

Dr Keerthi Kishore Nagalla

MBBS, MD (General Medicine) Consultant Physician & Diabetologist Reg. No. 64905

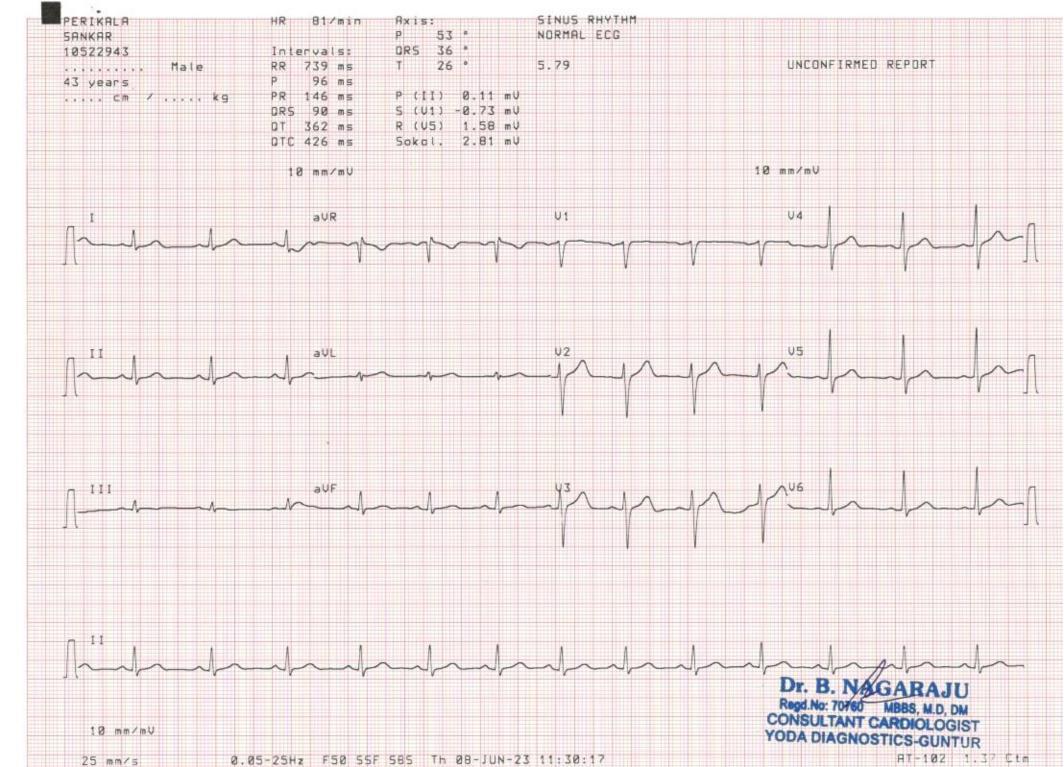
Name: Mr. Dezileala Sankar 23 Age: 43 Apazz Sex: Female Date:0.8./06./

R TEMP: Routine Health checkup B.P: 120/80 Hulling NO COMPLAINT PULSE: 79. 648 HIO Hypertension HEIGHT: ... 60. Cmg 1/ Low salt Diet / Diabetic Die: FBS-116mgld1 PPBS - 226mg/N/ Advice HEAC

Dr. KEERTHI KISHORE NAGALLA Regd.No: 64905 MBBS, M.D. General Medicine CONSULTANT GENERAL PHYSICIAN YODA DIAGNOSTICS-GUNTUR

CONTACT US

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Visit ID	: YGT15209	UHID/MR No	: YGT.0000015084
Patient Name	: Mr. PERIKALA SANKAR	Client Code	: 1409
Age/Gender	: 43 Y 0 M 0 D /M	Barcode No	: 10523951
DOB	:	Registration	: 08/Jun/2023 06:06PM
Ref Doctor	: SELF	Collected	: 08/Jun/2023 06:08PM
Client Name	: MEDI WHEELS	Received	: 08/Jun/2023 06:12PM
Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 08/Jun/2023 06:47PM
Hospital Name	:		

DEPARTMENT OF HAEMATOLOGY						
Test Name	Result	Unit	Biological. Ref. Range	Method		

ESR (ERYTHROCYTE SEDIMENTATION RATE)						
Sample Type : WHOLE BLOOD EDTA						
ERYTHROCYTE SEDIMENTATION RATE	30	mm/1st hr	0 - 15	Capillary Photometry		

COMMENTS:

ESR is an acute phase reactant which indicates presence and intensity of an inflammatory process. It is never diagnostic of a specific disease. It is used to monitor the course or response to treatment of certain diseases. Extremely high levels are found in cases of malignancy, hematologic diseases, collagen disorders and renal diseases.

Increased levels may indicate: Chronic renal failure (e.g., nephritis, nephrosis), malignant diseases (e.g., multiple myeloma, Hodgkin disease, advanced Carcinomas), bacterial infections (e.g., abdominal infections, acute pelvic inflammatory disease, syphilis, pneumonia), inflammatory diseases (e.g. temporal arteritis, polymyalgia rheumatic, rheumatoid arthritis, rheumatic fever, systemic lupus erythematosus [SLE]), necrotic diseases (e.g., acute myocardial infarction, necrotic tumor, gangrene of an extremity), diseases associated with increased proteins (e.g., hyperfibrinogenemia, macroglobulinemia), and severe anemias (e.g., iron deficiency or B12 deficiency).

Falsely decreased levels may indicate: Sickle cell anemia, spherocytosis, hypofibrinogenemia, or polycythemia vera.

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DEPARTMENT OF HAEMATOLOGY							
Test NameResultUnitBiological. Ref. RangeMethod							

BLOOD GROUP ABO & RH Typing Sample Type : WHOLE BLOOD EDTA					
ABO	" B"				
Rh Typing	POSITIVE (+)	/			
Method : Hemagglutination Tube method by forward and reverse grouping					

COMMENTS:

The test will detect common blood grouping system A, B, O, AB and Rhesus (RhD). Unusual blood groups or rare subtypes will not be detected by this method. Further investigation by a blood transfusion laboratory, will be necessary to identify such groups.

