

Regd. Office: Dr Laf Fathcato: Ltd, Block E. Sessor-18, Rohm, New Delhi-11008S Web: www.lapothibbs.com, CIN:1748990L1995PtcD65388

Name : Mr. SANDEEP KUMAR

Lab No. : 462656422 Ref By : IPSC HOSPITAL

Collected : 10/2/2024 9:04:00AM

A/c Status : P

Collected at : IPSC: PAIN & SPINE HOSPITAL

PLOT NO-453, POCKET-1, SECTOR-19, DWARKA NEW DELHI,New Delhi,South West110075DEL

,IND New Delhi Age : 37 Years Gender : Male

Reported : 10/2/2024 5:43:41PM

Report Status : Final
Processed at : DWARKA -2

Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Test Name	Results	Units	Bio. Ref. Interval
LIPID PROFILE, BASIC, SERUM (Spectrophotometry)			
Cholesterol Total	132	mg/dL	<200.00
Triglycerides	64	mg/dL	<150.00
HDL Cholesterol	34	mg/dL	>40.00
LDL Cholesterol,Direct	94	mg/dL	<100.00
VLDL Cholesterol	13	mg/dL	<30.00
Non-HDL Cholesterol	98	mg/dL	<130.00

Note

- 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.
- Lipid Association of India (LAI) recommends screening of all adults above the age of 20 years for Atherosclerotic Cardiovascular Disease (ASCVD) risk factors especially lipid profile. This should be done earlier if there is family history of premature heart disease, dyslipidemia, obesity or other risk factors

ASCVD Risk Stratification & Treatment goals in Indian population

- 1. Indians are at very high risk of developing ASCVD, they usually get the disease at an early age, have a more severe form of the disease and have poorer outcome as compared to the western populations
- 2. Many individuals remain asymptomatic before they get heart attack, ASCVD risk helps to identify high risk individuals even when there is no symptom related to heart disease
- 3. ASCVD risk category helps clinician to decide when to consider therapy and what should be the treatment goal

Treatment Goals as per Lipid Association of India 2020



Page 1 of 10



Regel Office: Dr Lai Fathcato Ltd, Block E, Sessor TB, Rohm, New Dehic 11008S Web: www.laipothlob.com, CIN: 1748990L1995PLD065888

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Delhi-110075

Test Report

Test Name		Results	Units	Bio. Ref. Interva	al
	CONSI	IDER THERAPY	TREATMENT	GOAL	
CATEGORY@	LDL CHOLESTEROL (LDL-C)(mg/dL)	NON HDL CHLOESTEROL (NON HDL-C) (mg/dL)		NON HDL CHLOESTEROL (NON HDL-C) (mg/dL	
Extreme (A)	>=50	>=80	<50 (Indispensable) <30 (Optional)	<80	
Extreme (B)	>=30	>=60	<30	<60	
Very High	>=50	>=80	<50	<80	
High	>=70	>=100	<70	<100	
Moderate	>=100	>=130	<100	<130	
Low	>=130*	>=160*	<100	<130	

^{*} In low risk patient, consider therapy after an initial non-pharmacological intervention for at least 3 months

@To know your risk category click on bit.ly link sent on your registered mobile number, answer the questionnaire, the ASCVD risk report can be downloaded from website

LIVER PANEL 1; LFT,SERUM (Reflectance Photometry)			
AST (SGOT)	27.7	U/L	<50
ALT (SGPT)	28.9	U/L	<50
AST:ALT Ratio	0.96		<1.00
GGTP	13.5	U/L	<55
Alkaline Phosphatase (ALP)	198.60	U/L	30 - 120
Bilirubin Total	1.81	mg/dL	0.30 - 1.20
Bilirubin Direct	0.31	mg/dL	<0.20
Bilirubin Indirect	1.50	mg/dL	<1.10
Total Protein	8.00	g/dL	6.40 - 8.30
Albumin	4.62	g/dL	3.50 - 5.20



Page 2 of 10



Regd. Office: Dr Lid Faths ato: Ltd. Block E. Sessor 18, Rohm, New Delhi-11008S Web: www.laipothlobs.com, CIN: 17489901,1995712065388

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Test Report

Test NameResultsUnitsBio. Ref. IntervalA: G Ratio1.370.90 - 2.00

Note

- 1. In an asymptomatic patient, Non alcoholic fatty liver disease (NAFLD) is the most common cause of increased AST, ALT levels. NAFLD is considered as hepatic manifestation of metabolic syndrome.
- 2. In most type of liver disease, ALT activity is higher than that of AST; exception may be seen in Alcoholic Hepatitis, Hepatic Cirrhosis, and Liver neoplasia. In a patient with Chronic liver disease, AST:ALT ratio>1 is highly suggestive of advanced liver fibrosis.
- 3. In known cases of Chronic Liver disease due to Viral Hepatitis B & C, Alcoholic liver disease or NAFLD, Enhanced liver fibrosis (ELF) test may be used to evaluate liver fibrosis.
- 4. In a patient with Chronic Liver disease, AFP and Des-gamma carboxyprothrombin (DCP)/PIVKA II can be used to assess risk for development of Hepatocellular Carcinoma.

KIDNEY PANEL; KFT,SERUM (Spectrophotometry, Indirect ISE)			
Creatinine	0.80	mg/dL	0.67 - 1.17
GFR Estimated	117	mL/min/1.73m2	>59
GFR Category	G1		
Urea	18.20	mg/dL	17.00 - 43.00
Urea Nitrogen Blood	8.50	mg/dL	6.00 - 20.00
BUN/Creatinine Ratio	11		
Uric Acid	8.60	mg/dL	3.50 - 7.20
Total Protein	8.00	g/dL	6.40 - 8.30
Albumin	4.62	g/dL	3.50 - 5.20
A : G Ratio	1.37		0.90 - 2.00
Calcium, Total	9.71	mg/dL	8.80 - 10.60
Phosphorus	3.36	mg/dL	2.40 - 4.40
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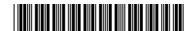
Delhi-110075

Test Report

Test Name Sodium	Results 141.00	Units mEq/L	Bio. Ref. Interval 136.00 - 146.00
Potassium	4.30	mEq/L	3.50 - 5.10
Chloride	102.10	mEq/L	101.00 - 109.00

