

LETTER OF APPROVAL / RECOMMENDATION

To,

The Coordinator, Mediwheel (Arcofemi Healthcare Limited) Helpline number: 011- 41195959

Dear Sir / Madam,

Sub: Annual Health Checkup for the employees of Bank of Baroda

This is to inform you that the following employee wishes to avail the facility of Cashless Annual Health Checkup provided by you in terms of our agreement.

PARTICULARS	EMPLOYEE DETAILS
NAME	MR. NUNSAVATHU PRAVEENKUMAR NAIK
EC NO.	107279
DESIGNATION	SMS CREDIT PROCESSING
PLACE OF WORK	VIJAYAWADA,RO VIJAYAWADA
BIRTHDATE	23-01-1990
PROPOSED DATE OF HEALTH CHECKUP	27-08-2022
BOOKING REFERENCE NO.	22S107279100023682E

This letter of approval / recommendation is valid if submitted along with copy of the Bank of Baroda employee id card. This approval is valid from 17-08-2022 till 31-03-2023 The list of medical tests to be conducted is provided in the annexure to this letter. Please note that the said health checkup is a cashless facility as per our tie up arrangement. We request you to attend to the health checkup requirement of our employee and accord your top priority and best resources in this regard. The EC Number and the booking reference number as given in the above table shall be mentioned in the invoice, invariably.

We solicit your co-operation in this regard.

Yours faithfully,

Sd/-

Chief General Manager HRM Department Bank of Baroda

(Note: This is a computer generated letter. No Signature required. For any clarification, please contact Mediwheel (Arcofemi Healthcare Limited))



E-mail:info@anuhospitals.com, www.anuneuroandcardiac.com

Name: N.Praveen Kumar Naik

Age/Sex: 32 yrs/M

Ref. By: Dr.D.Pujitha MD,(GEN)

Date: 08.10.2022

<u>ULTRASONOGRAPHY OF ABDOMEN</u>

LIVER:

12.7 cm Normal in size and texture.

No focal lesions noted. No intra-hepatic biliary dilatation.

PORTAL VEIN:

Normal in calibre.

GALLBLADDER:

Distended. Wall thickness is normal.

No calculi / peri cholecystic fluid collection.

CBD:

Normal in calibre.

PANCREAS:

Normal in size and texture.

No focal lesions / ductal dilatation / calcifications.

SPLEEN:

8.9 cm Normal in size and echotexture. No focal lesions.

RETROPERITONEUM: Aorta & IVC are normal in calibre.

No pre/para aortic lymphadenopathy. No obvious mass lesions at

adrenal region.

RIGHT KIDNEY:

11.3 x 4.3 cm Normal in size, position and texture. No focal lesions.

No calculi / hydronephrosis.

LEFT KIDNEY:

11.0 x 4.3 cm Normal in size, position and texture. No focal lesion

No calculi / hydronephrosis.

URINARY BLADDER: Distended, Mural thickness is normal, No Calculi,

PROSTATE

: Normal

CONCLUSION: NO ABNORMALITY NOTED.

SUGGEST CLINICAL CORRELATION.

UI. PAVAN KUMAR POLAGANI Dr Pavan Kumar Polagani MARES DNB 15 Consultant Radiologist 9 DIOLOGIST

^{*}No obvious pelvic pathology noted.

^{*}No free fluid noted in peritoneal cavity.



E-mail:info@anuhospitals.com, www.anuneuroandcardiac.com

2D - ECHO CARDIOGRAM & COLOUR DOPPLER REPORT

Patient's Name - N.PRAVEEN KUMAR

Age/Sex :-32Y/M Date: 08-10-2022 OP No: 67936

M-MODE:

LV: 4.9 X 3.1 cms EF: 60 %

LA: 3.0

cms

AO: 2.9

cms

IVS: 1.2

cms

PW: 1.1

cms

B-MODE:

LV: NO RWMA

LA: NORMAL

Mitral Valve

: NORMAL

RA: NORMAL

Aortic Valve

: NORMAL

RV: NORMAL

Tricuspid Valve: ; NORMAL

AO: NORMAL

Pulmonary Valve: ; NORMAL

PA: NORMAL

IAS: Intact

IVS: Intact

PERICARDIUM: NO PE

Colour Flow: __ MR: TRIVIAL

AR: NO

TR:TRIVIAL

PAH: NO

DOPPLER:

MV Flow: A>E AV Flow: 1.5 M/s, PV Flow: 1.1 M/s, RVSP: 20 mmHg

IMPRESSION

NO RWMA NORMAL LV FUNCTION TRIVIAL MR, TR, NO PAH **GRADE I DIASTOLIC DYSFUNCTION**

NO VEGETATION/CLOT/PE

DR. S. Viswanatha Kartik MD, DM,

Dept. of Cardiology

Consultant Interventional Cardiologist.

Dr. N. Anil Kumar MD., DM, FESC. FSCAI

Dept. of Cardiology,

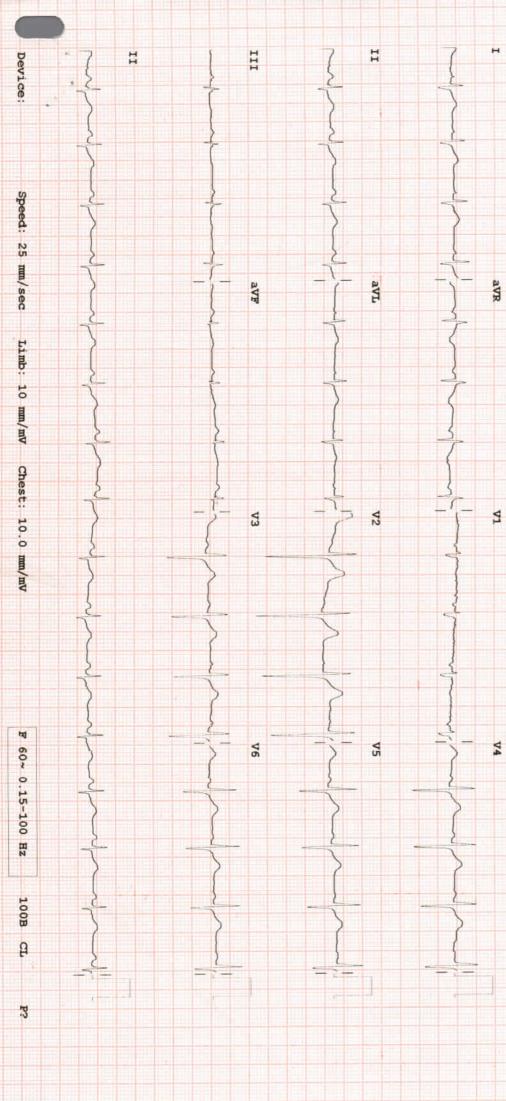
Consultant Interventional Cardiologist.

N. PRAVEEN KUMAR NAIK Male

10/8/2022 11:31:01 AM

T QRS PR QRSD QT QTc Rate --AXIS--132 71 321 404 81 21 95 . Sinus rhythm..... - NORMAL ECG -.....normal P axis, V-rate 50-99

12 Lead; Standard Placement aVR 7 Unconfirmed Diagnosis V4









Name

: Mr. N PRAVEEN KUMAR OP 67936

Visit No.

: V200010617

Age/Gender

: 32 Y/Male

Referred by

: Dr DR PUJITHA DUGGIRALA

External Visit ID

Patient No.

: P100008869

Registered On

: 08/10/2022 12:10

Collected On

: 08/10/2022 12:09

Reported On

0 - 15

: 08/10/2022 15:17

Final Report

HAEMATOLOGY

Test Name / Method

Results

Units

Reference Range

Sample Type

ERYTHROCYTE SEDIMENTATION

RATE-ESR Manual-Modified Westergren

10

mm/hr

Whole Blood

BLOOD GROUP & RH TYPING

method: Slide Agglutination/Reverse And Forward

"0" POSITIVE

Interpretation Notes:

*Suggested Gel card method for confirmation.