Disclaimer: There is no trackable record of previous ABO & RH test for this patient in this lab. Please correlate with previous blood group findings. Advsied cross matching before transfusion

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DEPARTMENT OF HAEMATOLOGY							
Test Name	Result	Unit	Biological. Ref. Range	Method			

CBC(COMPLETE BLOOD COUNT)						
Sample Type : WHOLE BLOOD EDTA						
HAEMOGLOBIN (HB)	15.0	g/dl	13.0 - 17.0	Cyanide-free SLS method		
RBC COUNT(RED BLOOD CELL COUNT)	5.05	million/cmm	4.50 - 5.50	Impedance		
PCV/HAEMATOCRIT	42.2	%	40.0 - 50.0	RBC pulse height detection		
MCV	83.6	fL	83 - 101	Automated/Calculated		
МСН	29.8	pg	27 - 32	Automated/Calculated		
MCHC	35.6	g/dl	32 - 35	Automated/Calculated		
RDW - CV	13.0	%	11.0-16.0	Automated Calculated		
RDW - SD	42.0	fl	35.0-56.0	Calculated		
MPV	9.6	fL	6.5 - 10.0	Calculated		
PDW	16.2	fL	8.30-25.00	Calculated		
PCT	0.231	%	0.15-0.62	Calculated		
TOTAL LEUCOCYTE COUNT	7,000	cells/ml	4000 - 11000	Flow Cytometry		
DLC (by Flow cytometry/Microscopy)						
NEUTROPHIL	54	%	40 - 80	Impedance		
LYMPHOCYTE	37	%	20 - 40	Impedance		
EOSINOPHIL	04	%	01 - 06	Impedance		
MONOCYTE	05	%	02 - 10	Impedance		
BASOPHIL	00	%	0 - 1	Impedance		
PLATELET COUNT	2.40	Lakhs/cumm	1.50 - 4.10	Impedance		

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DEPARTMENT	OF BIO	CHEM	ISTRY
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Result

1.80

Test Name

Unit

Biological. Ref. Range

Method

CLIA

0.30 - 5.60

THYROID PROFILE (T3,T4,TSH)						
Sample Type : SERUM						
T3	0.97	ng/ml	0.60 - 1.78	CLIA		
T4	10.61	ug/dl	4.82-15.65	CLIA		

ulU/mL

INTERPRETATION:

TSH

1. Serum T3, T4 and TSH are the measurements form three components of thyroid screening panel and are useful in diagnosing various disorders of thyroid gland function.

2. Primary hyperthyroidism is accompanied by elevated serum T3 and T4 values along with depressed TSH levels.

3. Primary hypothyroidism is accompanied by depressed serum T3 and T4 values and elevated serum TSH levels.

4. Normal T4 levels accompanied by high T3 levels are seen in patients with T3 thyrotoxicosis. Slightly elevated T3 levels may be found in pregnancy and in estrogen therapy while depressed levels may be encountered in severe illness, malnutrition, renal failure and during therapy with drugs like propanolol and propylthiouracil.

5. Although elevated TSH levels are nearly always indicative of primary hypothyroidism, rarely they can result from TSH secreting pituitary tumors (secondary hyperthyroidism).

6. Low levels of Thyroid hormones (T3, T4 & FT3, FT4) are seen in cases of primary, secondary and tertiary hypothyroidism and sometimes in non-thyroidal illness also. 7. Increased levels are found in Grave's disease, hyperthyroidism and thyroid hormone resistance.

8. TSH levels are raised in primary hypothyroidism and are low in hyperthyroidism and secondary hypothyroidism.

9. REFERENCE RANGE

PREGNANCY	TSH in uIU/ mL
1st Trimester	0.60 - 3.40
2nd Trimester	0.37 - 3.60
3rd Trimester	0.38 - 4.04

(References range recommended by the American Thyroid Association)

Comments:

1. During pregnancy, Free thyroid profile (FT3, FT4 & TSH) is recommended.

2. TSH levels are subject to circadian variation, reaches peak levels between 2-4 AM and at a minimum between 6-10 PM. The variation of the day has influence on the measured serum TSH concentrations.

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DEPARTMENT OF BIOCHEMISTRY					
Test Name	Result	Unit	Biological. Ref. Range	Method	

	LIVER FUNCT	TION TEST(LFT))			
Sample Type : SERUM						
TOTAL BILIRUBIN	1.30	mg/dl	0.3 - 1.2	JENDRASSIK & GROFF		
CONJUGATED BILIRUBIN	0.27	mg/dl	0 - 0.2	DPD		
UNCONJUGATED BILIRUBIN	1.03	mg/dl		Calculated		
S.G.O.T	18	U/L	< 50	KINETIC WITHOUT P5P- IFCC		
S.G.P.T	19	U/L	< 50	KINETIC WITHOUT P5P- IFCC		
ALKALINE PHOSPHATASE	96	U/L	30 - 120	IFCC-AMP BUFFER		
TOTAL PROTEINS	6.4	gm/dl	6.0 - 8.0	Biuret		
ALBUMIN	4.1	gm/dl	3.5 - 5.2	BCG		
GLOBULIN	2.3	gm/dl		Calculated		
A/G RATIO	1.78			Calculated		

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DEPARTMENT OF BIOCHEMISTRY

Result

Test Name

Unit

Biological. Ref. Range

Method

LIPID PROFILE					
Sample Type : SERUM					
TOTAL CHOLESTEROL	120	mg/dl	See Table		Cholesterol oxidase/peroxidase
H D L CHOLESTEROL	38	mg/dl	> 40		Enzymatic/ Immunoinhibiton
L D L CHOLESTEROL	59.8	mg/dl	See Table	e I	Enzymatic Selective Protein
TRIGLYCERIDES	111	mg/dl	See Table	e	GPO
VLDL	22.2	mg/dl	15 - 30		Calculated
T. CHOLESTEROL/ HDL RATIO	3.16		See Table	e	Calculated
TRIGLYCEIDES/ HDL RATIO	2.92	Ratio	< 2.0		Calculated
NON HDL CHOLESTEROL	82	mg/dl	< 130 C		Calculated
Interpretation NATIONAL LIPID ASSOCIATION RECOMMENDATIONS (NLA-2014)	TOTAL CHOLESTER	OL TRIGLYCE	RI DE LDL CHOLESTEROL	NON HD	L ROL
Optimal	<200	<150	<100	<130	
Above Optimal	-	-	100-129	130 - 159	
Borderline High	200-239	150-19		160 - 189	
High	>=240	200-49		190 - 219	9
Very High		>=500) >=190	>=220	

REMARKS	Cholesterol : HDL Ratio	
Low risk	3.3-4.4	
Average risk	4.5-7.1	
Moderate risk	7.2-11.0	
High risk	>11.0	

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. NLA-2014 identifies Non HDL Cholesterol(an indicator of all atherogenic lipoproteins 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.

