUSING, ELECTRICAL IMPEDENCE	3.08 ^L	Millions	cmm 3.50 - 5.00			
PACKED CELL VO	LUME (PCV) AUTOMATED HEMATOLOGY ANALYZER	27.8 ^L	%	40.0 - 54.0			
MEAN CORPUSCU	LAR VOLUME (MCV)	90.3	fL	80.0 - 100.0			
	LAR HAEMOGLOBIN (MCH) AUTOMATED HEMATOLOGY ANALYZER	30.9	pg	27.0 - 34.0			
	LAR HEMOGLOBIN CONC. (MCHC)	34.3	g/dL	32.0 - 36.0			
RED CELL DISTRI	BUTION WIDTH (RDW-CV)	15.5	%	11.00 - 16.00			
	BUTION WIDTH (RDW-SD)	53.1	fL	35.0 - 56.0			
MENTZERS INDEX	K	29.32	RATIO	BETA THALASSEMIA TRAIT 13.0 IRON DEFICIENCY ANEMIA: >13.0			
GREEN & KING IN by CALCULATED	IDEX	133.04	RATIO	BETA THALASSEMIA TRAIT <= 74.1 IRON DEFICIENCY ANEMIA: >= 74.1			
WHITE BLOOD C	ELLS (WBCS)						
TOTAL LEUCOCY by FLOW CYTOMETR	TE COUNT (TLC) y by sf cube & microscopy	4830	/cmm	4000 - 11000			
<u>DIFFERENTIAL L</u>	EUCOCYTE COUNT (DLC)						
NEUTROPHILS		65	%	50 - 70			

DR.VINAY CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY & MICROBIOLOGY) MBBS , MD (PATHOLOGY)

DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST



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	· · · · · · · · · · · · · · · · · · ·							
Test Name		Value	Unit	Biological Reference interval				
by FLOW CYTOMETR	RY BY SF CUBE & MICROSCOPY							
LYMPHOCYTES		22	%	20 - 40				
	RY BY SF CUBE & MICROSCOPY							
EOSINOPHILS	RY BY SF CUBE & MICROSCOPY	2	%	1 - 6				
MONOCYTES		11	%	2 - 12				
	RY BY SF CUBE & MICROSCOPY		10					
BASOPHILS		0	%	0 - 1				
•	RY BY SF CUBE & MICROSCOPY							
ABSOLUTE LEUR	<u>KOCYTES (WBC) COUNT</u>							
ABSOLUTE NEUT		3140	/cmm	2000 - 7500				
by FLOW CYTOMETR ABSOLUTE LYMP	RY BY SF CUBE & MICROSCOPY	- PNI	/cmm	800 - 4900				
	RY BY SF CUBE & MICROSCOPY	1063 ^L	/clilli	800 - 4900				
ABSOLUTE EOSIN		97	/cmm	40 - 440				
by FLOW CYTOMETR	RY BY SF CUBE & MICROSCOPY							
ABSOLUTE MONO		531	/cmm	80 - 880				
ABSOLUTE BASO	RY BY SF CUBE & MICROSCOPY	0	lomm	0 - 110				
	RY BY SF CUBE & MICROSCOPY	0	/cmm	0 - 110				
,	OTHER PLATELET PREDICTIV	<u>'E MARKERS.</u>						
PLATELET COUN		214000	/cmm	150000 - 450000				
	FOCUSING, ELECTRICAL IMPEDENCE	211000	, chilli	100000 100000				
PLATELETCRIT (- /	0.21	%	0.10 - 0.36				
	FOCUSING, ELECTRICAL IMPEDENCE	10	~	6.50 10.0				
MEAN PLATELET	VOLUME (MPV)	10	fL	6.50 - 12.0				
•	E CELL COUNT (P-LCC)	53000	/cmm	30000 - 90000				
	FOCUSING, ELECTRICAL IMPEDENCE							
	E CELL RATIO (P-LCR)	24.7	%	11.0 - 45.0				
	FOCUSING, ELECTRICAL IMPEDENCE RIBUTION WIDTH (PDW)	16.2	%	15.0 - 17.0				
	FOCUSING, ELECTRICAL IMPEDENCE	10.2	70	15.0 - 17.0				
-	UCTED ON EDTA WHOLE BLOOD							



