

TEST REPORT

Reg. No. : 403100837	Reg. Date : 23-Mar-2024 15:33	Ref.No :	Approved On : 23-Mar-2024 18:21
Name : Mr. KAMLESH SONAGRA			Collected On : 23-Mar-2024 16:39
Age : 31 Years	Gender: Male	Pass. No. :	Dispatch At :
Ref. By : APOLLO			Tele No. :
Location :			

Test Name	Results	Units	Bio. Ref. Interval
Complete Blood Count			
<u>Specimen: EDTA blood</u>			
Hemoglobin			
Hemoglobin(SLS method)	13.2	g/dL	13.0 - 17.0
Hematocrit (calculated)	40.4	%	40 - 50
RBC Count(Ele.Impedence)	4.82	X 10 ¹² /L	4.5 - 5.5
MCV (Calculated)	83.8	fL	83 - 101
MCH (Calculated)	27.4	pg	27 - 32
MCHC (Calculated)	32.7	g/dL	31.5 - 34.5
RDW (Calculated)	12.5	%	11.5 - 14.5
Differential WBC count (Impedance and flow)			
Total WBC count	H 10500	/μL	4000 - 10000
Neutrophils	61	%	38 - 70
Lymphocytes	30	%	21 - 49
Monocytes	06	%	3 - 11
Eosinophils	03	%	0 - 7
Basophils	00	%	0 - 1
Platelet			
Platelet Count (Ele.Impedence)	325000	/cmm	150000 - 410000
MPV	9.20	fL	6.5 - 12.0
Platelets appear on the smear	Adequate		
Malarial Parasites	Not Detected		
EDTA Whole Blood			

Note: All abnormal hemograms are reviewed and confirmed microscopically. Peripheral blood smear and malarial parasite examination are not part of CBC report.

Test done from collected sample.

This is an electronically authenticated report.



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Name : Mr. KAMLESH SONAGRA Collected On : 23-Mar-2024 16:39
Age : 31 Years Gender: Male Pass. No. : Dispatch At :
Ref. By : APOLLO Tele No. :
Location :

Test Name	Results	Units	Bio. Ref. Interval
ESR	05	mm/hr	17-50 Yrs : <12, 51-60 Yrs : <19, 61-70 Yrs : <20, >70 Yrs : <30

Method: Modified Westergren

EDTA Whole Blood

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TEST REPORT

Reg. No. : 403100837 Reg. Date : 23-Mar-2024 15:33 Ref.No : Approved On : 23-Mar-2024 18:12
Name : Mr. KAMLESH SONAGRA Collected On : 23-Mar-2024 16:39
Age : 31 Years Gender: Male Pass. No. : Dispatch At :
Ref. By : APOLLO Tele No. :
Location :

Test Name	Results	Units	Bio. Ref. Interval
BLOODGROUP & RH			
<u>Specimen: EDTA and Serum; Method: Gel card system</u>			
Blood Group "ABO" <i>Agglutination</i>	A		
Blood Group "Rh" <i>Agglutination</i>	Positive		
EDTA Whole Blood			

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TEST REPORT

Reg. No. : 403100837	Reg. Date : 23-Mar-2024 15:33	Ref.No :	Approved On : 24-Mar-2024 10:45
Name : Mr. KAMLESH SONAGRA			Collected On : 23-Mar-2024 16:39
Age : 31 Years	Gender: Male	Pass. No. :	Dispatch At :
Ref. By : APOLLO			Tele No. :
Location :			

Test Name	Results	Units	Bio. Ref. Interval
PERIPHERAL BLOOD SMEAR EXAMINATION			
<u>Specimen: Peripheral blood smear & EDTA blood, Method:Microscopy</u>			
RBC Morphology	RBCs are normocytic normochromic.		
WBC Morphology	Total WBC and differential count is within normal limit. No abnormal cells or blasts are seen.		
Differential Count	.		
Neutrophils	61	%	38 - 70
Lymphocytes	30	%	21 - 49
Monocytes	06	%	3 - 11
Eosinophils	03	%	0 - 7
Basophils	00	%	0 - 2
Platelets	Platelets are adequate with normal morphology.		
Parasite	Malarial parasite is not detected.		
Sample Type: EDTA Whole Blood			

Test done from collected sample.