3.Apolipoprotein B is an optional, secondary lipid target for treatment once LDL & Non HDL goals have been achieved 4. 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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DEPARTMENT	OF BIOCHEMISTRY
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Result

Test Name

Unit

Biological. Ref. Range

Method

HBA1C						
Sample Type : WHOLE BLOOD EDTA						
HBA1c RESULT	6.2	%	Normal Glucose tolerance (non-diabetic): <5.6% Pre-diabetic: 5.7-6.4% Diabetic Mellitus: >6.5%	HPLC		
ESTIMATED AVG. GLUCOSE	131	mg/dl				

Note:

1. Since HbA1c reflects long term fluctuations in the blood glucose concentration, a diabetic patient who is recently under good control may still have a high concentration of HbA1c. Converse is true for a diabetic previously under good control but now poorly controlled .

2. 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, targeting a goal of < 7.0 % may not be appropriate. HbA1c provides an index of average blood glucose levels over the past 8 - 12 weeks and is a much better indicator of long

term glycemic control .

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DEPARTMENT OF BIOCHEMISTRY					
Test NameResultUnitBiological. Ref. RangeMethod					

BLOOD UREA NITROGEN (BUN)						
Sample Type : Serum						
SERUM UREA		15	mg/dL	17 - 43		Urease GLDH
Blood Urea Nitrogen (BUN)		7.0	mg/dl	5 - 25		GLDH-UV

Increased In:

Impaired kidney function, Reduced renal blood flow {CHF, Salt and water depletion, (vomiting, diarrhea, diuresis, sweating), Shock}, Any obstruction of urinary tract, Increased protein catabolism, AMI, Stress

Decreased In:

Diuresis (e.g. with over hydration), Severe liver damage, Late pregnancy, Infancy, Malnutrition, Diet (e.g., low-protein and high-carbohydrate, IV feedings only), Inherited hyperammonemias (urea is virtually absent in blood)

Limitations:

Urea levels increase with age and protein content of the diet.

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DEPARTMENT OF BIOCHEMISTRY				
Test Name	Result	Unit	Biological. Ref. Range	Method

	FBS (GLUCO	OSE FASTING)		
Sample Type : FLOURIDE PLASMA				
FASTING PLASMA GLUCOSE	116	mg/dl	70 - 100	HEXOKINASE
INTERPRETATION:				
Increased In				
 Diabetes Mellitus 				
 Stress (e.g., emotion, burns, shock, 	anesthesia)			
 Acute pancreatitis 	anestnesia)			
Chronic pancreatitis				
 Wernicke encephalopathy (vitamin B1 	l deficiency)			
• Effect of drugs (e.g. corticosteroids,		l, phenytoin, thiazi	des)	
Decreased In				
Decreased III				
Pancreatic disorders				
Extrapancreatic tumors				
 Endocrine disorders 				
Malnutrition				
 Hypothalamic lesions 				
Alcoholism				
 Endocrine disorders 				

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	DE	PARTMENT O	F BIOCHEMI	ISTRY	
Test Name		Result	Unit	Biological. Ref. Range	Method

PPBS (POST PRANDIAL GLUCOSE)							
Sample Type : FLOURIDE PLASMA							
POST PRANDIAL PLASMA GLUCOSE	226	mg/dl	<140	HEXOKINASE			
INTERPRETATION:							
Increased In Diabetes Mellitus Stress (e.g., emotion, burns, shock, anesthes Acute pancreatitis Chronic pancreatitis Wernicke encephalopathy (vitamin B1 deficient Effect of drugs (e.g. corticosteroids, estrogent	ncy)	ytoin, thiazides)					
 Pancreatic disorders 							
Extrapancreatic tumors							
Endocrine disorders							
Malnutrition							
 Hypothalamic lesions Alcoholism 							
Endocrine disorders							

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DEPARTMENT OF BIOCHEMISTRY					
Test Name	Result	Unit	Biological. Ref. Range	Method	

SERUM CREATININE				
Sample Type : SERUM				
SERUM CREATININE	0.81	mg/dl	0.67 - 1.17	KINETIC-JAFFE
				· · · · · · · · · · · · · · · · · · ·

Increased In:

- Diet: ingestion of creatinine (roast meat), Muscle disease: gigantism, acromegaly,
- Impaired kidney function.

Decreased In:

- Pregnancy: Normal value is 0.4-0.6 mg/dL. A value >0.8 mg/dL is abnormal and should alert the clinician to further diagnostic evaluation.
- Creatinine secretion is inhibited by certain drugs (e.g., cimetidine, trimethoprim).

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Test Name	Result	Unit	Biological. Ref. Range	Method		

	URIC AC	ID -SERUM		
Sample Type : SERUM				
SERUM URIC ACID	5.4	mg/dl	3.5 - 7.20	URICASE - PAP
Uric acid is the final product of purine m	etabolism in the hum	an organism. U	ric acid measurements are u	sed in the diagnosis

Uric acid is the final product of purine metabolism in the human organism. Uric acid measurements are used in the diagnosis and treatment of numerous renal and metabolic disorders, including renal failure, gout, leukemia, psoriasis, starvation or other wasting conditions, and of patients receiving cytotoxic drugs.

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Test Name	Result	Unit	Biological. Ref. Range	Method	

	BUN/CREAT	ININE RATIC		
Sample Type : SERUM				
Blood Urea Nitrogen (BUN)	7.0	mg/dl	5 - 25	GLDH-UV
SERUM CREATININE	0.81	mg/dl	0.67 - 1.17	KINETIC-JAFFE
BUN/CREATININE RATIO	8.60	Ratio	6 - 25	Calculated

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Result

DEPARTMENT OF CLINICAL PATHOLOGY

Test Name

Unit

Biological. Ref. Range

Method

CU	E (COMPLETE U	RINE EXAMI	INATION)	
Sample Type : SPOT URINE				
PHYSICAL EXAMINATION				
TOTAL VOLUME	20 ML	ml		
COLOUR	PALE YELLOW			
APPEARANCE	CLEAR			
SPECIFIC GRAVITY	1.025		1.003 - 1.035	Bromothymol Blue
CHEMICAL EXAMINATION				
pH	6.8		4.6 - 8.0	Double Indicator
PROTEIN	NEGATIVE		NEGATIVE	Protein - error of Indicators
GLUCOSE(U)	NEGATIVE		NEGATIVE	Glucose Oxidase
UROBILINOGEN	NEGATIVE	mg/dl	< 1.0	Ehrlichs Reaction
KETONE BODIES	NEGATIVE		NEGATIVE	Nitroprasside
BILIRUBIN - TOTAL	NEGATIVE		Negative	Azo-coupling Reaction
BLOOD	NEGATIVE		NEGATIVE	Tetramethylbenzidine
LEUCOCYTE	NEGATIVE		Negative	by an azo-coupling reaction
NITRITE	NEGATIVE		NEGATIVE	Diazotization Reaction
MICROSCOPIC EXAMINATION				
PUS CELLS	1 - 2	cells/HPF	0-5	
EPITHELIAL CELLS	1 - 2	/hpf	0 - 15	
RBCs	NIL	Cells/HPF	Nil	
CRYSTALS	NIL	Nil	Nil	
CASTS	NIL	/HPF	Nil	
BUDDING YEAST	NIL		Nil	