NOTE: ABO group should be reconfirmed after 6 months of age in newborn, as the ABO antibodies are weak or absent in sera until 3-6 months of age.

*** End Of Report

PROCESSED BY

: MOGHAL HAJAVALI

MEERJA RAF

M.Sc,M.Phil,DCR

Dr.MUSTHAQ AHMED

M.Sc, PHD

SREE VANI BADDIPUTI

MBBS, MD.

Reg.No: 66636

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	Final R	eport		
Test Name / Method	Results	Units	, Reference Range	Sample Type
	HAEMAT	OLOGY		
Complete Blood Count				
HAEMOGLOBIN Photometry- SLS Method	15.8	gms/dl	13.0- 18.0	Whole Blood
TOTAL COUNT/WBC Automated -Electrical Impedence/Manual	9710	cells/cumm	4000- 11000	
DIFFERENTIAL COUNT (DC) Automated -Flow Cytometry/Manual				
DIFFERENTIAL COUNT (DC)				
NEUTROPHILS	53.4	%	40-75	
YMPHOCYTES	41.6	%	20-40	
OSINOPHILS	2.1	%	0-6	
MONOCYTES	2.7	%	1-10	
BASOPHILS	0.2	%	0-1	ĵ.
RED BLOOD COUNT - RBC method :Electrical Impedance	5.48	million/cumm	4.5- 6	
PACKED CELL VOLUME- PCV method : Calculated	52.9	%	' 34- 48	
MEAN CORPUSCULAR VOLUME-MCV method : Calculated	96.6	<i>f</i> L	80- 96	
MEAN CORPUSCULAR HAEMGLOBIN- MCH method : Calculated	28.8	pg	27- 32	
MEAN CORPUSCULAR HAEMOGLOBIN CONCENTRATIONMCHC method: Calculated	29.8	gm/dl	30- 35	
RDW Automated-Electrical Impedence	14.1	%	11.0 - 16.0	
PLATELET COUNT Automated -Electrical Impedence	3.13	Lakhs/cmm	1.5 - 4.1	
				ï

··· End Of Report ···

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Reported On

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External Visit ID

Final Report

Test Name / Method	' Results	Units	Reference Range	Sample Type
	CLINICAL BIO	CHEMISTRY		
POST PRANDIAL BLOOD SUGAR method : Hexokinase	188	mg/dl	80-140	FLOURIDE PLASMA
GLYCOSYLATED HEMOGLOBIN (HbA1c) *method: Turbidimetric Inhibition Immunoassay			a	Whole Blood
GLYCOSYLATED HEMOGLOBIN (HbA1c)	5.8	%	= 5.6 % - Normal<br 5.7 - 6.4 % -Prediabe >/= 6.5 % - Diabetes	etes
Estimated Average Glucose(eAG) Interpretation Notes :	119	mg/dl	1 1	

- Estimated average Glucose (eAG) is calculated as per Diabetic Control & Complication Trial (DCCT) guidlines.
- HbA1c is used for monitoring diabetic control. It reflects the mean plasma glucose over three months.
- HbA1c may be falsely low in diabetics with haemolytic disease. In these individuals a plasma fructosamine level may be used which evaluates diabetes over 15 days.
- Abnormal hemoglobins might affect the RBC or glycation rates. In these cases even analytically correct results do not reflect the same level of glycemic control.
- Trends in HbA1c are a better indicator of diabetic control than a solitary test.
- Values have to be correlated with the clinical findings.

BLOOD UREA NITROGEN-BUN method : Derived	16	mg/dl	6 - 20	SERUM
SERUM CREATININE method : Jaffe Kinetic	0.8	mg/dl	0.7 - 1.2	

Interpretation Notes:

- Creatinine is a waste product largely from muscle breakdown. High values, especially with high BUN levels, may indicate problems with the kidneys. Increased levels observed in Acute or chronic renal failure; urinary tract obstruction, nephrotoxic drugs.
- creatinine is widely used as a test of renal (Kidney) function both as a general screen, along with urine protein, for renal disease, and as a test for monitoring of renal function in patients with renal disorder. Decreased creatinine are seen in reduced muscle mass, possible drug effect.
- Values have to be correlated with the clinical findings.