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Name : Mr. KAMLESH SONAGRA			Collected On : 23-Mar-2024 16:39
Age : 31 Years	Gender : Male	Pass. No. :	Dispatch At :
Ref. By : APOLLO			Tele No. :
Location :			

Test Name	Results	Units	Bio. Ref. Interval
<u>FASTING PLASMA GLUCOSE</u>			
<u>Specimen: Fluoride plasma</u>			
Fasting Plasma Glucose <i>Hexokinase</i>	99.53	mg/dL	Normal: <=99.0 Prediabetes: 100-125 Diabetes :>=126

Flouride Plasma

Criteria for the diagnosis of diabetes:

1. HbA1c >= 6.5 *
- Or
2. Fasting plasma glucose >126 gm/dL. Fasting is defined as no caloric intake at least for 8 hrs.
- Or
3. Two hour plasma glucose >= 200mg/dL during an oral glucose tolerance test by using a glucose load containing equivalent of 75 gm anhydrous glucose dissolved in water.
- Or
4. In a patient with classic symptoms of hyperglycemia or hyperglycemic crisis, a random plasma glucose >= 200 mg/dL. *In the absence of unequivocal hyperglycemia, criteria 1-3 should be confirmed by repeat testing. American diabetes association. Standards of medical care in diabetes 2011. Diabetes care 2011;34:S11.

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TEST REPORT

Reg. No. : 403100837 Reg. Date : 23-Mar-2024 15:33 Ref.No : Approved On : 23-Mar-2024 20:07
Name : Mr. KAMLESH SONAGRA Collected On : 23-Mar-2024 16:39
Age : 31 Years Gender: Male Pass. No. : Dispatch At :
Ref. By : APOLLO Tele No. :
Location :

Test Name	Results	Units	Bio. Ref. Interval
POST PRANDIAL PLASMA GLUCOSE <u>Specimen: Fluoride plasma</u>			
Post Prandial Plasma Glucose <i>Hexokinase</i>	L 130.54	mg/dL	Normal: <=139 Prediabetes : 140-199 Diabetes: >=200
Flouride Plasma			

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TEST REPORT

Reg. No. : 403100837	Reg. Date : 23-Mar-2024 15:33	Ref.No :	Approved On : 23-Mar-2024 18:30
Name : Mr. KAMLESH SONAGRA			Collected On : 23-Mar-2024 16:39
Age : 31 Years	Gender: Male	Pass. No. :	Dispatch At :
Ref. By : APOLLO			Tele No. :
Location :			

Test Name	Results	Units	Bio. Ref. Interval
LIPID PROFILE			
CHOLESTEROL	186.00	mg/dL	Desirable <=200 Borderline high risk 200 - 240 High Risk >240
Triglyceride <i>Enzymatic Colorimetric Method</i>	92.00	mg/dL	<150 : Normal, 150-199 : Border Line High, 200-499 : High, >=500 : Very High
Very Low Density Lipoprotein(VLDL) <i>Calculated</i>	18	mg/dL	0 - 30
Low-Density Lipoprotein (LDL) <i>Calculated Method</i>	114.88	mg/dL	< 100 : Optimal, 100-129 : Near Optimal/above optimal, 130-159 : Borderline High, 160-189 : High, >=190 : Very High
High-Density Lipoprotein(HDL)	53.12	mg/dL	<40 >60
CHOL/HDL RATIO <i>Calculated</i>	3.50		0.0 - 3.5
LDL/HDL RATIO <i>Calculated</i>	2.16		1.0 - 3.4
TOTAL LIPID <i>Calculated</i>	516.00	mg/dL	400 - 1000
Serum			

As a routine test to determine if your cholesterol level is normal or falls into a borderline-, intermediate- or high-risk category.
 To monitor your cholesterol level if you had abnormal results on a previous test or if you have other risk factors for heart disease.
 To monitor your body's response to treatment, such as cholesterol medications or lifestyle changes.
 To help diagnose other medical conditions, such as liver disease.
 Note : biological reference intervals are according to the national cholesterol education program (NCEP) guidelines.