OTHER

BACTERIA

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Nil

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NIL

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Ref Doctor	: SELF	Collected	: 08/Jun/2023 06:08PM
Client Name	: MEDI WHEELS	Received	: 08/Jun/2023 06:12PM
Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 08/Jun/2023 06:17PM
Hospital Name	:		

DEPARTMENT OF CLINICAL PATHOLOGY					
Test Name	Result	Unit	Biological. Ref. Range	Method	

*** End Of Report ***

Verified By : G SUMALATHA

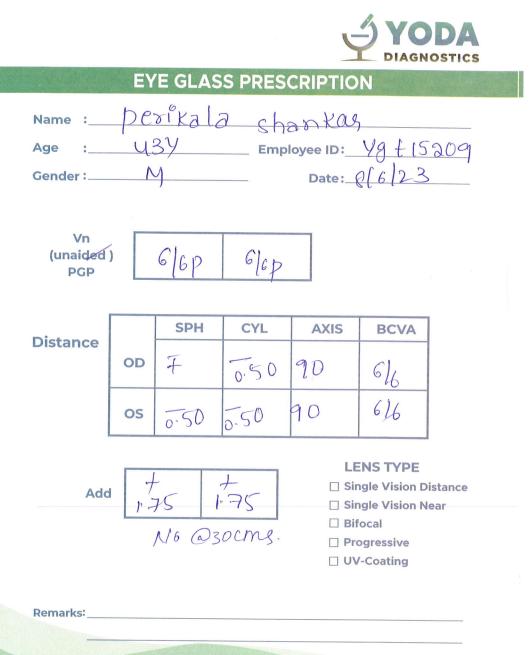


Approved By :

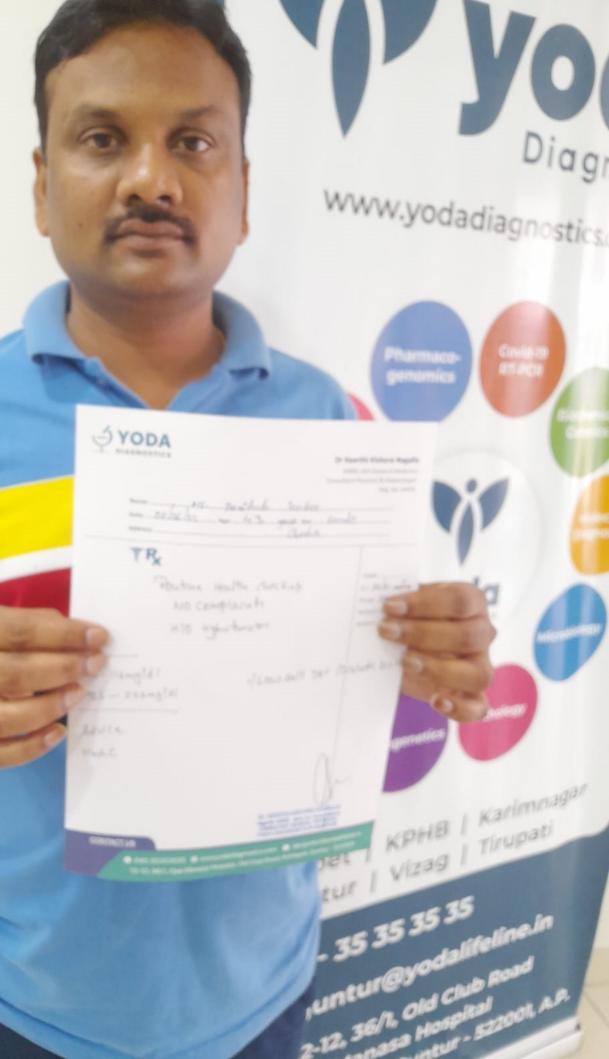
e falte 7.05

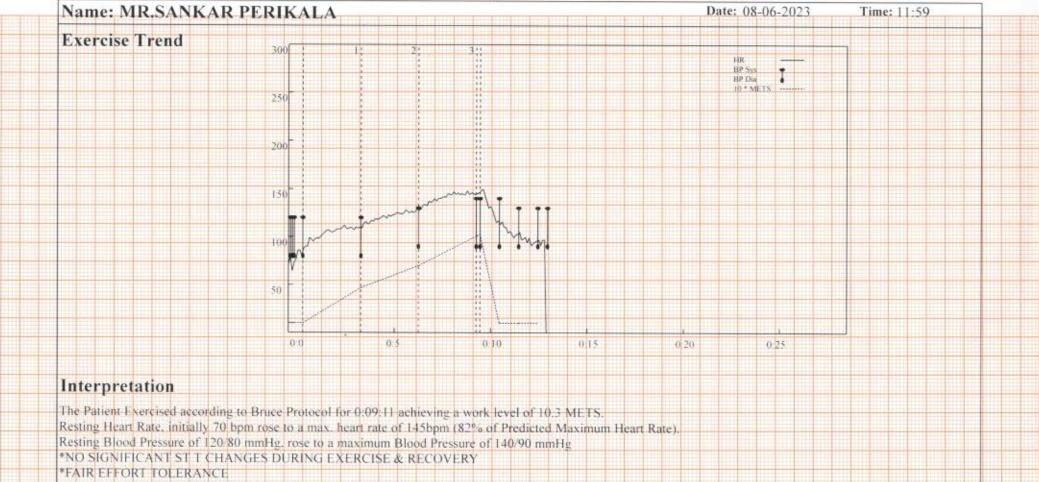
Dr. Sumalatha MBBS,DCP Consultant Pathologist



