A PIONEER DIAGNOSTIC CENTRE

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

NAME	: Mr. KULWANT SINGH				
AGE/ GENDER	: 70 YRS/MALE		PATIENT ID	: 1587697	
COLLECTED BY	:		REG. NO./LAB NO.	: 1224091	30021
REFERRED BY	:		<b>REGISTRATION DATE</b>	:13/Sep/20	24 03:49 PM
BARCODE NO.	: 12504675		COLLECTION DATE	: 13/Sep/20	24 04:51PM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITU	JΤΕ	<b>REPORTING DATE</b>	:13/Sep/20	24 05:33PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBAI	LA CITY - HA	RYANA		
Test Name		Value	Unit	Bio	blogical Reference interval
	SWAS	THYA WE	LLNESS PANEL: 1.0		
	CON	IPLETE BL	OOD COUNT (CBC)		
RED BLOOD CELLS (F	RBCS) COUNT AND INDICES				
HAEMOGLOBIN (HB) by CALORIMETRIC	)	9.8 <sup>L</sup>	gm/dL	12	2.0 - 17.0
RED BLOOD CELL (RE	BC) COUNT FOCUSING, ELECTRICAL IMPEDENCE	2.86 <sup>L</sup>	Millions/c	mm 3.	50 - 5.00
PACKED CELL VOLUN	/IE (PCV)	27.7 <sup>L</sup>	%	40	0.0 - 54.0
MEAN CORPUSCULA		96.8 P	KR fl	80	0.0 - 100.0
MEAN CORPUSCULA	AUTOMATED HEMATOLOGY ANALYZER R HAEMOGLOBIN (MCH) AUTOMATED HEMATOLOGY ANALYZER	34.3 <sup>H</sup>	pg	27	2.0 - 34.0
MEAN CORPUSCULA	R HEMOGLOBIN CONC. (MCHC)	35.4	g/dL	32	2.0 - 36.0
RED CELL DISTRIBUT	TON WIDTH (RDW-CV)	15.7	%	11	.00 - 16.00
RED CELL DISTRIBUT	TION WIDTH (RDW-SD) automated hematology analyzer	56.3 <sup>H</sup>	fL	35	.0 - 56.0
MENTZERS INDEX by CALCULATED		33.85	RATIO		TA THALASSEMIA TRAIT: < 13 ON DEFICIENCY ANEMIA: >13.
GREEN & KING INDE by calculated	X	53.19	RATIO		TA THALASSEMIA TRAIT:<= 65 ON DEFICIENCY ANEMIA: > 65
WHITE BLOOD CELLS	<u>S (WBCS)</u>				
TOTAL LEUCOCYTE C by FLOW CYTOMETRY DIFFERENTIAL LEUCO	Y BY SF CUBE & MICROSCOPY	10910	/cmm	40	00 - 11000
NEUTROPHILS by flow cytometr	Y BY SF CUBE & MICROSCOPY	82 <sup>H</sup>	%	50	- 70
LYMPHOCYTES	Y BY SF CUBE & MICROSCOPY	10 <sup>L</sup>	%	20	) - 40
EOSINOPHILS	Y BY SF CUBE & MICROSCOPY	1	%	1.	- 6



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Test Name		Value	Unit	Biological Reference interval
MONOCYTES		7	%	2 - 12
BASOPHILS	Y BY SF CUBE & MICROSCOPY Y BY SF CUBE & MICROSCOPY <b>TES (WBC) COUNT</b>	0	%	0 - 1
ABSOLUTE NEUTRO		8946 <sup>H</sup>	/cmm	2000 - 7500
ABSOLUTE LYMPHO	<b>Y BY SF CUBE &amp; MICROSCOPY</b> CYTE COUNT Y BY SF CUBE & MICROSCOPY	1091 <sup>L</sup>	/cmm	800 - 4900
ABSOLUTE EOSINOP		109	/cmm	40 - 440
ABSOLUTE MONOCY		764	KR /cmm	80 - 880
ABSOLUTE BASOPHI	L COUNT y by sf cube & microscopy	0	/cmm	0 - 110
PLATELETS AND OTI	HER PLATELET PREDICTIVE MARKER	<u>RS.</u>		
PLATELET COUNT (P	LT) FOCUSING, ELECTRICAL IMPEDENCE	99000 <sup>L</sup>	/cmm	150000 - 450000
PLATELETCRIT (PCT)	OCUSING, ELECTRICAL IMPEDENCE	0.11	%	0.10 - 0.36
MEAN PLATELET VO		11	fL	6.50 - 12.0
PLATELET LARGE CEI		35000	/cmm	30000 - 90000
PLATELET LARGE CE		35.2	%	11.0 - 45.0
PLATELET DISTRIBU		16.1	%	15.0 - 17.0