URIC ACID Method: Uricase-POD 6.5

mg/dl

3.4 - 7.0 .

" End Of Report "

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Reg.No: 66636







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Final Report

Test Name / Method	Results	Units	Reference Range	Sample Type
	CLINICAL BIO	CHEMISTRY		
THYROID PROFILE				
TRIIODO THYRONINE-T3 TOTAL Method: ECLIA	1.50	ng/ml	0.80 - 2.0	SERUM
THYROXINE -T4 TOTAL Method: ECLIA	8.50	ug/dl	5.1 - 14.1	1
THYROID STIMULATING HORMONE - TSH (Ultra Sensitive) Method : ECLIA	1.17	mIU/mI	• 0.40 - 4.20	

*** End Of Report ***

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Test Name / Method	Results	Units	Reference Range	Sample Type
	CLINICAL BIO	CHEMISTRY	1	
Lipid Profile				
CHOLESTEROL TOTAL Method: CHOD-POD	211	mg/dl	200-239: Borderline >240: Elevated <200: Normal	SERUM
TRIGLYCERIDES Method: GPO/PAP	171	mg/dl	<150: Normal 151-200: Borderline 201-499:High >500:Very High	
HDL CHOLESTEROL method : Direct	48	mg/dl	>55 NoRisk 35-55 Moderate Risk <35 High Risk	
LDL CHOLESTEROL method: Direct	129	mg/dl	<100: Optimal 101-129: Near/Above Optimal 130-159: Borderline 160-189: High >190: Very High	
VLDL CHOLESTEROL method : Calculated	34	mg/dl	7.0-40.0	
CHOL/HDL RATIO method : Calculated	4.3		0.0-4.5	
LDL/HDL RATIO method : Calculated	2.6		0.0-3.5	

*** End Of Report **

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Final Report

Test Name / Method	Results	Units	Reference Range	Sample Type
	CLINICAL BIO	CHEMISTRY		
Liver Function Test				
TOTAL BILIRUBIN method : Diazonium	1.10	mg/dl	0.0-1.2	SERUM
BILIRUBIN DIRECT method : Diazonium	0.50	mg/dl	0 - 0.3	
BILIRUBIN INDIRECT method : Calculated	0.60	mg/dl	0.0-1.0	
SGOT(AST) Without P5p	35	U/L	Upto 40	
SGPT(ALT) Without P5p	53	UIL	Upto 41	
ALKALINE PHOSPHATASE Method: AMP	116	IU/L	35 - 140	
GAMMA GT Method : Glupa C	48	U/L	8 - 61	
TOTAL PROTEIN method: Biuret	7.2	g/dl	6.4 - 8.7	
ALBUMIN Method : BCG	3.8	g/dl	3.5-5.2	
GLOBULIN method : Derived	3.4	gm/dl	2.5-3.8	
A/G RATIO method : Calculated	1,1		1.0-2.1	