Note

- 1. Estimated GFR (eGFR) calculated using the 2021 CKD-EPI creatinine equation and GFR Category reported as per KDIGO guideline 2012.
- eGFR category G1 or G2 does not fulfil the criteria for CKD, in the absence of evidence of kidney damage
- 3. The BUN-to-creatinine ratio is used to differentiate prerenal and postrenal azotemia from renal azotemia. Because of considerable variability, it should be used only as a rough guide. Normally, the BUN/creatinine ratio is about 10:1





Regd. Office: Dr Lid Fork.abs Ltd, Block E, Seisor-18, Reiner, New Delhi-110085 Web: www.lapothabs.com, CIN:1748990(1995FLD05788)

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West110075DEL,IND

New Delhi

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Delhi-110075

Test Report

Test Name	Results	Units	Bio. Ref. Interval
HbA1c (GLYCOSYLATED HEMOGLOBIN), BLOOD (HPLC, NGSP Certified)			
HbA1c	5.4	%	4 - 5.6
Estimated average glucose (eAG)	108	mg/dL	

Interpretation

HbA1c result is suggestive of non diabetic adults (>=18 years)/ well controlled Diabetes in a known Diabetic Interpretation as per American Diabetes Association (ADA) Guidelines

Reference Group	Non diabetic adults >=18 years		Diagnosing Diabetes	Therapeutic goals for glycemic control
HbA1c in %	4.0-5.6	5.7-6.4	>= 6.5	<7.0

Note: Presence of Hemoglobin variants and/or conditions that affect red cell turnover must be considered, particularly when the HbA1C result does not correlate with the patient's blood glucose levels.

FACTORS THAT INTERFERE WITH HbA1C MEASUREMENT	FACTORS THAT AFFECT INTERPRETATION OF HBA1C RESULTS
Hemoglobin variants,elevated fetal hemoglobin (HbF) and chemically modified derivatives of hemoglobin (e.g. carbamylated Hb in patients with renal failure) can affect the accuracy of HbAlc measurements	Any condition that shortens erythrocyte survival or decreases mean erythrocyte age (e.g.,recovery from acute blood loss, hemolytic anemia, HbSS, HbCC, and HbSC) will falsely lower HbAlc test results regardless of the assay method used.Iron deficiency anemia is associated with higher HbAlc





Regd. Office: Dr Lid FathLabs Ltd., Block E. Sessor IB., Roman, New Defect 110085 Whole www.ispethiobs.com, CIN: 174899011995FLCW5388

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Test Report

Test Name	Results	Units	Bio. Ref. Interval
PSA (PROSTATE SPECIFIC ANTIGEN), TOTAL, (ECLIA)	SERUM		
PSA, TOTAL	0.245	ng/mL	<1.40

Note

- 1. This is a recommended test for detection of prostate cancer along with Digital Rectal Examination (DRE) in males above 50 years of age.
- 2. False negative / positive results are observed in patients receiving mouse monoclonal antibodies for diagnosis or therapy
- 3. PSA levels may appear consistently elevated / depressed due to the interference by heterophilic antibodies & nonspecific protein binding
- 4. Immediate PSA testing following digital rectal examination, ejaculation, prostatic massage, indwelling catheterization, ultrasonography and needle biopsy of prostate is not recommended as they falsely elevate levels
- 5. PSA values regardless of levels should not be interpreted as absolute evidence of the presence or absence of disease. All values should be correlated with clinical findings and results of other investigations
- 6. Sites of Non-prostatic PSA production are breast epithelium, salivary glands, peri-urethral & anal glands, cells of male urethra & breast milk
- 7. Physiological decrease in PSA level by 18% has been observed in hospitalized / sedentary patients either due to supine position or suspended sexual activity
- 8. The concentration of PSA in a given specimen, determined with assays from different manufacturers, may not be comparable due to differences in assay methods, calibration, and reagent specificity.

Recommended Testing Intervals

- Pre-operatively (Baseline)
- 2-4 days post-operatively
- Prior to discharge from hospital
- Monthly followup if levels are high or show a rising trend

Post Surgery	Frequency of testing
1st year	Every 3 months
2nd year	Every 4 months
3rd year onwards	Every 6 months



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Test Report

Test Name Results Units Bio. Ref. Interval

Clinical Use

- An aid in the early detection of Prostate cancer when used in conjunction with Digital rectal
 examination in males more than 50 years of age and in those with two or more affected first degree
 relatives.
- · Followup and management of Prostate cancer patients
- Detect metastatic or persistent disease in patients following surgical or medical treatment of Prostate cancer

Increased Levels

- Prostate cancer
- Benign Prostatic Hyperplasia
- Prostatitis
- · Genitourinary infections

THYROID PROFILE, FREE, SERUM (ECLIA)			
Free Triiodothyronine (T3, Free)	3.41	pg/mL	2.50 - 4.30
Free Thyroxine (T4, Free)	1.25	ng/dL	0.93 - 1.70
TSH, Ultrasensitive	4.140	μIU/mL	0.27 - 4.20

Note

- 1. 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.
- 2. TSH Values <0.03 μ IU/mL need to be clinically correlated due to presence of a rare TSH variant in some individuals



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Test Report

Test Name	Results	Units	Bio. Ref. Interval
STOOL EXAMINATION, ROUTINE; STOOL, R/E (Light microscopy)			
Colour	Yellow Brown		Brown
Form and Consistency	Semi Formed		Semi Solid
Mucus	Absent		Absent
Visible Blood	Absent		Absent
Reaction	Acidic		Alkaline
Charcot-Leyden Crystals	None Seen		None Seen
Pus Cells	1-2	/hpf	0 - 5
RBC	None Seen	/hpf	None Seen
Macrophages	None Seen		None Seen
Trophozoites	None Seen		None Seen
Cysts	None Seen		None Seen
Helminthic Ova	None Seen		None Seen
Larva	None Seen		None Seen
Other Observations	None Seen		None Seen





Regd. Office: I7-Lid FathLato Ltd, Block E, Selsov Ht, Rohm, New Delhi-11008S Web: www.lapothlabs.com, CIN:1748990L1995PL006S988

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Test Report

Test Name	Results	Units	Bio. Ref. Interval
URINE EXAMINATION, ROUTINE; URINE, R/E (Automated Strip test, Chemical, Light microscopy)			
Physical			
Colour	Yellow		Pale yellow
Specific Gravity	1.030		1.001 - 1.030
рН	5.5		5.0 - 8.0
Chemical			
Proteins	Trace		Negative
Glucose	Negative		Negative
Ketones	Negative		Negative
Bilirubin	Negative		Negative
Urobilinogen	Negative		Negative
Leucocyte Esterase	Positive		Negative
Nitrite	Negative		Negative
Microscopy			
R.B.C.	Negative		0.0 - 2.0 RBC/hpf
Pus Cells	8-10 WBC/HPF		0-5 WBC / hpf
Epithelial Cells	3-5 Epi Cells/hpf		0.0 - 5.0 Epi cells/hpf
Casts	None seen		None seen/Lpf
Crystals	None seen		None seen
Others	None seen		None seen
Result Rechecked, Please Correlate Clinically.			