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Fest Name		Value	Unit	Biological Reference interval
	ERYTHROC	CYTE SEDIMEN	NTATION RATE (ESR)
ERYTHROCYTE SH	EDIMENTATION RATE (ESR)	84 ^H	mm/1st hr	0 - 20
		۰.		
	GATION BY CAPILLARY PHOTOMETRY	0.		
NTERPRETATION:	ic test because an elevated result of	ften indicates the p	resence of inflammatic	on associated with infection, cancer and aut
NTERPRETATION: I. ESR is a non-specif mmune disease, but	ic test because an elevated result of does not tell the health practitioner	ften indicates the p	inflammation is in the	body or what is causing it.
NTERPRETATION: I. ESR is a non-specif mmune disease, but 2. An ESR can be affe	ic test because an elevated result of does not tell the health practitioner	ften indicates the p	inflammation is in the	on associated with infection, cancer and aut body or what is causing it. ically used in conjunction with other test su
NTERPRETATION: L. ESR is a non-specif mmune disease, but 2. An ESR can be affe as C-reactive protein 3. This test may also	ic test because an elevated result of does not tell the health practitioner cted by other conditions besides inf be used to monitor disease activity	ften indicates the p r exactly where the flammation. For this	inflammation is in the s reason, the ESR is typ	body or what is causing it.
NTERPRETATION: 1. ESR is a non-specif mmune disease, but 2. An ESR can be affe- as C-reactive protein 3. This test may also l systemic lupus erythe	ic test because an elevated result of does not tell the health practitioner cted by other conditions besides inf be used to monitor disease activity ematosus	ften indicates the p r exactly where the flammation. For this	inflammation is in the s reason, the ESR is typ	body or what is causing it. ically used in conjunction with other test su
NTERPRETATION: 1. ESR is a non-specif mmune disease, but 2. An ESR can be affe- is C-reactive protein 3. This test may also l systemic lupus erythe CONDITION WITH LON A low ESR can be see	ic test because an elevated result of does not tell the health practitioner cted by other conditions besides inf be used to monitor disease activity ematosus W ESR n with conditions that inhibit the no	ften indicates the pire exactly where the flammation. For this and response to the provide the provide the provide the provide the provided the prov	inflammation is in the s reason, the ESR is typ erapy in both of the ab	body or what is causing it. ically used in conjunction with other test su ove diseases as well as some others, such a ch as a high red blood cell count
NTERPRETATION: 1. ESR is a non-specif mmune disease, but 2. An ESR can be affe- is C-reactive protein 3. This test may also i condition with LOV A low ESR can be see polycythaemia). sign	ic test because an elevated result of does not tell the health practitioner cted by other conditions besides inf be used to monitor disease activity ematosus N ESR n with conditions that inhibit the no lificantly high white blood cell coun	ften indicates the pire exactly where the flammation. For this and response to the private sedimentation of the private sedimentatio	inflammation is in the s reason, the ESR is typ erapy in both of the ab	body or what is causing it. ically used in conjunction with other test su ove diseases as well as some others, such a
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Test Name		Value	Unit	Biological Reference interval				
	CLINIC	CAL CHEMISTRY	Y/BIOCHEMIS	STRY				
		GLUCOSE FAS	STING (F)					
GLUCOSE FASTIN by GLUCOSE OXIDAS	G (F): PLASMA E - PEROXIDASE (GOD-POD)	113.12 ^H	mg/dL	NORMAL: < 100.0 PREDIABETIC: 100.0 - 125.0 DIABETIC: > 0R = 126.0				
INTERPRETATION	H AMERICAN DIABETES ASSOCIA							

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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Test Name		Value	Unit	Biological Reference interval				
		LIPID PRO	OFILE : BASIC					
CHOLESTEROL TO by CHOLESTEROL OX		162.01	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)	131.21	mg/dL	OPTIMAL: < 150.0 BORDERLINE HIGH: 150.0 - 199.0 HIGH: 200.0 - 499.0				
HDL CHOLESTER(by SELECTIVE INHIBIT	DL (DIRECT): SERUM	44.46	mg/dL	VERY HIGH: > OR = 500.0 LOW HDL: < 30.0 BORDERLINE HIGH HDL: 30. 60.0				
LDL CHOLESTERC by CALCULATED, SPE		91.31	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		117.55	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		26.24	mg/dL	VERY HIGH: > OR = 220.0 0.00 - 45.00				
TOTAL LIPIDS: SE by CALCULATED, SPE	RUM	455.23	mg/dL	350.00 - 700.00				
CHOLESTEROL/HI	DL RATIO: SERUM	3.64	RATIO	LOW RISK: 3.30 - 4.40 AVERAGE RISK: 4.50 - 7.0				