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Age : 31 Years	Gender : Male	Pass. No. :	Dispatch At :
Ref. By : APOLLO			Tele No. :
Location :			

Test Name	Results	Units	Bio. Ref. Interval
<u>LIVER FUNCTION TEST</u>			
TOTAL PROTEIN <small>Biuret Colorimetric</small>	7.6	g/dL	6.4 - 8.3
ALBUMIN <small>Bromocresol Green(BCG)</small>	4.8	g/dL	3.2 - 5.0
GLOBULIN <small>Calculated</small>	2.80	g/dL	2.4 - 3.5
ALB/GLB <small>Calculated</small>	1.71		1.2 - 2.2
SGOT <small>Pyridoxal 5 Phosphate Activation, IFCC</small>	24.5	U/L	0 - 40
SGPT <small>Pyridoxal 5 Phosphate Activation, Ifcc</small>	26.3	U/L	0 - 41
Alkaline Phosphatase <small>ENZYMATIC COLORIMETRIC IFCC, PNP, AMP BUFFER</small>	56.4	U/L	40 - 130
TOTAL BILIRUBIN <small>Diazo</small>	0.69	mg/dL	0.0 - 1.2
DIRECT BILIRUBIN <small>Diazo Reaction</small>	0.10	mg/dL	0 - 0.3
INDIRECT BILIRUBIN <small>Calculated</small>	0.59	mg/dL	0.0 - 1.00
Serum			

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Name : Mr. KAMLESH SONAGRA **Collected On** : 23-Mar-2024 16:39
Age : 31 Years **Gender**: Male **Pass. No.** : **Dispatch At** :
Ref. By : APOLLO **Tele No.** :
Location :

Test Name	Results	Units	Bio. Ref. Interval
HEMOGLOBIN A1C (HBA1C) <i>High Performance Liquid Chromatography (HPLC)</i>	5.60	%	Normal: ≤ 5.6 Prediabetes: 5.7-6.4 Diabetes: ≥ 6.5 6-7 : Near Normal Glycemia, <7 : Goal , 7-8 : Good Control ,>8 : Action Suggested.
Mean Blood Glucose <i>(Calculated)</i>	114	mg/dL	
Sample Type: EDTA Whole Blood			

Criteria for the diagnosis of diabetes

- HbA1c ≥ 6.5 * Or Fasting plasma glucose >126 gm/dL. Fasting is defined as no caloric intake at least for 8 hrs. Or
- Two hour plasma glucose ≥ 200 mg/dL during an oral glucose tolerance test by using a glucose load containing equivalent of 75 gm anhydrous glucose dissolved in water. Or
- In a patient with classic symptoms of hyperglycemia or hyperglycemic crisis, a random plasma glucose ≥ 200 mg/dL. *In the absence of unequivocal hyperglycemia, criteria 1-3 should be confirmed by repeat testing. American diabetes association. Standards of medical care in diabetes 2011. Diabetes care 2011:34:S11.

Limitation of HbA1c

- In patients with Hb variants even analytically correct results do not reflect the same level of glycemic control that would be expected in patients with normal population.
 - Any cause of shortened erythrocyte survival or decreased mean erythrocyte survival or decreased mean erythrocyte age eg. hemolytic diseases, pregnancy, significant recent/chronic blood loss etc. will reduce exposure of RBC to glucose with consequent decrease in HbA1c values.
 - Glycated HbF is not detected by this assay and hence specimens containing high HbF ($>10\%$) may result in lower HbA1c values than expected. Importance of HbA1C (Glycated Hb.) in Diabetes Mellitus
- HbA1C, also known as glycated hemoglobin, is the most important test for the assessment of long term blood glucose control(also called glycemic control).
 - HbA1C reflects mean glucose concentration over past 6-8 weeks and provides a much better indication of longterm glycemic control than blood glucose determination.
 - HbA1c is formed by non-enzymatic reaction between glucose and Hb. This reaction is irreversible and therefore remains unaffected by short term fluctuations in blood glucose levels.
 - Long term complications of diabetes such as retinopathy (Eye-complications), nephropathy (kidney-complications) and neuropathy (nerve complications), are potentially serious and can lead to blindness, kidney failure, etc.
 - Glycemic control monitored by HbA1c measurement using HPLC method (GOLD STANDARD) is considered most important. (Ref. National Glycohaemoglobin Standardization Program - NGSP)
- Note : Biological reference intervals are according to American Diabetes Association (ADA) Guidelines.

Test done from collected sample.