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Test Report

Test Name Results Units Bio. Ref. Interval

DMC NO 69098

Dr. Arohi Gupta MBBS,MD Pathology Chief of Laboratory Dr Lal PathLabs Ltd

-----End of report ----



IMPORTANT INSTRUCTIONS

•Test results released pertain to the specimen submitted. •All test results are dependent on the quality of the sample received by the Laboratory. •Laboratory investigations are only a tool to facilitate in arriving at a diagnosis and should be clinically correlated by the Referring Physician. •Report delivery may be delayed due to unforeseen circumstances. Inconvenience is regretted. •Certain tests may require further testing at additional cost for derivation of exact value. Kindly submit request within 72 hours post reporting. •Test results may show interlaboratory variations. •The Courts/Forum at Delhi shall have exclusive jurisdiction in all disputes /claims concerning the test(s). • or results of test(s). •Test results are not valid for medico legal purposes. •This is computer generated medical diagnostic report that has been validated by Authorized Medical Practitioner/Doctor. •The report does not need physical signature.

(#) Sample drawn from outside source.

If Test results are alarming or unexpected, client is advised to contact the Customer Care immediately for possible remedial action.

Tel: +91-11-49885050,Fax: - +91-11-2788-2134, E-mail: lalpathlabs@lalpathlabs.com



Page 10 of 10



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4/0 abdominal . Terberculous

"A Unit of Surange Healthcare North India Pvt. Ltd."

MR. SHADEEP KIMAR 37400001M

for Health Check up No complaints

No 4/0 any orne chrome Heness No 4/0 any surgenical P/A

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W-- 113kg

PR - 99/snus

CHEST (I/E) -

BP - 1/8/81 ming

Example Spain Specialist





"A Unit of Surange Healthcare North India Pvt. Ltd"

Radiology No.

8940/OPDPB23DL

Patient Name

Guardian Name

Referred By

Mr. SANDEEP KUMAR

: Dr. .

Date

10-Feb-2024

Age/Sex

37Y Male

UHID No.

8557/UHID23DL

Mobile No.

7349380636

X-RAY CHEST

Indication:-Routine check up.

Image quality:-

No evidence of rotation.

PA view. Normal penetration.

Airway:-

Trachea central.

Carina & bronchi are normal.

No hilar abnormality.

Lung fields:- Clear.

Cardiac:-

Cardiac borders are visible.

Normal heart size.

Diaphragm:- Costophrenic angles on right & left are normal.

Cardiophrenic angles on right & left are normal.

Diaphragm portion are normal.

Bony cage:- No evidence of bony lesion/fracture seen.

No evidence of cervical ribs seen.

Impression: No significant abnormality detected.

Dr. Harshita Surange MBBS,DMRD(RADIODIAGNOSIS DIPLOMA IN MSK, UCAM(Spain) Reg.No. MCI/16522,DMC/18402









Regil, Office: Dr List FuthLate Ltd., Block-E., Sector-18, Rorini, New Detro 110005 C WWW. LIQUETURES. COM. CWI. L.74079CK PROFECOLSTON

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LIPID PROFILE, BASIC (Spectrophotometry)	SERUM			
Cholesterol Total		132	mg/dL	<200.00
Triglycerides		64 +	mg/dL	<150.00
HDL Cholesterol		34	mg/dL	>40.00
LDL Cholesterol, Direct		94	mg/dL	<100.00
VLDL Cholesterol		13	mg/dL	<30.00
Non-HDL Cholesterol		98	mg/dL	<130.00

Note

- 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. Lipid Association of India (LAI) recommends screening of all adults above the age of 20 years for Atherosclerotic Cardiovascular Disease (ASCVD) risk factors especially lipid profile. This should be done earlier if there is family history of premature heart disease, dyslipidemia, obesity or other risk factors

ASCVD Risk Stratification & Treatment goals in Indian population

- Indians are at very high risk of developing ASCVD, they usually get the disease at an early age, have a more severe form of the disease and have poorer outcome as compared to the western populations
- 2. Many individuals remain asymptomatic before they get heart attack, ASCVD risk helps to identify high risk individuals even when there is no symptom related to heart disease
- 3. ASCVD risk category helps clinician to decide when to consider therapy and what should be the treatment goal

Treatment Goals as per Lipid Association of India 2020





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Test Name		Results	Units	Bio. Ref. Interval
ASCVD RISK	CONS	IDER THERAPY	TREATMENT	GOAL
CATEGORY®	LDL CHOLESTEROL (LDL-C)(mg/dL)	NON HDL CHLOESTEROL (NON HDL-C) (mg/dL)	LDL CHOLESTEROL (LDL-C)(mg/dL)	NON HDL CHLOESTEROL (NON HDL-C) (mg/dL
Extreme (A)	>=50	>=80	<50 (Indispensable) <30 (Optional)	<80
Extreme (B)	>=30	>=60	<30	<60
very High	>=50	>=80	<\$0	<80
нigh	>=70	>=100	<70	<100
Moderate	>=100	>=130	<100	<130
Low	>=130*	>=160*	<100	<130

^{*} In low risk patient, consider therapy after an initial non-pharmacological intervention for at least 3 months

@To know your risk category click on bit.ly link sent on your registered mobile number, answer the questionnaire, the ASCVD risk report can be downloaded from website

