

. 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 spouse of our employee wishes to avail the facility of Cashless Annual Health Checkup provided by you in terms of our agreement.

OF HEALTH CHECK UP BENEFICIARY
KOTA RAVI KIRAN KOTA RAVI KIRA
14-07-1990
22-10-2022
22D172896100027672S
SPOUSE DETAILS
MS. SRAVANI BANDI
172896
SINGLE WINDOW OPERATOR A
GANNAVARAM
18-06-1990

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 12-10-2022 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's spouse 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.

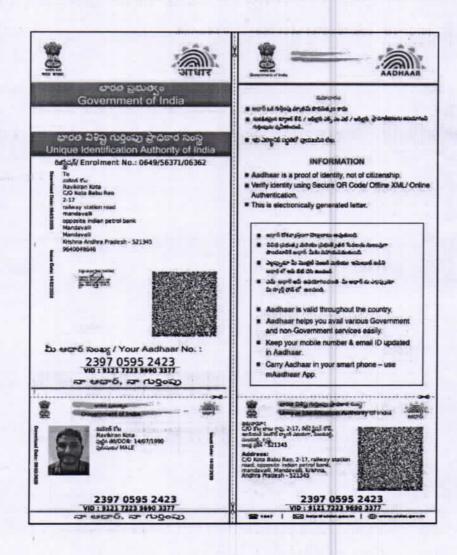
We solicit your co-operation in this regard.

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)).







#3-20/14, Main Road, Enikepadu, Vijayawada - 521108. Ph: 0866 - 2843133, 2843733, E-mail: info@anuhospitals.com www.anuneuroandcardiac.com

Name

: Mr. KOTA RAVI KIRAN OP MR 68759

Visit No.

: V200012080

Age/Gender

: 33 Y/Male

Referred by

: Dr DR SOUMYA MEDARAMETLA

External Visit ID

Patient No.

: P100009620

Registered On

: 22/10/2022 10:31

Collected On

: 22/10/2022 10:31

Reported On

: 22/10/2022 13:07

Final Report

	Final R	eport		O In Trans
Test Name / Method	Results	Units	Reference Range	Sample Type
Test Name / Method	HAEMATOLOGY			
ERYTHROCYTE SEDIMENTATION RATE-ESR Manual-Modified Westergren	. 15	mm/hr	0 - 15	Whole Blood

BLOOD GROUP & RH TYPING

"B"

method: Slide Agglutination/Reverse And Forward

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 RAFI

M.Sc,M.Phil,DCR

Dr.MUSTHAQ AHMED

M.Sc, PHD

SREE VANI BADDIPUTI

MBBS, MD.

Reg.No: 66636

Page 1 of 11

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NAME OF TAXABLE PARTY.	-	- 12	

	Final R	eport	The second secon	O 1 T
Test Name / Method	Results	Units	Reference Range	Sample Type
est Name / Method	HAEMAT	OLOGY		
Complete Blood Count HAEMOGLOBIN Photometry- SLS Method	15.1	gms/dl	13.0- 18.0	Whole Blood
TOTAL COUNT/WBC Automated -Electrical Impedence/Manual	8700	cells/cumm	4000- 11000	
DIFFERENTIAL COUNT (DC) Automated -Flow Cytometry/Manual				
DIFFERENTIAL COUNT (DC)		%	40-75	
NEUTROPHILS	47		20-40	
LYMPHOCYTES	40	%	0-6	
EOSINOPHILS	06	%	1-10	
MONOCYTES	07	%	0-1	
BASOPHILS	00	%		
RED BLOOD COUNT - RBC method :Electrical Impedance	6.23	million/cumm	4.5- 6	
PACKED CELL VOLUME- PCV method : Calculated	49.6	%	34- 48	
MEAN CORPUSCULAR VOLUME-MCV method : Calculated	79.6	fL	80- 96	
MEAN CORPUSCULAR HAEMGLOBIN-	24.2	pg	27- 32	
method: Calculated MEAN CORPUSCULAR HAEMOGLOBIN CONCENTRATIONMCHC method: Calculated	30.4	gm/dl	30- 35	
RDW Automated-Electrical Impedence	14.0	%	11.0 - 16.0	
PLATELET COUNT Automated -Electrical Impedence	3.13	Lakhs/cmm	1.5 - 4.1	