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TEST REPORT

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Name : Mr. KAMLESH SONAGRA			Collected On : 23-Mar-2024 16:39
Age : 31 Years	Gender: Male	Pass. No. :	Dispatch At :
Ref. By : APOLLO			Tele No. :
Location :			

Bio-Rad CDM System
Bio-Rad Variant V-II Instrument #1

PATIENT REPORT
V2TURBO_A1c_2.0

Patient Data

Sample ID: 140303500662
 Patient ID:
 Name:
 Physician:
 Sex:
 DOB:

Analysis Data

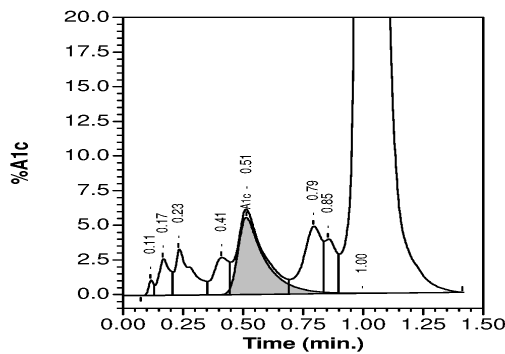
Analysis Performed: 23/03/2024 23:15:36
 Injection Number: 12752
 Run Number: 551
 Rack ID:
 Tube Number: 2
 Report Generated: 23/03/2024 23:18:02
 Operator ID:

Comments:

Peak Name	NGSP %	Area %	Retention Time (min)	Peak Area
Unknown	---	0.2	0.114	3196
A1a	---	0.9	0.165	16330
A1b	---	1.7	0.231	31251
LA1c	---	1.6	0.409	28817
A1c	5.6	---	0.513	85832
P3	---	2.9	0.792	52000
P4	---	1.4	0.853	25570
Ao	---	86.5	0.996	1560877

Total Area: 1,803,873

HbA1c (NGSP) = 5.6 %



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TEST REPORT

Reg. No. : 403100837 **Reg. Date** : 23-Mar-2024 15:33 **Ref.No** : **Approved On** : 23-Mar-2024 23:11
Name : Mr. KAMLESH SONAGRA **Collected On** : 23-Mar-2024 16:39
Age : 31 Years **Gender**: Male **Pass. No.** : **Dispatch At** :
Ref. By : APOLLO **Tele No.** :
Location :

Test Name	Results	Units	Bio. Ref. Interval
THYROID FUNCTION TEST			
T3 (triiodothyronine), Total <small>CMIA</small>	1.09	ng/mL	0.70 - 2.04
T4 (Thyroxine), Total <small>CMIA</small>	7.44	µg/dL	4.6 - 10.5
TSH (Thyroid stimulating hormone) <small>CMIA</small>	1.239	µIU/mL	0.35 - 4.94

Sample Type: Serum

Comments:

Thyroid stimulating hormone (TSH) is synthesized and secreted by the anterior pituitary in response to a negative feedback mechanism involving concentrations of FT3 (free T3) and FT4 (free T4). Additionally, the hypothalamic tripeptide, thyrotropin-releasing hormone (TRH), directly stimulates TSH production. TSH stimulates thyroid cell production and hypertrophy, also stimulate the thyroid gland to synthesize and secrete T3 and T4. Quantification of TSH is significant to differentiate primary (thyroid) from secondary (pituitary) and tertiary (hypothalamus) hypothyroidism. In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hypothyroidism, TSH levels are low.

TSH levels During Pregnancy :

- First Trimester : 0.1 to 2.5 µIU/mL
- Second Trimester : 0.2 to 3.0 µIU/mL
- Third trimester : 0.3 to 3.0 µIU/mL

Reference : Carl A.Burtis,Edward R.Ashwood,David E.Bruns. Tietz Textbook of Clinical Chemistry and Molecular Diagnostics. 5th Edition. Philadelphia: WB Saunders,2012:2170

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TEST REPORT

Reg. No. : 403100837	Reg. Date : 23-Mar-2024 15:33	Ref.No :	Approved On : 23-Mar-2024 18:36
Name : Mr. KAMLESH SONAGRA			Collected On : 23-Mar-2024 16:39
Age : 31 Years	Gender: Male	Pass. No. :	Dispatch At :
Ref. By : APOLLO			Tele No. :
Location :			