LIVER PANEL 1; LFT,SERUM (Reflectance Photometry)			
AST (SGOT)	27.7	U/L	<50
ALT (SGPT)	28.9	U/L -	<50
AST:ALT Ratio	0.96		<1.00
GGTP	13.5	U/L	<55
Alkaline Phosphatase (ALP)	198.60	U/L	30 - 120
Bilirubin Total	1.81	mg/dL	0.30 + 1.20
Bilirubin Direct	0.31	,mg/dL	<0.20
Bilirubin Indirect	1.50	mg/dL	<1.10
Total Protein	8.00	g/dL	6.40 - 8.30
Albumin	4.62	g/dL	3.50 - 5.20

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Age : 37 Years Gender Male

Reported : 10/2/2024 5:43:41PM

Report Status : Final Processed at : DWARKA-2

Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Test Name Results Units A: G Ratio 1.37

Bio. Ref. Interval

0.90 - 2.00

Note

- 1. In an asymptomatic patient, Non alcoholic fatty liver disease (NAFLD) is the most common cause of increased AST, ALT levels. NAFLD is considered as hepatic manifestation of metabolic syndrome.
- 2. In most type of fiver disease, ALT activity is higher than that of AST; exception may be seen in Alcoholic Hepatitis, Hepatic Cirrhosis, and Liver neoplasia. In a patient with Chronic liver disease, AST:ALT ratio>1 is highly suggestive of advanced liver fibrosis.
- 3. In known cases of Chronic Liver disease due to Viral Hepatitis B & C, Alcoholic liver disease or NAFLD, Enhanced liver fibrosis (ELF) test may be used to evaluate liver fibrosis.
- 4. In a patient with Chronic Liver disease, AFP and Des-gamma carboxyprothrombin (DCP)/PIVKA II can be used to assess risk for development of Hepatocellular Carcinoma.

	THE RESERVE THE PARTY OF THE PA	HI H	Page 3 of 10
Phosphorus	3.36	mg/dL	2.40 - 4.40
Calcium, Total	9.71	mg/dL	8.80 - 10.60
A : G Ratio	1.37		0.90 - 2.00
Albumin *	4.62	g/dL	3.50 - 5.20
Total Protein	8.00	g/dL	6.40 - 8.30
Uric Acid	8.60	mg/dL	3.50 - 7.20
BUN/Creatinine Ratio	11		
Urea Nitrogen Blood	8.50	mg/dL	6.00 - 20.00
Urea	18.20	mg/dL	17.00 - 43.00
GFR Category	G1		
GFR Estimated	117	mL/min/1.73m2	>59
Creatinine	0.80	mg/dL	0.67 - 1,17
KIDNEY PANEL; KFT,SERUM (Spectrophotometry, Indirect ISE)			



Regd, Office: Dr Let francais Ltd, Block-E, Setter-18, Roters, New Clefs-1 (1006) Web: www.lepstockis.com, ON: L74999Ck.19999C065369

Name : Mr. SANDEEP KUMAR

Lab No. : 462656422

Ref By : IPSC HOSPITAL

Collected : 10/2/2024 9:04:00AM

A/c Status : P

Collected at : IPSC: PAIN & SPINE HOSPITAL

PLOT NO-453, POCKET-1, SECTOR-19, DWARKA

NEW DELHI, New Delhi, South West 10075DEL

,IND * New Delhi Age : 37 Years Gender : Male

Reported : 10/2/2024 5:43:41PM

Report Status : Final

Processed at : DWARKA -2

Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Test Name	Results	Units	Bio. Ref. Interval
Sodium	141.00	mEq/L	136.00 - 146.00
Potassium	4,30	mEq/L	3.50 - 5.10
Chloride	102.10	mEq/L	101.00 - 109.00

Note

- Estimated GFR (eGFR) calculated using the 2021 CKD-EPI creatinine equation and GFR Category reported as per KDIGO guideline 2012.
- eGFR category G1 or G2 does not fulfil the criteria for CKD, in the absence of evidence of kidney damage *
- The BUN-to-creatinine ratio is used to differentiate prerenal and postrenal azotemia from renal
 azotemia. Because of considerable variability, it should be used only as a rough guide. Normally, the
 BUN/creatinine ratio is about 10:1



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Name ; Mr. SANDEEP KUMAR

Lab No. : 462656422

Ref By : IPSC HOSPITAL

Collected : 10/2/2024 9:04:00AM

A/c Status ; P

Collected at : IPSC : PAIN & SPINE HOSPITAL

PLOT NO-453, POCKET-1, SECTOR-19, DWARKA NEW DELHI, New Delhi, South

West110075DEL JND

New Delhi

Age : 37 Years Gender : Male

Reported : 10/2/2024 5:43:41PM

Report Status : Final Processed at : DWARKA -2

Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Test Name	Results	Units	Bio. Ref. Interval
HbA1c (GLYCOSYLATED HEMOGLOBIN), BLOOD (HPLC, NGSP Certified)			
HbA1c	5.4	%	4 - 5.6
Estimated average glucose (eAG)	108	mg/dL	

Interpretation

HbA1c result is suggestive of non diabetic adults (>=18 years)/ well controlled Diabetes in a known Diabetic Interpretation as per American Diabetes Association (ADA) Guidelines

Reference Group	Non diabetic	At risk	Diagnosing	Therapeutic goals
	adults >=18 years	(Prediabetes)	Diabetes	for glycemic control
HbAlc in %	4.0-5.6	5.7-6.4	>= 6.5	<7.0

Note: Presence of Hemoglobin variants and/or conditions that affect red cell turnover must be considered, particularly when the HbA1C result does not correlate with the patient's blood glucose levels.

FACTORS THAT INTERFERE WITH HBA1C FACTORS THAT AFFECT INTERPRETATION OF HBA1C RESULTS

Hemoglobin variants, elevated fetal hemoglobin (HbF) and chemically modified derivatives of hemoglobin (e.g. carbamylated Hb in patients with renal failure) can affect the accuracy of HbAlc measurements Any condition that shortens erythrocyte survival or decreases mean erythrocyte age (e.g., recovery from acute blood loss, hemolytic anemia, HbSS, HbCC, and HbSC) will falsely lower HbAlc test results regardless of the assay method used. Iron deficiency anemia is associated with higher HbAlc



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Name -

: Mr. SANDEEP KUMAR

Lab No.