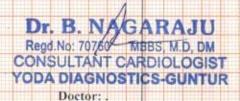


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*TEST IS NEGATIVE FOR EXERCISE INDUCID ISCHEMIA

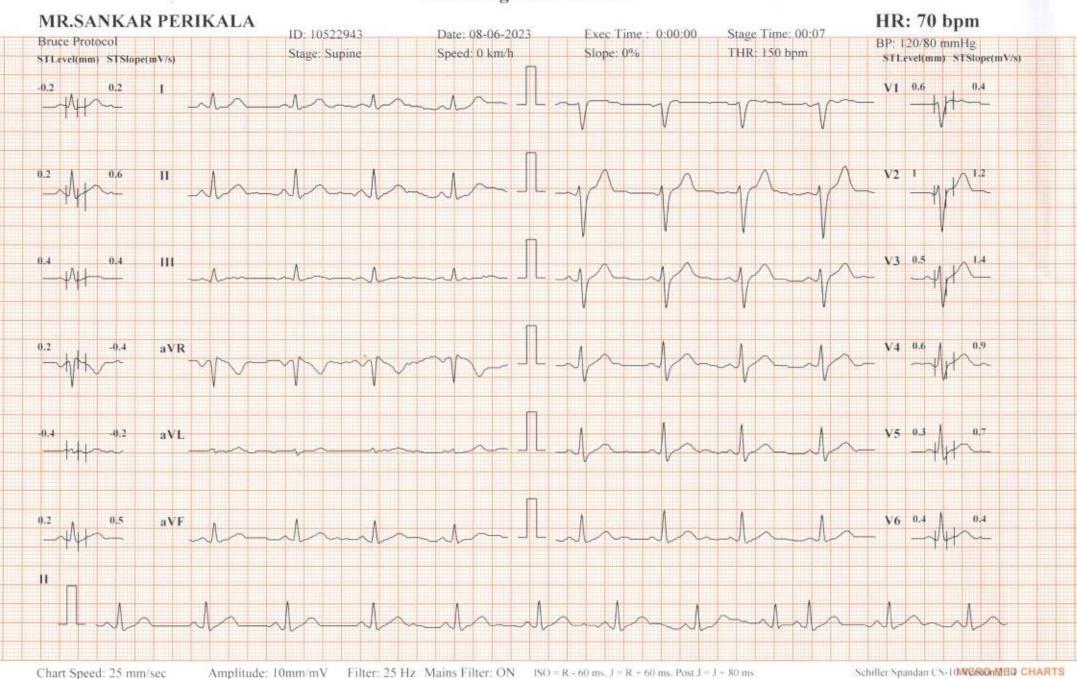


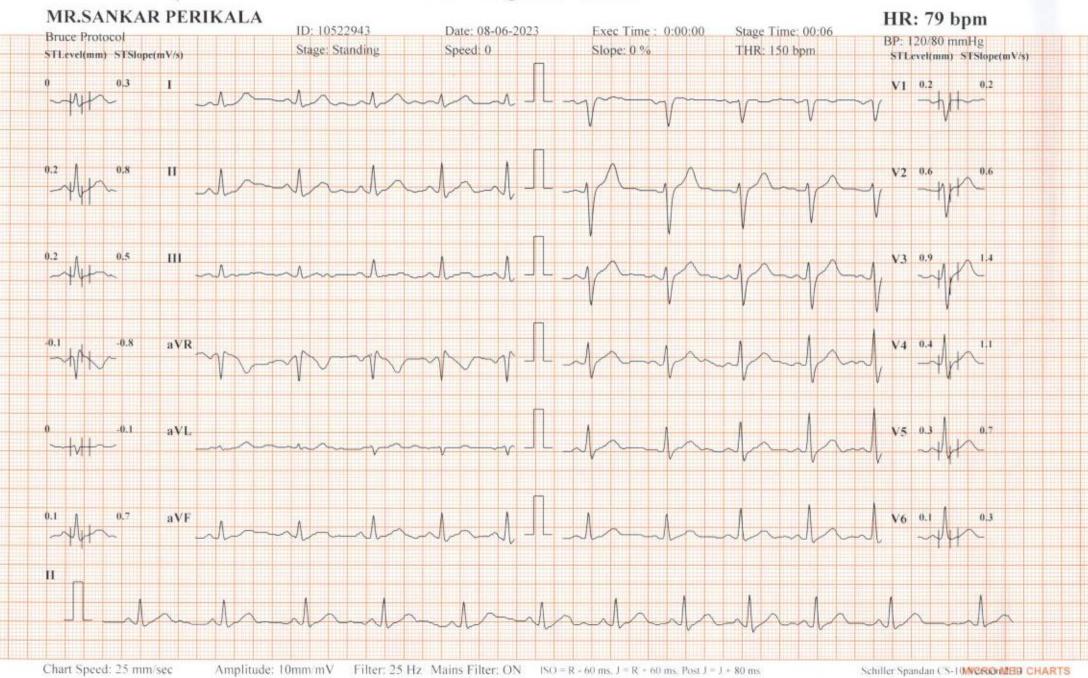
Ref. Doctor: DR SELF

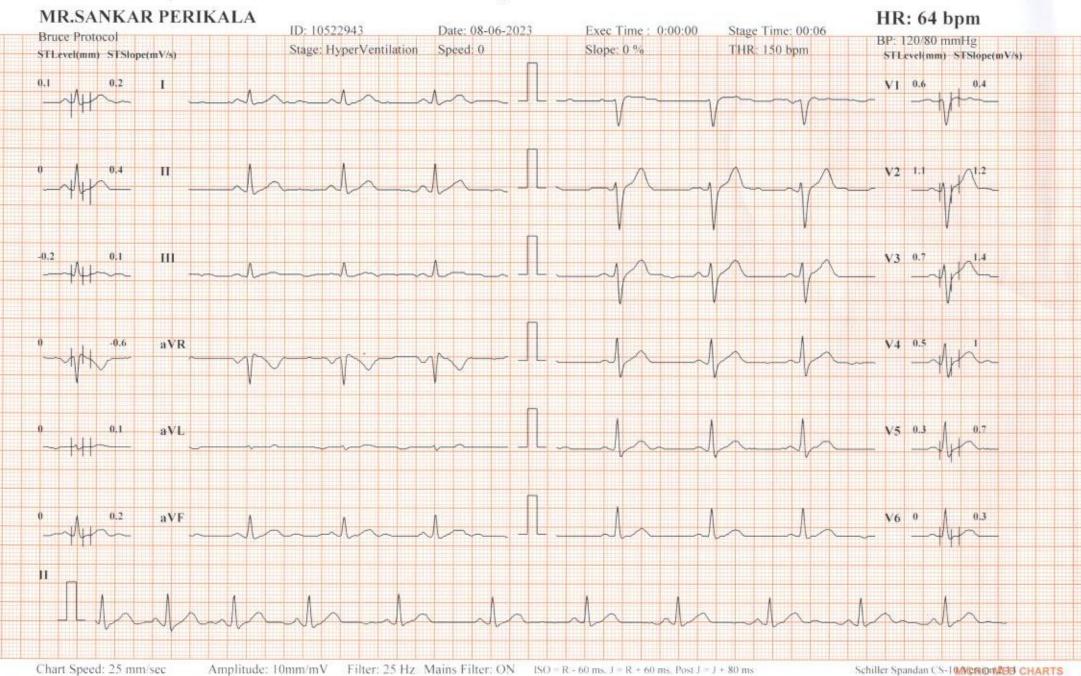
Schiller Spandan CS-10 Version 2 14

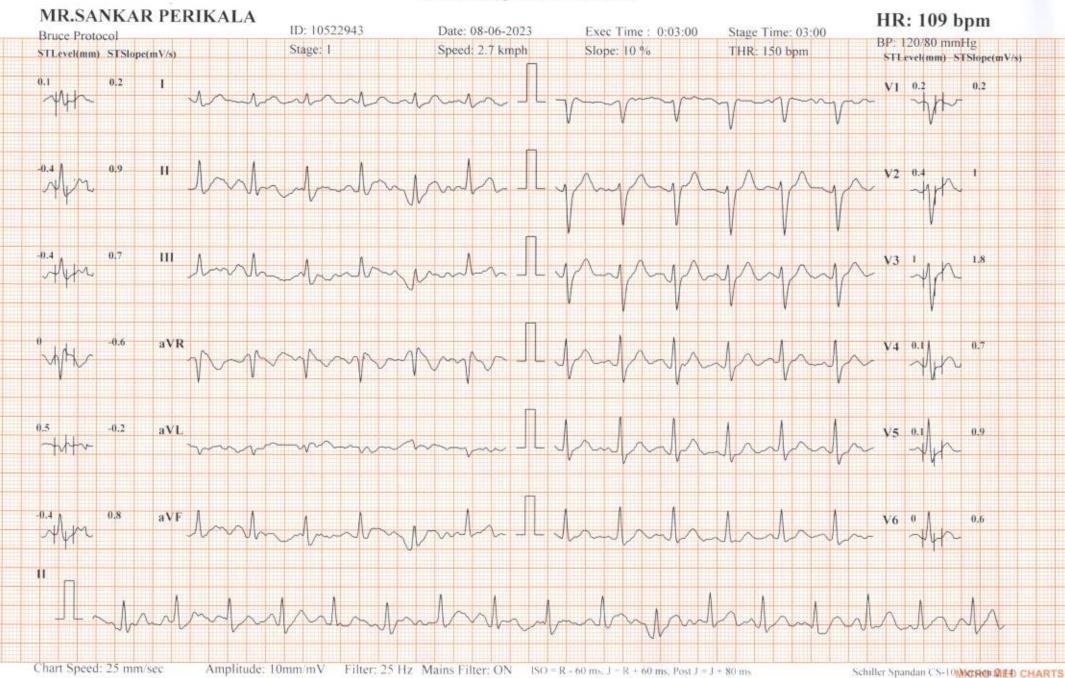
Summary Report edited by User MICRO MED CHARTS

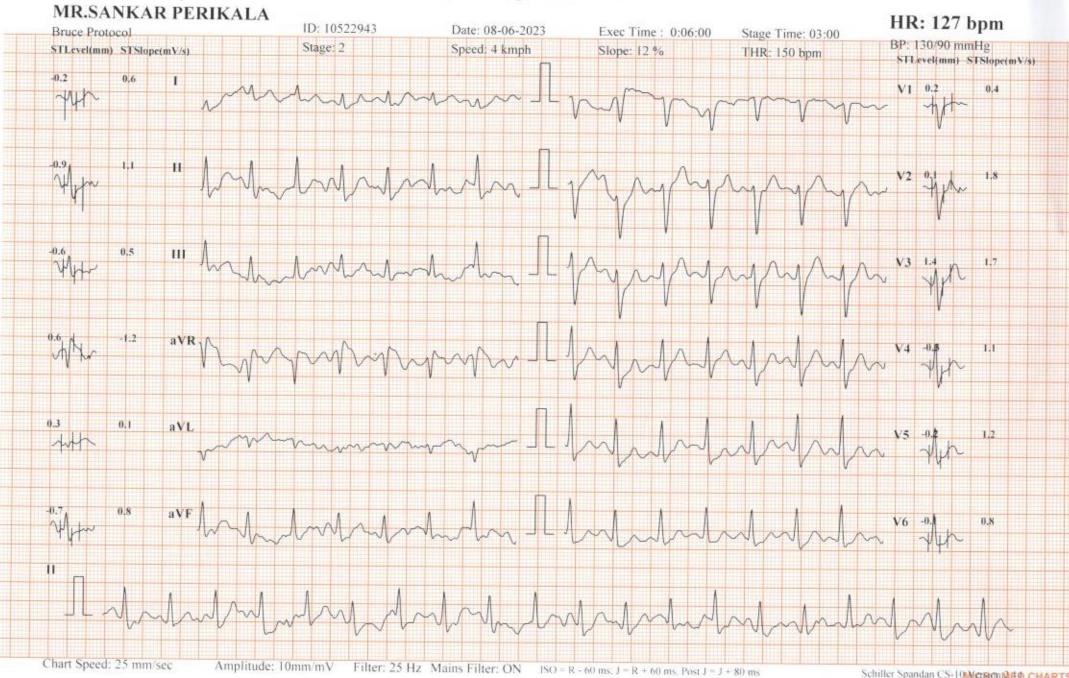
ledica Fest rotoco xercis lax BI	History: HT tions: YES Details: d: Bruce e Time:			Height:	60 ems		I Provide Laboration				
ledica Fest rotoco xercis lax BI	tions: YES Details: I: Bruce e Time:						Weight:	71 Kg	ID: 10522943		
Fest rotoco xercis 1ax BI	Details: bl: Bruce e Time:							9			
rotoco xercis lax BI	l: Bruce e Time:										
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xercis 1ax BI	e Time:			Deadland	M	1.77					
lax BI		0:09:11		Predicted			A CD P	1.5.0105	2.7	Target HR	: 150