Test Name	Results	Units	Bio. Ref. Interval
<u>URINE ROUTINE EXAMINATION</u>			
<u>Physical Examination</u>			
Colour	Pale Yellow		
Clarity	Clear		
<u>CHEMICAL EXAMINATION (by strip test)</u>			
pH	6.0		4.6 - 8.0
Sp. Gravity	1.020		1.002 - 1.030
Protein	Nil		Absent
Glucose	Nil		Absent
Ketone	Nil		Absent
Bilirubin	Nil		Nil
Nitrite	Negative		Nil
Leucocytes	Nil		Nil
Blood	Absent		Absent
<u>MICROSCOPIC EXAMINATION</u>			
Leucocytes (Pus Cells)	1-2		0 - 5/hpf
Erythrocytes (RBC)	Nil		0 - 5/hpf
Casts	Nil	/hpf	Absent
Crystals	Nil		Absent
Epithelial Cells	Nil		Nil
Monilia	Nil		Nil
T. Vaginalis	Nil		Nil
Urine			

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Name : Mr. KAMLESH SONAGRA Collected On : 23-Mar-2024 16:39
Age : 31 Years Gender: Male Pass. No. : Dispatch At :
Ref. By : APOLLO Tele No. :
Location :

Test Name	Results	Units	Bio. Ref. Interval
Creatinine	0.96	mg/dL	0.67 - 1.5

Serum

Creatinine is the most common test to assess kidney function. Creatinine levels are converted to reflect kidney function by factoring in age and gender to produce the eGFR (estimated Glomerular Filtration Rate). As the kidney function diminishes, the creatinine level increases; the eGFR will decrease. Creatinine is formed from the metabolism of creatine and phosphocreatine, both of which are principally found in muscle. Thus the amount of creatinine produced is, in large part, dependent upon the individual's muscle mass and tends not to fluctuate much from day-to-day. Creatinine is not protein bound and is freely filtered by glomeruli. All of the filtered creatinine is excreted in the urine.

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TEST REPORT

Reg. No. : 403100837 Reg. Date : 23-Mar-2024 15:33 Ref.No : Approved On : 23-Mar-2024 18:43
Name : Mr. KAMLESH SONAGRA Collected On : 23-Mar-2024 16:39
Age : 31 Years Gender: Male Pass. No. : Dispatch At :
Ref. By : APOLLO Tele No. :
Location :

Test Name	Results	Units	Bio. Ref. Interval
Urea	21.5	mg/dL	<= 65 YEARS AGE: <50 mg/dL; >65 YEARS AGE: <71 mg/dL

UREASE/GLDH**Serum**

Useful screening test for evaluation of kidney function. Urea is the final degradation product of protein and amino acid metabolism. In protein catabolism, the proteins are broken down to amino acids and deaminated. The ammonia formed in this process is synthesized to urea in the liver. This is the most important catabolic pathway for eliminating excess nitrogen in the human body. Increased blood urea nitrogen (BUN) may be due to prerenal causes (cardiac decompensation, water depletion due to decreased intake and excessive loss, increased protein catabolism, and high protein diet), renal causes (acute glomerulonephritis, chronic nephritis, polycystic kidney disease, nephrosclerosis, and tubular necrosis), and postrenal causes (eg, all types of obstruction of the urinary tract, such as stones, enlarged prostate gland, tumors). The determination of serum BUN currently is the most widely used screening test for the evaluation of kidney function. The test is frequently requested along with the serum creatinine test since simultaneous determination of these 2 compounds appears to aid in the differential diagnosis of prerenal, renal and postrenal hyperuremia.

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TEST REPORT

Reg. No. : 403100837	Reg. Date : 23-Mar-2024 15:33	Ref.No :	Approved On : 23-Mar-2024 22:53
Name : Mr. KAMLESH SONAGRA			Collected On : 23-Mar-2024 16:39
Age : 31 Years	Gender: Male	Pass. No. :	Dispatch At :
Ref. By : APOLLO			Tele No. :
Location :			

Test Name	Results	Units	Bio. Ref. Interval
<u>ELECTROLYTES</u>			
Sodium (Na+) <small>Method:ISE</small>	144.00	mmol/L	136 - 145
Potassium (K+) <small>Method:ISE</small>	4.3	mmol/L	3.5 - 5.1
Chloride(Cl-) <small>Method:ISE</small>	104.00	mmol/L	98 - 107

Sample Type: Serum

Comments

The electrolyte panel is ordered to identify electrolyte, fluid, or pH imbalance. Electrolyte concentrations are evaluated to assist in investigating conditions that cause electrolyte imbalances such as dehydration, kidney disease, lung diseases, or heart conditions. Repeat testing of the electrolyte or its components may be used to monitor the patient's response to treatment of any condition that may be causing the electrolyte, fluid or pH imbalance.