: 462656422

Ref By Collected : IPSC HOSPITAL : 10/2/2024 9:04:00AM

A/c Status

: P

Collected at

. IPSC : PAIN & SPINE HOSPITAL

PLOT NO-453, POCKET-1, SECTOR-19,

DWARKA NEW DELHI, New Delhi, South

West110075DEL ,IND

New Delhi

Age

37 Years

Gender

Male

Reported : 10/2/2024 5:43:41PM

Report Status : Final

Processed at

: DWARKA -2

Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Test Name

Results

Units

Bio. Ref. Interval

PSA (PROSTATE SPECIFIC ANTIGEN), TOTAL, SERUM

(ECLIA)

PSA, TOTAL

0.245

ng/mL

<1.40

Note

- This is a recommended test for detection of prostate cancer along with Digital Rectal Examination (DRE) in males above 50 years of age.
- False negative / positive results are observed in patients receiving mouse monoclonal antibodies for diagnosis or therapy
- PSA levels may appear consistently elevated / depressed due to the interference by heterophilic antibodies & nonspecific protein binding
- Immediate PSA testing following digital rectal examination, ejaculation, prostatic massage, indwelling catheterization, ultrasonography and needle biopsy of prostate is not recommended as they falsely elevate levels
- PSA values regardless of levels should not be interpreted as absolute evidence of the presence or absence of disease. All values should be correlated with clinical findings and results of other investigations
- Sites of Non-prostatic PSA production are breast epithelium, salivary glands, peri-urethral & anal glands, cells of male urethra & breast milk
- Physiological decrease in PSA level by 18% has been observed in hospitalized / sedentary patients either due to supine position or suspended sexual activity
- The concentration of PSA in a given specimen, determined with assays from different manufacturers, may not be comparable due to differences in assay methods, calibration, and reagent specificity.

Recommended Testing Intervals

- Pre-operatively (Baseline)
- · 2-4 days post-operatively
- Prior to discharge from hospital
- Monthly followup if levels are high or show a rising trend

Post Surgery	Frequency of testing
1st year	Every 3 months
2nd year	Every 4 months ·
3rd year onwards	Every 6 months

Page 6 of 10



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Name

: Mr. SANDEEP KUMAR

Lab No.

: 462656422

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PLOT NO-453, POCKET-1, SECTOR-19,

DWARKA NEW DELHI, New Delhi, South

West110075DEL ,IND

New Delhi

Age

: 37 Years

Gender

Male

Reported : 10/2/2024 5:43:41PM

Report Status

: Final .

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Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Test Name

Results

Units

Bio. Ref. Interval

Clinical Use

- An aid in the early detection of Prostate cancer when used in conjunction with Digital rectal
 examination in males more than 50 years of age and in those with two or more affected first degree
 relatives.
- · Followup and management of Prostate cancer patients
- Detect metastatic or persistent disease in patients following surgical or medical treatment of Prostate cancer *

Increased Levels

- Prostate cancer
- Benign Prostatic Hyperplasia
- Prostatitis
- Genitourinary infections

THYPAIN	DDOE!! !	EDEE	SERUM

(ECLIA)

Free Triiodothyronine (T3, Free)

3.41

pg/mL

2.50 - 4.30

Free Thyroxine (T4, Free)

1.25

ng/dL

0.93 - 1.70

TSH, Ultrasensitive

4.140

µIU/mL

0.27 - 4.20

Note

- 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 Values <0.03 μIU/mL need to be clinically correlated due to presence of a rare TSH variant in some individuals

Page 7 of 10



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Name

: Mr. SANDEEP KUMAR

Lab No.

: 462656422

Ref By Collected : IPSC HOSPITAL : 10/2/2024 9:04:00AM

A/c Status

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NEW DELHI, New Delhi, South West110075DEL

PLOT NO-453, POCKET-1, SECTOR-19, DWARKA

New Delhi

Age

: 37 Years

Gender Reported

Male

: 10/2/2024 5:43:41PM

Final

Report Status Processed at

DWARKA -2

Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Yellow Brown

Semi Formed

Absent

Absent

Acidic

1-2

None Seen

Test Name

STOOL EXAMINATION, ROUTINE; STOOL, R/E

(Light microscopy)

Colour

Form and Consistency

Visible Blood

Reaction

Mucus

Charcot-Leyden Crystals

Pus Cells

RBC

Macrophages

Trophozoites

Cysts

Helminthic Ova

Larva

Other Observations

Results

Units

/hpf

/hpf

Bio. Ref. Interval

Brown

Semi Solid

Absent

Absent.

Alkaline

None Seen

0 - 5

None Seen



Regul. Office: Dr Lat Publisher Ltd. Block-E, Sector-18, Roman, False Clerk-110005 Web: www.inputPlates.com, CWI-C24899CL1995PLC065366

Name

: Mr. SANDEEP KUMAR

Lab No.

462656422

Ref By Collected : IPSC HOSPITAL : 10/2/2024 9:04:00AM

A/c Status

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PLOT NO-453, POCKET-1, SECTOR-19, DWARKA NEW DELHI, New Delhi, South West 110075DEL

New Delhi

Age

37 Years

Gender

Male

: 10/2/2024 5:43:41PM Reported

Report Status

Processed at

: Final

: DWARKA -2

Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Test Name	Results	Units	Bio. Ref. Interval
JRINE EXAMINATION, ROUTINE; URINE, R/E Automated Strip test, Chemical, Light microscopy)			
hysical			
Colour	Yellow		Pale yellow
Specific Gravity	1.030	17	1.001 - 1.030
н .	5.5		5.0 - 8.0
Chemical			
Proteins	Trace		Negative
Slucose .	Negative	-	Negative
Cetones	Negative		Negative
Bilirubin	Negative		Negative
Irobilinogen	Negative		Negative
eucocyte Esterase	Positive		Negative
litrite	Negative		Negative
Microscopy	*** :€:		
R.B.C.	Negative		0.0 - 2.0 RBC/hpf
Pus Cells	8-10 WBC/HPF		0-5 WBC / hpf
Epithelial Cells	3-5 Epi Cells/hpf		0.0 - 5.0 Epi cells/hpf
Casts	None seen		' None seen/Lpf
Crystals	None seen		None seen
Others	None seen		None seen
Result Rechecked, Please Correlate Clinically.			



Regal, Office: Dr Lai Particale Ltd. Block E, Sector 18, Robert, New Dette 110005 Web: www.squetisch.com, CM: L74074DL17759,C065300

Name

: Mr. SANDEEP KUMAR

Lab No.