F14-122.074).	140/90		Achieved Max HR: 145 (82% of Predicted MHR) Max BP x HR: 20300							
ost To	rmination Crit			Max BP x	HR: 205	00				Max Mets:	10.3
roto	col Details	Stage Name	Stage Time	METS	Speed	Conde	Union Data	Top	0.000	1000	Two as a second se
		17		11215	kmph	Grade	Heart Rate	8P mmttg	RPP	ST Level	ST Slope mV/S
		Supine	00:07	1	0	0	70	120/80	8400	1 V2	1.4 V3
		Standing	00.06	1	.0	0	79	120/80	9480	0.9 ¥3	1.4 V3
		HyperVentilation	00:06	1	0	0	64	120/80	7680	1.1.V2	14 V3
		PreTest	00.28	1	1.6	0	86	120/80	10320	1.6.111	1.2 V2
		Stage: 1	03-00	4:7	2.7	10	109	120/80	13080	1 V3	1.8 V3
		Stage 2	.03.00 ,	7	4	12	127	130/90	16510	14.V3	1.8 V2
		Stage: 3	03.00	10.1	5.5	14	145	140/90	20300	-1.7.11	2.2 VS
		Peak Exercise	00.11	10.3	6.8	16	145	140/90	20300	12.V2	2.7 V3
		Recovery I	01.00	1	0	0	115	140.90	16100	1.2 V2	2.5 V3
		Recovery2	01:00	1	0	0	102	130/90	13260	0.6 VI	1.8 V3
		Recovery3	01.00	1	0	0	.95	1.30/0	12350	0.7 V3	1.4 V3