----- End Of Report -----

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Approved On: 23-Mar-2024 22:53



NAME	SONAGARA KAMLESHKUMAR M.		
AGE/ SEX	31 yrs /M	DATE	23.03.2024
REF. BY	HEATH CHECKUP	DONE BY	Dr. Parth Thakkar Dr. Abhimanyu Kothari

2D ECHO CARDIOGRAPHY & COLOR DOPPLER STUDY

FINDINGS:-

- Normal LV systolic function, LVEF= 60%.
- No RWMA at rest.
- Normal LV Compliance.
- LV & LA are of normal size.
- RA & RV are of normal size.
- Intact IAS & IVS.
- All valves are structurally normal.
- Trivial MR, No AR, No PR.
- No TR, No PAH, RVSP=26 mmHg.
- No Clots or vegetation.
- No evidence of pericardial effusion..
- IVC is normal in size and preserved respiratory variation.



MEASUREMENTS:-

LVIDD	30 (mm)	LA	30 (mm)
LVIDS	20 (mm)	AO	21(mm)
LVEF	60%	AV cusp	
IVSD / LVPWD	10/10 (mm)	EPSS	

DOPPLER STUDY:-

Valve	Velocity (M/sec)	Max gradient (MmHg)	Mean gradient (Mm Hg)	Valve area Cm ²
Aortic	0.9	5		
Mitral	E:0.5 A:0.7			
Pulmonary	0.8	3.0		
Tricuspid	1.1	20		

CONCLUSION:-

- Normal LV systolic function, LVEF= 60%.
- No RWMA at rest.
- Normal LV Compliance.
- All valves are structurally normal.
- Trivial MR, No AR, No PR/PS.
- No TR, No PAH, RVSP=26 mmHg.
- Normal IVC,

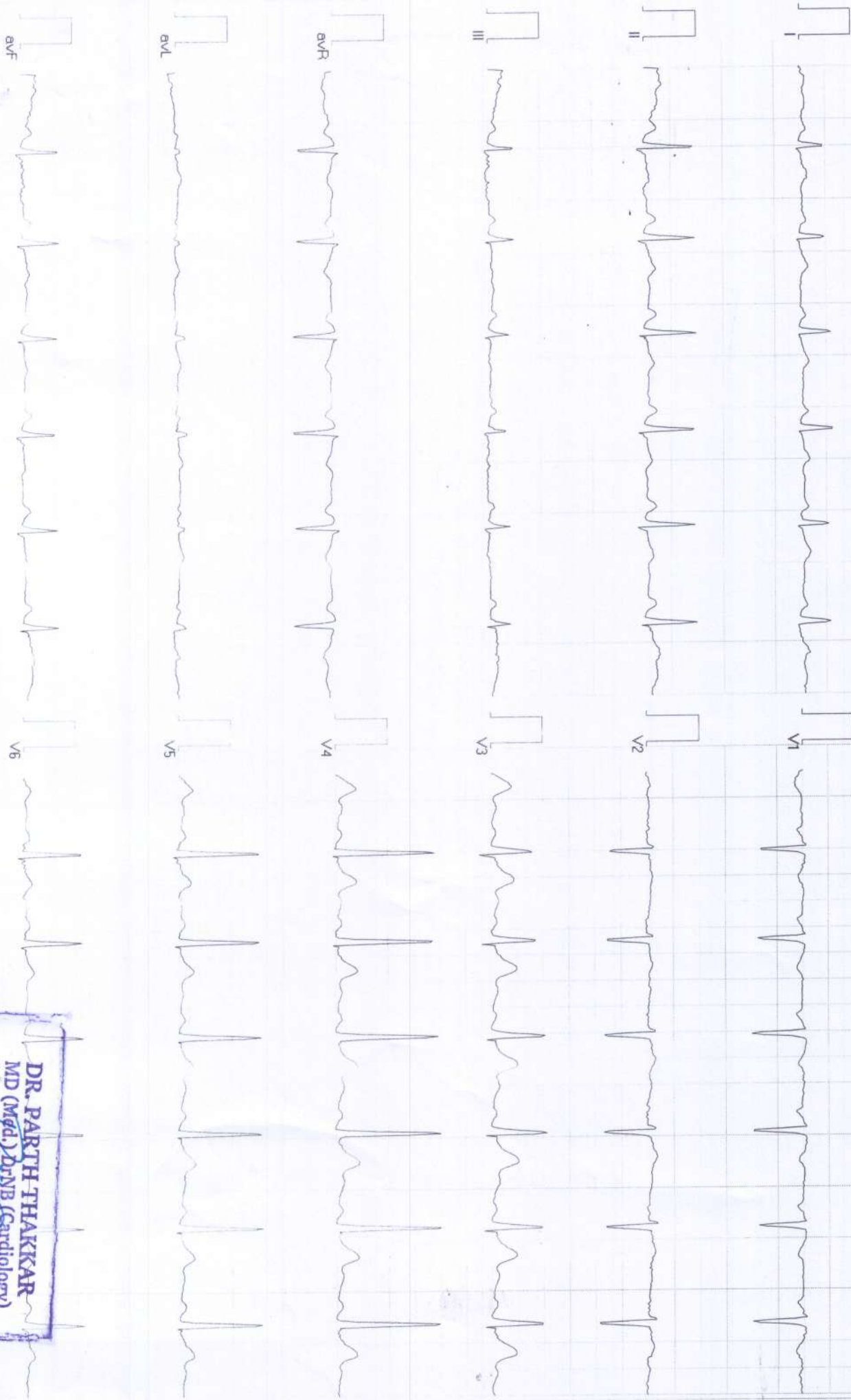
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 9714675115