462656422

Ref By

: IPSC HOSPITAL

Collected

: 10/2/2024 9:04:00AM

A/c Status Collected at

: P

: IPSC : PAIN & SPINE HOSPITAL

PLOT NO-453, POCKET-1, SECTOR-19, DWARKA

NEW DELHI, New Delhi, South West110075DEL

JIND

New Deihi

Age ; 37 Years

Gender : Male

Reported : 10/2/2024 5:43:41PM

Report Status : Final

Processed at : DWARKA -2

Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Test Name

Results

Units

Bio, Ref. Interval

DME NO 69098

Dr. Aroni Gupta MBBS MD Pathology Chief of Laboratory Dr Laf PathLaba Ltd

-End of report



IMPORTANT INSTRUCTIONS

*Test results released pertain to the specimen submitted.*All test results are dependent on the quality of the sample received by the Laboratory.

*Laboratory investigations are only a tool to facilitate in arriving at a diagnosis and should be clinically correlated by the Referring Physician.*Report delivery may be delayed due to unforeseen circumstances. Inconvenience is regretted.*Certain tests may require further testing at additional cost for derivation of exact value. Kindly submit request within 72 hours post reporting.*Test results may show interlaboratory variations.*The Courts/Forum at Dethi, shall have exclusive jurisdiction in all disputes/claims concerning the test(s). & or results of test(s).*Test results are not valid for medico legal purposes.*This is computer generated medical diagnostic report that has been validated by Authorized Medical Practitionen/Doctor.*The report does not need physical signature.

(#) Sample drawn from outside source.

If Test results are alarming or unexpected, client is advised to contact the Customer Care immediately for possible remedial action.

Tel: +91-11-49885050,Fax: - +91-11-2788-2134, E-mail: talpathlabs@lalpathlabs.com



Regal: Office : Dr Lie Familiato Ltd., Block-E, Sector-18, Roman, New Clem-1 (CRIS) e www.laquehoec.com, CIN: L748990X.199595.Couq3ee

Name

: Mr. SANDEEP KUMAR

Lab No.

: 462656437

Ref By

. IPSC HOSPITAL

Collected

: 10/2/2024 12:53:00PM

A/c Status

Collected at : IPSC : PAIN & SPINE HOSPITAL

PLOT NO-453, POCKET-1, SECTOR-19, DWARKA

NEW DELHI, New Delhi, South West110075DEL

New Delhi

Age

: 37 Years

Gender

. Male

Reported : 10/2/2024 5:59:10PM

: Final

Report Status Processed at

: DWARKA-2

Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Test Name

Results

Units

Bio. Ref. Interval

GLUCOSE, POST PRANDIAL (PP), 2 HOURS,

PLASMA

(Hexokinase)

81.40

mg/dL

70.00 - 140.00

DMC NO 69098

Dr. Arshi Gusta MBBS,MD Pathology Chief of Laboratory Dr Lai PathLate Ltd

End of report



IMPORTANT INSTRUCTIONS

*Test results released pertain to the specimen submitted.*All test results are dependent on the quality of the sample received by the Laboratory. *Liberatory investigations are only a tool to facilitate in arriving at a diagnosis and should be clinically correlated by the Referring Physician .*Report delivery may be delayed due to unforeseen circumstances. Inconvenience is regretted. *Certain tests may require further testing at additional cost for derivation of exact value. Kindly submit request within 72 hours post reporting, *Test results may show interlaboratory variations, *The Courts/Forum at Delhi shall have exclusive jurisdiction in all disputes/claims concerning the test(s) & or results of test(s).*Test results are not valid for medico legal purposes. *This is computer generated medical diagnostic report that has been validated by Authorized Medical Practitioner /Doctor. *The report does not need physical signature.

(W) Sample drawn from outside source. If Yest results are alarming or unexpected, client is advised to contact the Customer Cars immediately for possible remedial action.

Tel: +91-11-49885050,Fax: - +91-11-2788-2134, E-mail: lalpathlabs@lalpathlabs.com



Regal, Office: Dr Lat Publicate 120, Block E, Sector-18, Rohms, New Debt-110065 IN WASHINGTON COM: LTATIFICA PROPERTIES

Name

: Mr. SANDEEP KUMAR

Lab No.

: 462656449

Ref By

: IPSC HOSPITAL

Collected

: 10/2/2024 1:46:00PM

A/c Status

Collected at : IPSC : PAIN & SPINE HOSPITAL

PLOT NO-453, POCKET-1, SECTOR-19, DWARKA

NEW DELHI, New Delhi, South West110075DEL

New Delhi

Age

+ 37 Years

Gender

: Male

Reported : 10/2/2024 6:58:39PM

: Final Report Status

Processed at : DWARKA-2

Plot No. 60, Sector 12 B, Dwarka-New

Delhi-110075

Test Report

Test Name	Results	Units	Bio. Ref. Interval
ERYTHROCYTE SEDIMENTATION RATE (ESR)	18	mm/hr	0 - 15
(Capillary photometry)			

Note

1. C-Reactive Protein (CRP) is the recommended test in acute inflammatory conditions.

Test conducted on EDTA whole blood at 37°C.