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Test Name		Value	Unit	Biological Reference interval
	ERYTHR	OCYTE SEDIMEN	TATION RATE (ESR	)
	ENTATION RATE (ESR)	98 <sup>H</sup>	mm/1st hi	0 - 20
1. ESR is a non-specific immune disease, but do 2. An ESR can be affect as C-reactive protein	bes not tell the health practition ed by other conditions besides in	er exactly where the nflammation. For this	inflammation is in the s reason, the ESR is typi	n associated with infection, cancer and aut body or what is causing it. cally used in conjunction with other test su ove diseases as well as some others, such a

systemic lupus erythematosus

#### **CONDITION WITH LOW ESR**

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.
4. If the ESR is elevated, it is typically a result of two types of proteins, globulins or fibrinogen.
5. 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 explicit contraceptives are the process. aspirin, cortisone, and quinine may decrease it



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Test Name		Value	Unit	Biological Reference interval
	CLIN	ICAL CHEMISTRY	/BIOCHEMISTRY	Y
		GLUCOSE FAS	TING (F)	
GLUCOSE FASTING ( by GLUCOSE OXIDAS	F): PLASMA ie - peroxidase (god-pod)	107.67 <sup>H</sup>	mg/dL	NORMAL: < 100.0 PREDIABETIC: 100.0 - 125.0 DIABETIC: > 0R = 126.0
	H AMERICAN DIABETES ASSOCIAT			DIADE 110. > UK = 120.0

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.
A fasting plasma glucose level in excess of 125 mg/dl on both occasions is confirmatory for diabetic state.



TEST PERFORMED AT KOS DIAGNOSTIC LAB. AMBALA CANTT

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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AN	MBALA CITY - HA	ARYANA	
Test Name		Value	Unit	Biological Reference interval
		LIPID PR	OFILE : BASIC	
CHOLESTEROL TOTA by CHOLESTEROL OX		76.84	mg/dL	OPTIMAL: < 200.0 BORDERLINE HIGH: 200.0 - 239.0 HIGH CHOLESTEROL: > OR = 240.0
TRIGLYCERIDES: SER by GLYCEROL PHOSP	RUM PHATE OXIDASE (ENZYMATIC)	43.79	mg/dL	OPTIMAL: < 150.0 BORDERLINE HIGH: 150.0 - 199.0 HIGH: 200.0 - 499.0 VERY HIGH: > OR = 500.0
HDL CHOLESTEROL ( by SELECTIVE INHIBI		19.23 <sup>L</sup>	mg/dL	LOW HDL: < 30.0 BORDERLINE HIGH HDL: 30.0 - 60.0 HIGH HDL: > OR = 60.0
LDL CHOLESTEROL: S by CALCULATED, SPE		48.85	mg/dL	OPTIMAL: < 100.0 ABOVE OPTIMAL: 100.0 - 129.0 BORDERLINE HIGH: 130.0 - 159.0 HIGH: 160.0 - 189.0 VERY HIGH: > OR = 190.0
NON HDL CHOLESTE by calculated, spe		57.61	mg/dL	OPTIMAL: < 130.0 ABOVE OPTIMAL: 130.0 - 159.0 BORDERLINE HIGH: 160.0 - 189.0 HIGH: 190.0 - 219.0 VERY HIGH: > OR = 220.0
VLDL CHOLESTEROL: by CALCULATED, SPE		8.76	mg/dL	0.00 - 45.00
TOTAL LIPIDS: SERU	M	197.47 <sup>L</sup>	mg/dL	350.00 - 700.00
CHOLESTEROL/HDL I by CALCULATED, SPE	RATIO: SERUM	4	RATIO	LOW RISK: 3.30 - 4.40 AVERAGE RISK: 4.50 - 7.0 MODERATE RISK: 7.10 - 11.0 HIGH RISK: > 11.0
LDL/HDL RATIO: SER by CALCULATED, SPE		2.54	RATIO	LOW RISK: 0.50 - 3.0 MODERATE RISK: 3.10 - 6.0 HIGH RISK: > 6.0