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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.)**



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

			HIGH RISK: > 11.0
LDL/HDL RATIO: SERUM	2.05	RATIO	LOW RISK: 0.50 - 3.0
by CALCULATED, SPECTROPHOTOMETRY			MODERATE RISK: 3.10 - 6.0
			HIGH RISK: > 6.0
TRIGLYCERIDES/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	2.95 ^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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NAME	: Mr. NARMAIL SINGH						
AGE/ GENDER	: 69 YRS/MALE		PATIENT ID	: 1816215			
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BARCODE NO.	: 12507873		COLLECTION DATE	: 03/Apr/2025 09:48AM			
CLIENT CODE.			REPORTING DATE	: 03/Apr/2025 05:02PM			
CLIENT ADDRESS			ARYANA				
Test Name	,	Value	Unit	Biological Reference interval			
	LIVER FU	JNCTIO	N TEST (COMPLETE)			
BILIRUBIN TOTAL by DIAZOTIZATION, SF	: SERUM PECTROPHOTOMETRY	0.46	mg/dL	INFANT: 0.20 - 8.00 ADULT: 0.00 - 1.20			
	Г (CONJUGATED): SERUM	0.16	mg/dL	0.00 - 0.40			
BILIRUBIN INDIRE by CALCULATED, SPE	CT (UNCONJUGATED): SERUM	0.3	mg/dL	0.10 - 1.00			
SGOT/AST: SERUN by IFCC, WITHOUT PY	1 RIDOXAL PHOSPHATE	11.34	U/L	7.00 - 45.00			
SGPT/ALT: SERUM by IFCC, WITHOUT PY	[RIDOXAL PHOSPHATE	7.8		0.00 - 49.00			
AST/ALT RATIO: S by CALCULATED, SPE		1.45	RATIO	0.00 - 46.00			
ALKALINE PHOSPI by PARA NITROPHEN PROPANOL	HATASE: SERUM YL PHOSPHATASE BY AMINO METHYL	87.08	U/L	40.0 - 130.0			
GAMMA GLUTAM by SZASZ, SPECTROF	YL TRANSFERASE (GGT): SERUM PHTOMETRY	14.04	U/L	0.00 - 55.0			
TOTAL PROTEINS by BIURET, SPECTRO		5.25 ^L	gm/dL	6.20 - 8.00			
ALBUMIN: SERUM by BROMOCRESOL G		3.39 ^L	gm/dL	3.50 - 5.50			
GLOBULIN: SERUN by CALCULATED, SPE		1.86 ^L	gm/dL	2.30 - 3.50			
A : G RATIO: SERU by CALCULATED, SPE		1.82	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





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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	NC)ST	IC	S	IG	NIF	FIC	:А	١N	ICI	E:

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, AMBA	AR ROAD, AMBALA CITY - HARYANA		1	
Test Name		Value	Unit	Biological Reference interva	
	KIDNEY	FUNCTI	ON TEST (COMPLETI	E)	
JREA: SERUM		18.64	mg/dL	10.00 - 50.00	
	E DEHYDROGENASE (GLDH)	10.0.			
CREATININE: SERUN		0.69	mg/dL	0.40 - 1.40	
by ENZYMATIC, SPECTR	GEN (BUN): SERUM	8.71	mg/dL	7.0 - 25.0	
by CALCULATED, SPECT		0.71	ing/ull	1.0 25.0	
BLOOD UREA NITROGEN (BUN)/CREATININE		12.62 RATIO	10.0 - 20.0		
RATIO: SERUM by CALCULATED, SPECT	POPHOTOMETRY				
UREA/CREATININE F		27.01	RATIO		
by CALCULATED, SPECT					
URIC ACID: SERUM		2.42 ^L	mg/dL	3.60 - 7.70	
by URICASE - OXIDASE F	PEROXIDASE	8.88	mg/dL	8.50 - 10.60	
by ARSENAZO III, SPECT	ROPHOTOMETRY	0.00	ingut	0.50 10.00	
PHOSPHOROUS: SER		2.67	mg/dL	2.30 - 4.70	
-	E, SPECTROPHOTOMETRY				
ELECTROLYTES		1 4 1 . 2 5	1.7	125.0 150.0	
SODIUM: SERUM by ISE (ION SELECTIVE E	ELECTRODE	141.25	mmol/L	135.0 - 150.0	
POTASSIUM: SERUM		4.63	mmol/L	3.50 - 5.00	
by ISE (ION SELECTIVE E	ELECTRODE)				
CHLORIDE: SERUM by ISE (ION SELECTIVE E		105.94	mmol/L	90.0 - 110.0	
	ERULAR FILTERATION RATE				
	RULAR FILTERATION RATE	100.2			
(eGFR): SERUM					
by CALCULATED					
INTERPRETATION:	n pro, and post ronal azotomia				
	n pre- and post renal azotemia. 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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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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Test Name		Value	Unit	Biological Reference interv
		CLINICAL P.	ATHOLOGY	
	URINE R	OUTINE & MICR	OSCOPIC EXAMI	NATION
PHYSICAL EXAMI	NATION			
QUANTITY RECIEV		25	ml	
by DIP STICK/REFLECT	ANCE SPECTROPHOTOMETRY	YELLOW		PALE YELLOW
	ANCE SPECTROPHOTOMETRY			
FRANSPARANCY by DIP STICK/REFLECT	ANCE SPECTROPHOTOMETRY	HAZY		CLEAR
SPECIFIC GRAVITY		1.02		1.002 - 1.030
by DIP STICK/REFLECT CHEMICAL EXAM	ANCE SPECTROPHOTOMETRY			
REACTION		ALKALINE		
•	ANCE SPECTROPHOTOMETRY			
PROTEIN by DIP STICK/REFLECT	ANCE SPECTROPHOTOMETRY	TRACE		NEGATIVE (-ve)
SUGAR		NEGATIVE	(-ve)	NEGATIVE (-ve)
by DIP STICK/REFLECT	ANCE SPECTROPHOTOMETRY	7.5		5.0 - 7.5
	ANCE SPECTROPHOTOMETRY			
BILIRUBIN by DIP STICK/REFLECT	ANCE SPECTROPHOTOMETRY	NEGATIVE	(-ve)	NEGATIVE (-ve)
NITRITE		NEGATIVE	(-ve)	NEGATIVE (-ve)
by DIP STICK/REFLECT UROBILINOGEN	ANCE SPECTROPHOTOMETRY.	NOT DETE	CTED EU/dL	0.2 - 1.0
	ANCE SPECTROPHOTOMETRY	NOT DETEN	ETED EU/dL	0.2 - 1.0
KETONE BODIES	ANCE SPECTROPHOTOMETRY	NEGATIVE	(-ve)	NEGATIVE (-ve)
BLOOD		NEGATIVE	(-ve)	NEGATIVE (-ve)
•	ANCE SPECTROPHOTOMETRY		. ,	
ASCORBIC ACID	ANCE SPECTROPHOTOMETRY	NEGATIVE	(-ve)	NEGATIVE (-ve)



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Test Name		Value	Unit	Biological Reference interval
RED BLOOD CELL	S (RBCs) CENTRIFUGED URINARY SEDIMENT	NEGATIVE (-ve)	/HPF	0 - 3
PUS CELLS by MICROSCOPY ON (CENTRIFUGED URINARY SEDIMENT	8-10	/HPF	0 - 5
EPITHELIAL CELL by MICROSCOPY ON (S CENTRIFUGED URINARY SEDIMENT	4-6	/HPF	ABSENT
CRYSTALS by MICROSCOPY ON (CENTRIFUGED URINARY SEDIMENT	CALCIUM OXALA	ГЕ (++)	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 TRICHOMONAS VAGINALIS (PROTOZOA)

by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT

* * * End Of Report *

ABSENT

NEGATIVE (-ve)



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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.)**



NEGATIVE (-ve)

ABSENT