CONCEPT DIAGNOSTIC

2022 / KAMLESHBHAI SONAGARA / 31 Yrs / M / Non Smoker
Heart Rate : 81 bpm / Tested On : 23-Mar-24 17:55:06 / HF 0.05 Hz - LF 35 Hz / Notch 50 Hz / Sn 1.00 Cm/mV / Sw 25 mm/s

ECG



Normal

Allergens ECG (Piscis)(PIS218210312)

Reported by
DR. PARTH THAKKAR
MD (Med.), DNB (Cardiology)
Interventional Cardiologist
G-32646



NAME :	KAMLESH SONAGARA	AGE/SEX:	31 Y/M
REF. BY:	HEALTH CHECK UP	DATE :	23-Mar-24

USG ABDOMEN & PELVIS

LIVER: normal in size & **shows increased echogenicity**. No evidence of dilated IHBR. No evidence of focal or diffuse lesion. CBD & Portal vein appears normal.

GALL-BLADDER: normal, No evidence of Gall Bladder calculi.

PANCREAS: appears normal in size & echotexture, No evidence of peri-pancreatic fluid collection.

SPLEEN: normal in size & shows normal echogenicity.

KIDNEYS: Both kidneys appear normal in size & echotexture.
Right kidney measures 98 x 47 mm. Left kidney measures 107 x 56 mm.
No evidence of calculus or hydronephrosis on either side.

URINARY BLADDER: appears normal and shows normal distension & normal wall thickness. No evidence of calculus or mass lesion.

PROSTATE: normal in size & echotexture.

No evidence of Ascites. No evidence of significant lymphadenopathy.

USG WITH HIGH FREQUENCY SOFT TISSUE PROBE:

Visualized bowel loops appears normal in caliber. No evidence of focal or diffuse wall thickening. No collection in RIF. Gaseous distension of the large bowel loops noted.

IMPRESSION:

- **Grade-I fatty liver.**

Dr. TEJAS PATEL
DNB RADIODIAGNOSIS

Dr. Tejas Patel
Diplomate N. B.
G-33659



NAME :	KAMLESHKUMAR SONAGARA	DATE :	23/03/2024
AGE/SEX:	31Y/M	REG.NO :	00
REFERRED BY: HEALTH CHECK UP			

X-RAY CHEST PA VIEW

- Both lung fields are clear.
- No evidence of consolidation or Koch's lesion seen.
- Heart size is within normal limit.
- Both CP angles are clear.
- Both dome of diaphragm appear normal.
- Bony thorax under vision appears normal.

Dr. Vidhi Shah
M.D. Radiologist
G - 41469

Dr. VIDHI SHAH
MD RADIODIAGNOSIS