Dr. Arohi Gupta MBBS.MD Pathology Chief of Laboratory Dr Lai PathLabs Ltd



IMPORTANT INSTRUCTIONS

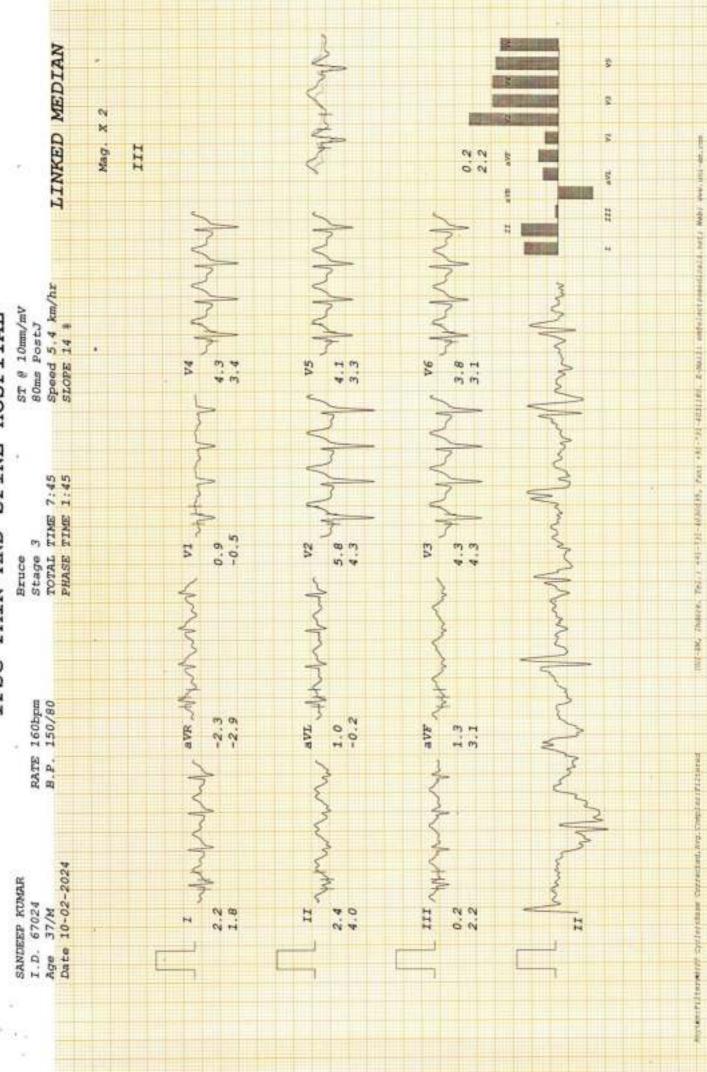
*Test results released pertain to the specimen aubmitted *All test results are dependent on the quality of the sample received by the Laboratory *Laboratory investigations are only a tool to facilitate in arriving at a diagnosis and should be clinically correlated by the Referring Physician .*Report delivery may be delayed due to unforeseen circumstances, inconvenience is regretted. *Certain tests may require further testing at additional cost for derivation of exact value. Kindly submit request within 72 hours post reporting *Test results may show interlaboratory variations *The Courts/Forum as Delhi shall have exclusive jurisdiction in all disputes/claims concerning the test(s) & or results of test(s).*Test results are not valid for medico legal purposes.*This is computer generated medical diagnostic report that has been validated by Authorized Medical Practitioner/Doctor.*The report does not need physical signature.

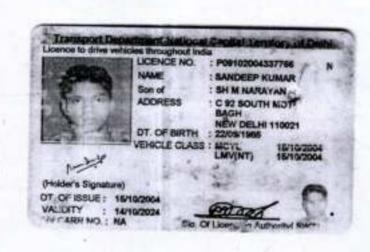
(#) Sample drawn from outside source.

If Test results are alarming or unexpected, client is advised to contact the Customer Care immediately for possible remedial action.

Tel: +91-11-49885050,Fax: - +91-11-2788-2134, E-mail: lalpathlabs@lalpathlabs.com

ID: 16 mr sandcep kumar Male 37Years Req. No. ;	10-02-2024 09:12:52 AM	Diagnosis Information: Sinus rhythm Normal ECG Normal ECG Dr. ANIL SAHOO Dr. ANIL SAHOO Report Confirmed by: Report Confirmed b
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	(Y 4 4
0.67-45Hz AC50 25mm/s H	10mm/mV 2*5.0s+ir CARDIART 9	D VI.44 Glasgow V28.6.7 \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \









7349380636 Mobile No. . .7G : Referred By SSSYUHID23DL UHID No. **Guardian Name** 37Y Male xs2\sgA Mr. SANDEEP KUMAR Patient Name 10-Feb-2024 Date 8940\OPDP823DL Radiology No.

ULTRASOUND OF WHOLE ABDOMAN

in caliber, Intrahepatic bile ducts and CBD are not dilated. Hepatic portal veins and the IVC appear normal The liver is normal in size (1307, in RML) contour however is increased in echotexture.

calculus / wall thickness noted. Gall bladder is adequately distended with normal intraluminal fluid contents. No evidence of

Pancreas is of normal size and contour with normal echotexture.

Cortical thickness is normal. Right kidney is normal in size and position. It shows normal movements with respiration.

No calculus, mass or hydronephyrotic changes seen.

Right kidney measures-99x51mm

Renal artery pulsation appear normal.

Cortical thickness is normal. Left kidney is normal in size and position . It shows normal movements with respiration.

No calculus, mass or hydronephyrotic changes seen.

Left kidney measures-111x50mm

Renal artery pulsation appear normal.

Reg. No. MCI/16522, DMC/18402 DIPLOMA IN MSK, UCAM(Spain) MBBS, DMRD(RADIODIAGNOSIS Dr. Harshita Surange









 Radiology No.
 : 8940/OPDP823DL
 Date
 : 10-Feb-2024

 Patient Name
 : Mr. SANDEEP KUMAR
 Age/Sex
 : 37Y Male

 Guardian Name
 : 8557/UHID23DL

 Referred By
 : Dr.,
 : 7349380636

Spleen is of normal size and shape. Echotexture is normal. No focal lesion is seen.

No evidence of retro-peritoneal lymphadenopathy/ ascites/ pleural effusion noted.

Urinary bladder does not show any calculus or mass lesion. No significant wall thickening noted.

Prostate is of normal size for age with regular contours and normal echo-texture. It measures 26x32x40 mm which is equal to 17cc gms.