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: 08/10/2022 12:09

: 32 Y/Male

Referred by

: Dr DR PUJITHA DUGGIRALA

Reported On

: 08/10/2022 19:32

External Visit ID

Final Report

	i iliai is			
Test Name / Method	Results	Units	Reference Range	Sample Type
	CLINICAL PA	ATHOLOGY		
URINE ROUTINE/ANALYSIS method: Macroscopic Examination				URINE
PHYSICAL EXAMINATION				
COLOUR Method: Macroscopic examination	YELLOW		Pale Yellow/Clear	
VOLUME Method: Macroscopic examination	15ml	ml	"	
APPEARANCE Method: Macroscopic examination	CLEAR		Clear	
CHEMICAL EXAMINATION				
pH Method: Reagent Strip Method (Double Indicator)	6.0		4.6-8.0	
SPECIFIC GRAVITY Method: Reagent Strip Method (Ion exchange)	1.020		1.005-1.030	
PROTEIN Method:Reagent Strip Method (Protien Error of indicator/SSA Test)	Nil		Nil	
GLUCOSE Method:Reagent Strip Method (GOD-POD/Benedict's Semiquantitative method)	Nil	%	Nil	
KETONES Method:Reagent Strip Method(Nitroprusside reaction/Dumn and Shipleys Reaction)	Negative		, Negative	
LEUCOCYTE ESTERASE	Absent		Absent	
UROBILINOGEN Method:Reagent Strip Method (Modified Ehrlich Reaction/Ehrlich Reagent)	Nil		<1.0 mg/dL	
BILIRUBIN Method:Reagent Strip Method (Diazonium Method/FOUCHET'S METHOD)	Negative		Negative	
MICROSCOPIC EXAMINATION				
RBCs Method:Microscopic Examination	Nil	/HPF	Nil	
EPITHELIAL CELLS Method:Microscopic Examination	1-2	/HPF	1-2/HPF	
BLOOD Method:Reagent Strip Method (Diazonium Method)	Negative		Negative	
PUS CELLS Method:Microscopic Examination	2-4	/HPF	Nil	

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NITRITES

Method:Reagent Strip Method (Diazonium

Method)

BACTERIA

Method:Microscopic Examination

CRYSTALS

Method:Microscopic Examination

Method:Microscopic Examination

OTHERS

Not Seen

Not seen

Not seen

Negative

Negative

: P100008869

: 08/10/2022 12:10

: 08/10/2022 12:09

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Not Seen

Not Seen

Not Seen

End Of Report

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TERMS & CONDITIONS OF REPORTING

- · It is presumed that the specimen belongs to the patient named or identified in the test request form.
- . The report results are for information and interpretation for your referring doctor and can be correlated with the patient's clinical history.
- Biological Reference Range/Interval is suggested for your Gender and Age on the basis of available literature. All reference ranges are to be reconsidered by doctor's advice for your specific care.
- . Test requested might not be performed for the following reasons:
 - a) Specimen quality insufficient (inadequate collections/spillage in transit)
 - b) Specimen quality unacceptable (haemolysed/clotted/lipemic etc.)
 - c) Incorrect specimen type.
 - d) Test cancelled either or request of patient or doctor, or because of incorrect test code, test name of specimen received. Reference may be provided to a new Accession number. Under "COMMENT" if the specimen has been re-accessioned for a different test. It is expected that a fresh specimen will be sent for the purpose of reporting on the same parameter (s), if required.
- This Medical Report is a professional opinion, not a diagnosis. Test results are not valid for medico legal purposes.
- . The report will carry the name and age provided at the time of registration. To maintain confidentiality, certain reports may not be e-mailed at the discretion of the management.
- All the notes and interpretation beneath the test result in the report provided are for educational purpose only. It is not intended to be a substitute for doctor's consultation.
- Reports that carries a 'PRELIMINARY' status signifies that results are yet to be reported for one or more of the test, or else as is the case with many microbiology tests, a "FINAL" culture, identification or drug susceptibility result might be pending. In such case, the descriptor "RESULTS" column and will be replaced by the test results whenever the latter are ready. The report will, when completed, acquire a "FINAL" status.
- Results of tests may vary from laboratory to laboratory and in some parameters from time to time for the same patients. Test results and reference range may also vary depending on the technology and methodology used. Laboratory test results may also vary depending on the age, sex, time of the day sample has been taken, diet, medication and limitation of modern technology.
- . In case of any unexpected or alarming test results, please contact us immediately for re-confirmation, further discussion, clarifications and rectifications, if needed only.
- In case of any discrepancy due to typing error, kindly get it rectified immediately. If the collection date was not stated in the Test Requisition Form, the same will not be printed on the report.
- The Lab or its employees/representatives assume any liability or responsibility for any loss or damage that may be incurred by any person as a result of interpreting the meaning of this report.
- In case of any issues or suggestions about your test results, please email us on lab@tridentdiagnostics.com
- Our liability is limited to the amount of investigations booked with us.
- . The courts (forums) at Bengaluru shall have exclusive jurisdiction in all disputes/claims concerning the tests and the results of the tests.

Page 11 of 11