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Test Name	Value	Unit	Biological Reference interval

**TRIGLYCERIDES/HDL RATIO: SERUM** RATIO 3.00 - 5.00 2.28<sup>L</sup> by CALCULATED, SPECTROPHOTOMETRY

#### **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 interval
	LIV	ER FUNCTIO	N TEST (COMPLETE)	
BILIRUBIN TOTAL: S		1.02	mg/dL	INFANT: 0.20 - 8.00 ADULT: 0.00 - 1.20
	CONJUGATED): SERUM	0.55 <sup>H</sup>	mg/dL	0.00 - 0.40
•	(UNCONJUGATED): SERUM	0.47	mg/dL	0.10 - 1.00
SGOT/AST: SERUM		33.03	U/L	7.00 - 45.00
SGPT/ALT: SERUM	/RIDOXAL PHOSPHATE /RIDOXAL PHOSPHATE	23.94	KR U/L	0.00 - 49.00
AST/ALT RATIO: SER	M	1.38	RATIO	0.00 - 46.00
ALKALINE PHOSPHA		104.76	U/L	40.0 - 130.0
GAMMA GLUTAMYL by SZASZ, SPECTRO	TRANSFERASE (GGT): SERUM	15.26	U/L	0.00 - 55.0
TOTAL PROTEINS: S		8.13 <sup>H</sup>	gm/dL	6.20 - 8.00
ALBUMIN: SERUM	GREEN	2.4 <sup>L</sup>	gm/dL	3.50 - 5.50
GLOBULIN: SERUM	ECTROPHOTOMETRY	5.73 <sup>H</sup>	gm/dL	2.30 - 3.50
A : G RATIO: SERUN		0.42 <sup>L</sup>	RATIO	1.00 - 2.00

by CALCULATED, SPECTROPHOTOM

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).

PROGNOSTIC	SIGNIFICANCE:

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
	KIC	ONEY FUNCTIO	N TEST (COMPLETE)	
UREA: SERUM by UREASE - GLUTAM	ATE DEHYDROGENASE (GLDH)	59.61 <sup>H</sup>	mg/dL	10.00 - 50.00
CREATININE: SERUM by ENZYMATIC, SPECT		0.58	mg/dL	0.40 - 1.40
BLOOD UREA NITRO		27.86 <sup>H</sup>	mg/dL	7.0 - 25.0
BLOOD UREA NITRO RATIO: SERUM by CALCULATED, SPE	GEN (BUN)/CREATININE	48.03 <sup>H</sup>	RATIO	10.0 - 20.0
UREA/CREATININE R.	ATIO: SERUM	102.78	RATIO	
URIC ACID: SERUM	E PEROXIDASE	6.34	mg/dL	3.60 - 7.70
CALCIUM: SERUM by arsenazo III, spec	CTROPHOTOMETRY	8.62	mg/dL	8.50 - 10.60
PHOSPHOROUS: SERI	MU	2.77	mg/dL	2.30 - 4.70

CALCIUM: SERUM by ARSENAZO III, SPECTROPHOTOMETRY	8.62	mg/dL	8.50 - 10.60
PHOSPHOROUS: SERUM by phosphomolybdate, spectrophoto	2.77 DMETRY	mg/dL	2.30 - 4.70
<u>ELECTROLYTES</u>			
SODIUM: SERUM by ISE (ION SELECTIVE ELECTRODE)	134.8 <sup>L</sup>	mmol/L	135.0 - 150.0
POTASSIUM: SERUM by ISE (ION SELECTIVE ELECTRODE)	4.2	mmol/L	3.50 - 5.00
CHLORIDE: SERUM by ISE (ION SELECTIVE ELECTRODE)	101.1	mmol/L	90.0 - 110.0
ESTIMATED GLOMERULAR FILTERATIO	<u>N RATE</u>		

ESTIMATED GLOMERULAR FILTERATION RATE 104.9 (eGFR): SERUM 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.