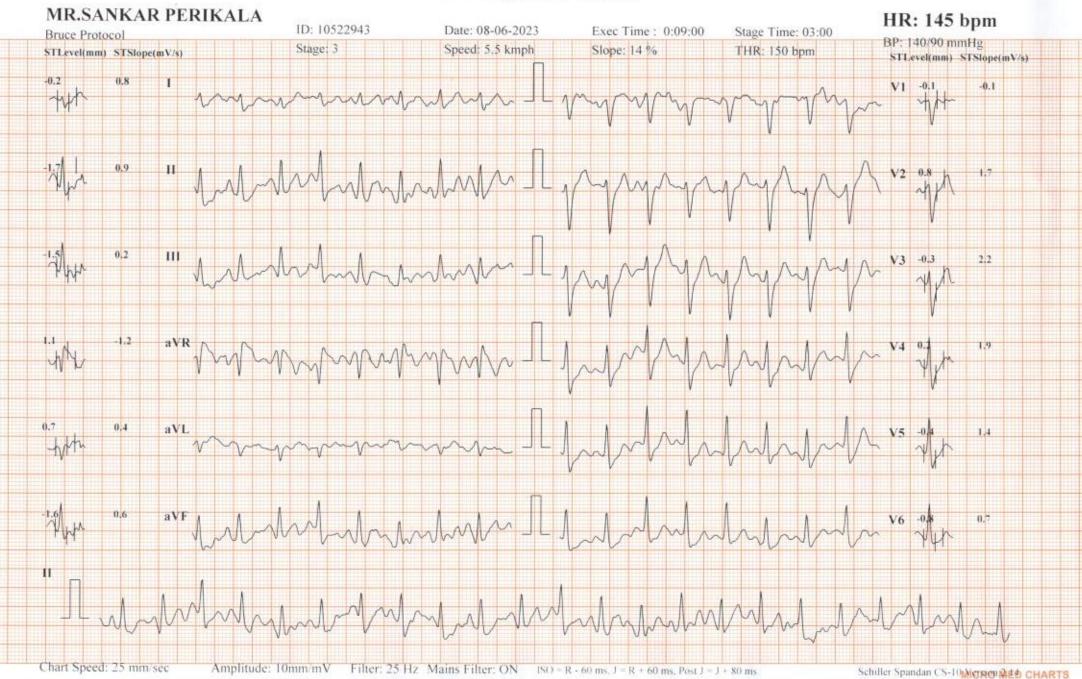


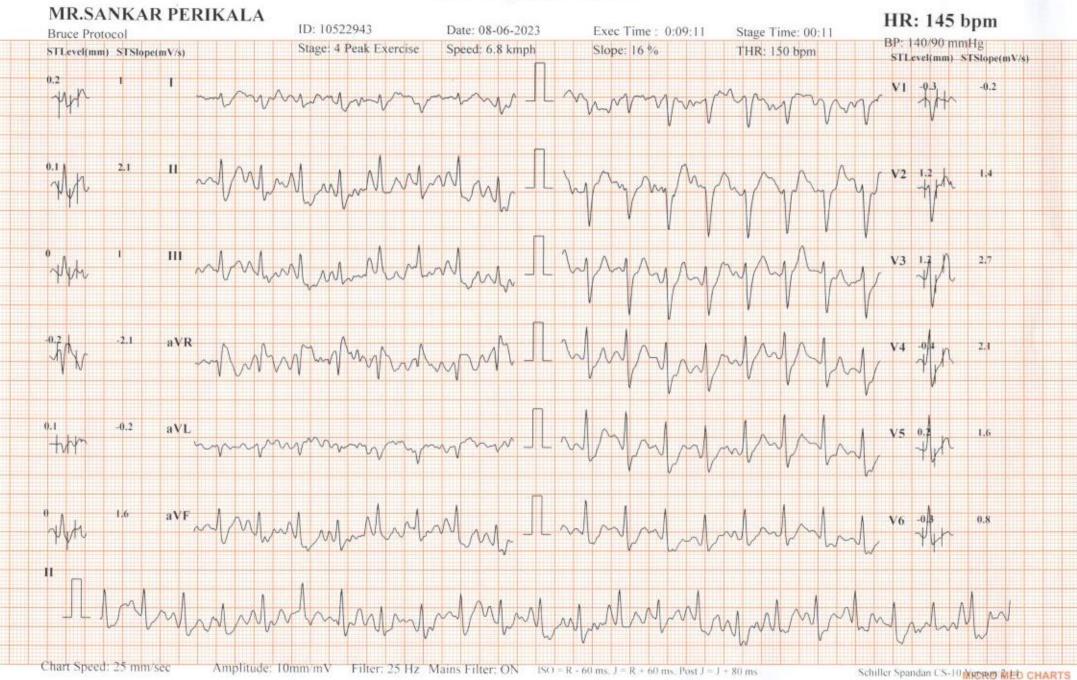


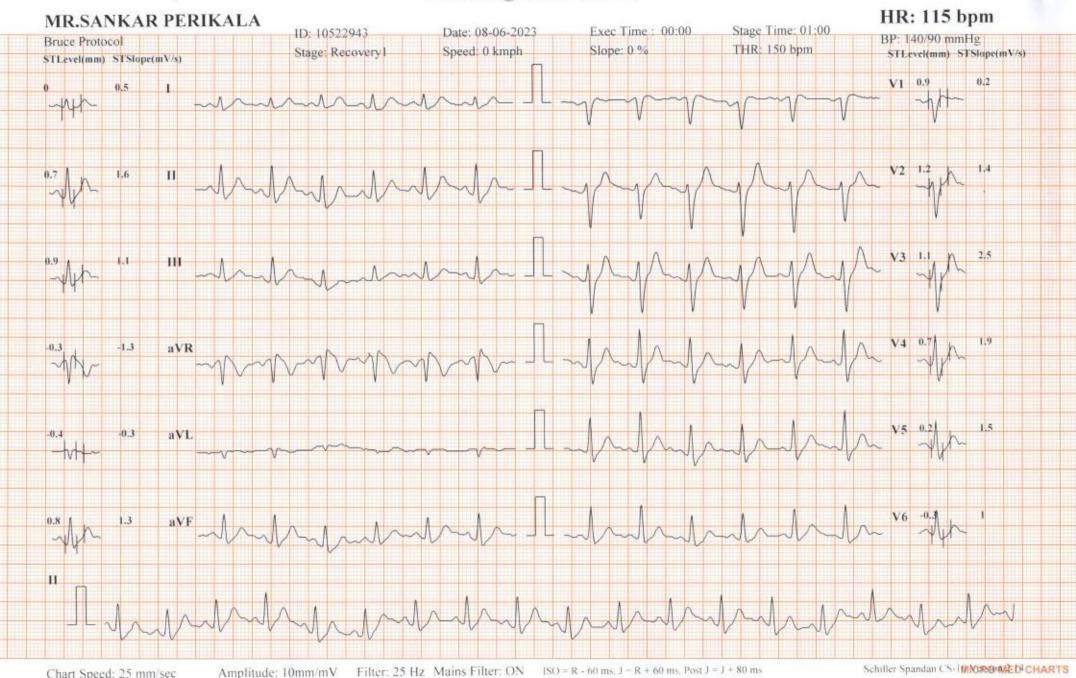