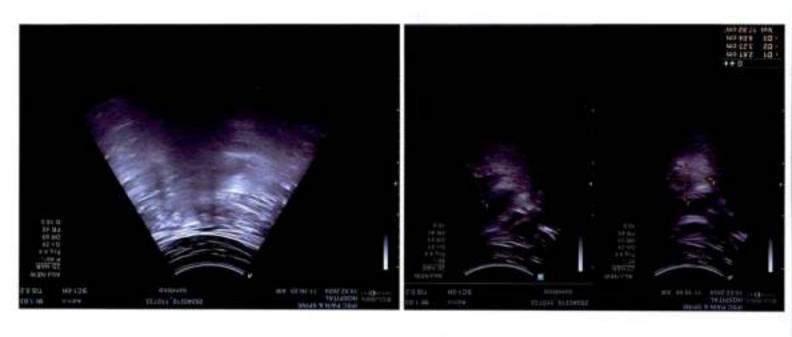
Impressions: Fatty live grade I

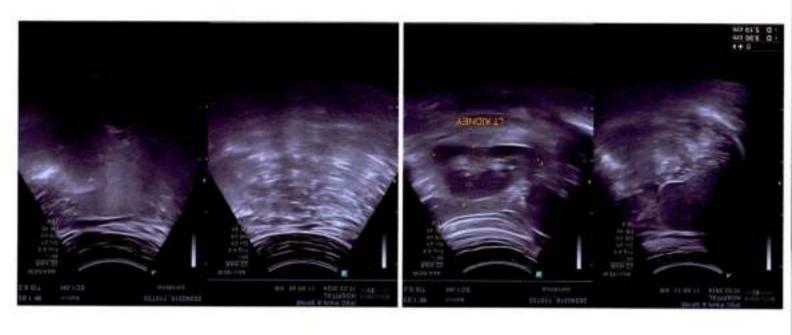


DIPLOMA IN MSK, UCAM(Spain) DIPLOMA IN MSK, UCAM(Spain) PRES, MOI/16522, DMC/18402 Reg. No. MCI/16522









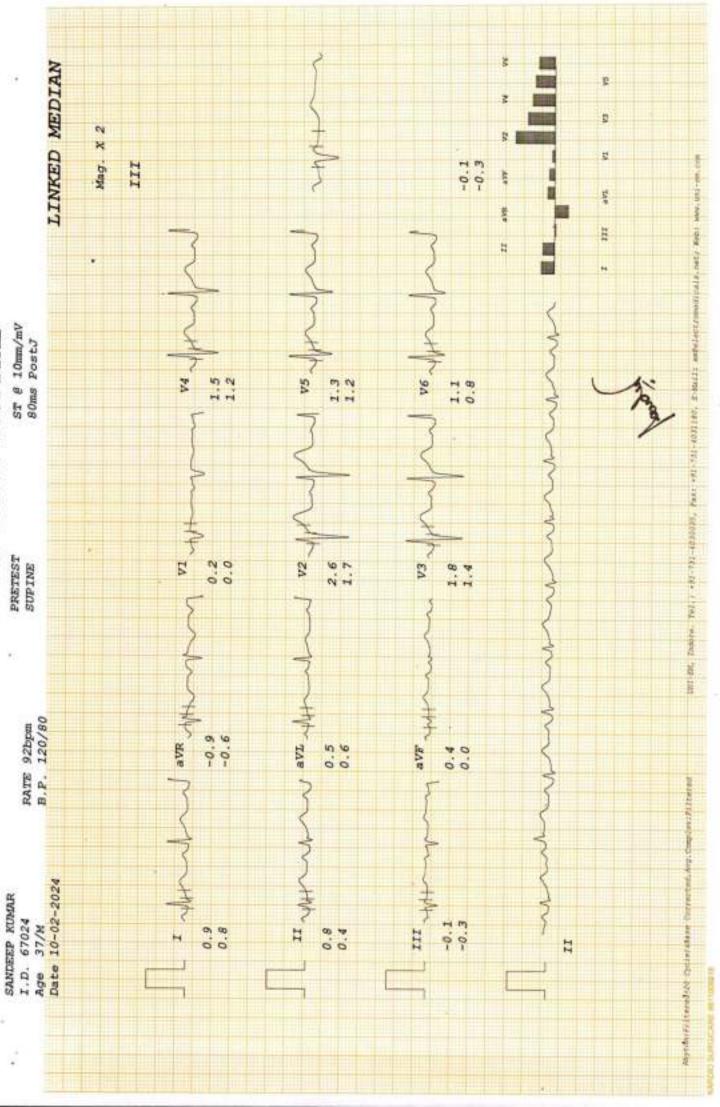


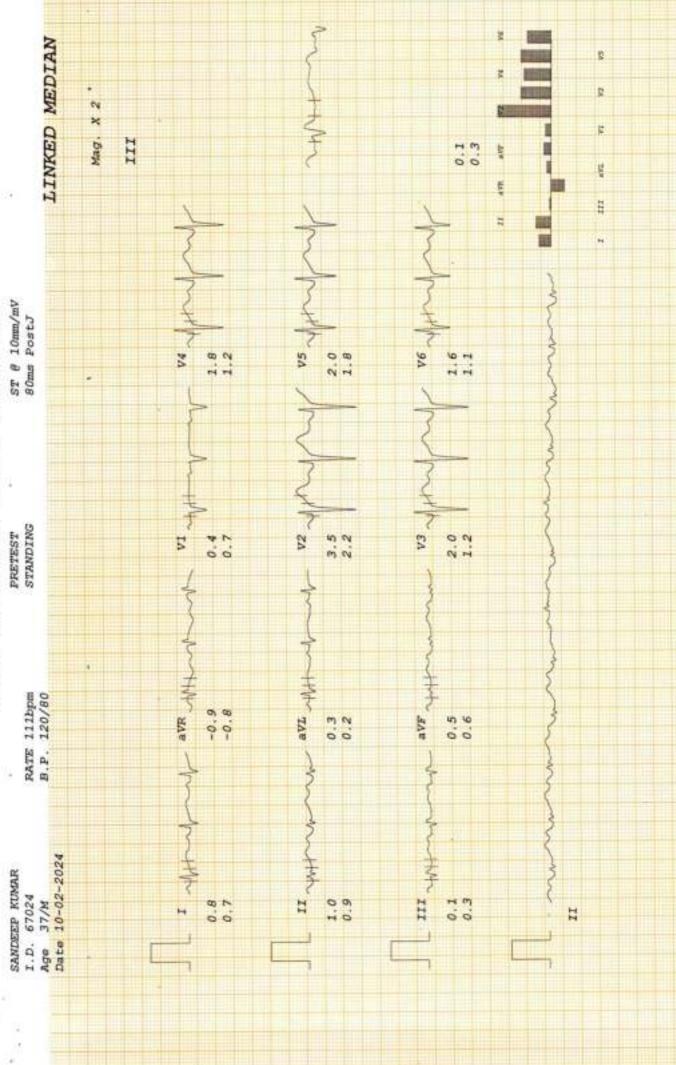
PLOT-453 NEAR SBI BANK SECTOR-19

DWARKA NEW DELHI-110075, PH: 9555437357

SANDEEP KUMAR

SE
2:55 2:55 2.7 10 2:55 2:55 3.4 14 8:0 5:4 8:0 5:4 14 8:0 5:10:10:10:10:10:10:10:10:10:10:10:10:10:
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