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Test Name	Value	Unit	Biological Reference interval

3. GI haemorrhage.

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

NAME	: Mr. KULWANT SINGH		
AGE/ GENDER	: 70 YRS/MALE	PATIENT ID	: 1587697
<b>COLLECTED BY</b>	:	<b>REG. NO./LAB NO.</b>	: 122409130021
<b>REFERRED BY</b>	:	<b>REGISTRATION DATE</b>	: 13/Sep/2024 03:49 PM
BARCODE NO.	: 12504675	<b>COLLECTION DATE</b>	: 13/Sep/2024 04:51PM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITUTE	<b>REPORTING DATE</b>	: 13/Sep/2024 05:47PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBALA CITY	- HARYANA	

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

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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AM			
Test Name		Value	Unit	Biological Reference interval
		CLINICAL PA	THOLOGY	
	URINF RC	OUTINE & MICRO	DSCOPIC EXAMINAT	ΓΙΟΝ
PHYSICAL EXAMINAT				
QUANTITY RECIEVED		20	ml	
	ANCE SPECTROPHOTOMETRY	20		
COLOUR		PALE YELLOW	I	PALE YELLOW
	ANCE SPECTROPHOTOMETRY			
TRANSPARANCY		TURBID		CLEAR
SPECIFIC GRAVITY	TANCE SPECTROPHOTOMETRY	1.02 PK		1.002 - 1.030
	ANCE SPECTROPHOTOMETRY	1.02		1.002 - 1.030
CHEMICAL EXAMINA				
REACTION		ACIDIC		
by DIP STICK/REFLECT	ANCE SPECTROPHOTOMETRY			
PROTEIN		NEGATIVE (-v	re)	NEGATIVE (-ve)
	TANCE SPECTROPHOTOMETRY			
SUGAR		NEGATIVE (-v	re)	NEGATIVE (-ve)
pH	TANCE SPECTROPHOTOMETRY	6		5.0 - 7.5
1	ANCE SPECTROPHOTOMETRY	0		3.0 - 7.3
BILIRUBIN		NEGATIVE (-v	ve)	NEGATIVE (-ve)
by DIP STICK/REFLECT	ANCE SPECTROPHOTOMETRY			
NITRITE		NEGATIVE (-v	re)	NEGATIVE (-ve)
	ANCE SPECTROPHOTOMETRY.			0.2 1.0
UROBILINOGEN	ANCE SPECTROPHOTOMETRY	NOT DETECTI	ED EU/dL	0.2 - 1.0
KETONE BODIES		NEGATIVE (-v	re)	NEGATIVE (-ve)
	ANCE SPECTROPHOTOMETRY		,	
BLOOD		1+		NEGATIVE (-ve)
•	TANCE SPECTROPHOTOMETRY			
ASCORBIC ACID	ANCE SPECTROPHOTOMETRY	NEGATIVE (-v	e)	NEGATIVE (-ve)
MICROSCOPIC EXAM				

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**NOT VALID FOR MEDICO LEGAL PURPOSE** 



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Test Name				
		Value	Unit	Biological Reference interval
RED BLOOD CELLS (F	RBCs) Centrifuged urinary sediment	6-8	Unit /HPF	Biological Reference interval 0 - 3
by MICROSCOPY ON OPUS CELLS				-
by MICROSCOPY ON O PUS CELLS by MICROSCOPY ON O EPITHELIAL CELLS	CENTRIFUGED URINARY SEDIMENT	6-8	/HPF	0 - 3

CRYSTALS NEGATIVE (-ve) NEGATIVE (-ve) by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT CASTS NEGATIVE (-ve) NEGATIVE (-ve) by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT NEGATIVE (-ve) BACTERIA NEGATIVE (-ve) by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT OTHERS **NEGATIVE** (-ve) NEGATIVE (-ve) by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT TRICHOMONAS VAGINALIS (PROTOZOA) ABSENT ABSENT

by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT

\* \* \* End Of Report \*



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