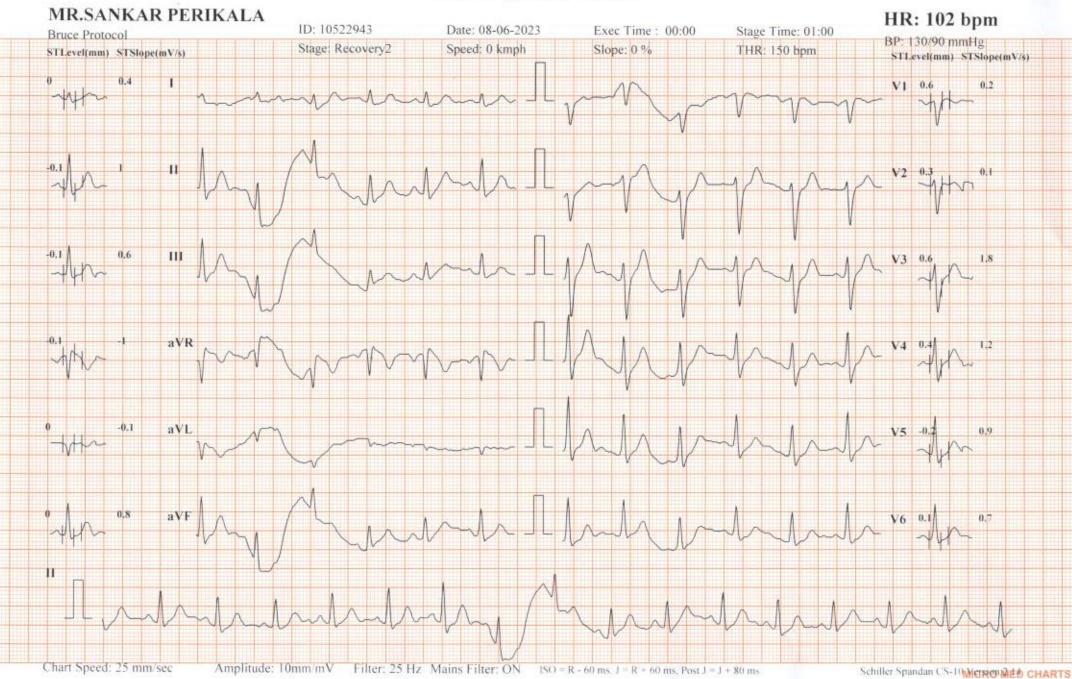


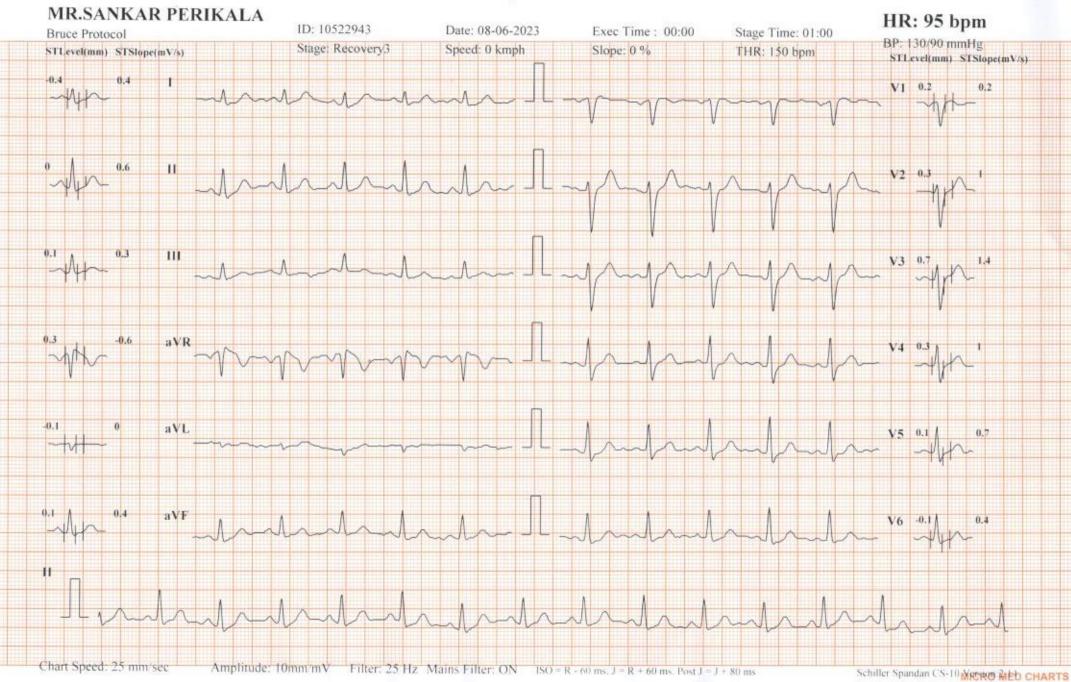














PERIKALA SANKAR 43Y MALE 10522943 CHEST PA 08-Jun-23 YODA DIAGNOSTICS