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: 22/10/2022 13:07 Reported On

> 5.7 - 6.4 % - Prediabetes >/= 6.5 % - Diabetes

Test Name / Method	Results	Units	Reference Range	Sample Type
restraine / mones	CLINICAL BIO	CHEMISTRY		
FASTING BLOOD SUGAR method : Hexokinase	89	mg/dl	Normal: 70 - 99 Pre-Diabetic : 100 - 125 Diabetic : >126	FLOURIDE PLASMA
FASTING URINE SUGAR method : Reagent Strip	NIL	%	Nil	URINE
POST PRANDIAL BLOOD SUGAR method : Hexokinase		mg/dl	80-140	FLOURIDE PLASMA
POST PRANDIAL URINE SUGAR method : Reagent Strip	15	%	Nil	URINE
GLYCOSYLATED HEMOGLOBIN (HbA1c) *method: Turbidimetric Inhibition Immunoassay	1			Whole Blood
GLYCOSYLATED HEMOGLOBIN (HbA1c)	5.9	%	= 5.6 % - Normal</td <td>etas</td>	etas

Final Report

Interpretation Notes:

- 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

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- 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.
- e to be correlated with the clinical findings.

GLYCOSYLATED HEMOGLOBIN (HbA1c)

Estimated Average Glucose(eAG)

BLOOD UREA NITROGEN-BUN method : Derived	19	mg/dl	6 - 20	SERUM

BUN/Creatinine Ratio

method: Calculated

mg/dl

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SERUM CREATININE

method : Jaffe Kinetic

0.97

mg/dl

0.7 - 1.2

SERUM

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

8.3

mg/dl

3.4 - 7.0

*** End Of Report

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	Final R	leport	-	O-male Tune
- IN / Mathod	Results	Units	Reference Range	Sample Type
Fest Name / Method	CLINICAL BIO	CHEMISTRY		
Lipid Profile CHOLESTEROL TOTAL Method: CHOD-POD	214	mg/dl	200-239: Borderline >240: Elevated <200: Normal	SERUM
TRIGLYCERIDES Method:GPO/POD	174	mg/dl	<150: Normal 151-200: Borderline 201-499:High >500:Very High	
HDL CHOLESTEROL Direct Method	43	mg/dl	>55 NoRisk 35-55 Moderate Risk <35 High Risk	
LDL CHOLESTEROL Direct Method	161	mg/dl	<100: Optimal 101-129: Near/Above Optimal 130-159: Borderline 160-189: High >190: Very High	
VLDL CHOLESTEROL method : Calculated	10	mg/dl	7.0-40.0	
CHOL/HDL RATIO method : Calculated	4.9		0.0-4.5	
LDL/HDL RATIO method : Calculated	3.7		0.0-3.5	

· Fnd Of Report ***

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	F	inal	Rep	0	r	ι
-		- COLUMN				

	Final R	eport	Dengo	Sample Type
	Results	Units	Reference Range	Ouripio 17F
Test Name / Method	CLINICAL BIO	CHEMISTRY		
THYROID PROFILE TRIIODO THYRONINE-T3 TOTAL	1.10	ng/ml	0.80 - 2.0	SERUM
Method : ECLIA	7.61	ug/dl	5.1 - 14.1	
THYROXINE -T4 TOTAL Method: ECLIA THYROID STIMULATING HORMONE -	2.10	mIU/mI	0.40 - 4.20	
THYROID STIMULATING HORANOTTEST (Ultra Sensitive) Method: ECLIA				

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	Final R		D. forence Pango	Sample Type
Mothod	Results	Units	Reference Range	Campio Typi
est Name / Method	CLINICAL BIO	CHEMISTRY		
Liver Function Test FOTAL BILIRUBIN method : Diazonium	0.55	mg/dl	0.0-1.2	SERUM
BILIRUBIN DIRECT	0.2	mg/dl	0 - 0.3	
method : Diazonium BILIRUBIN INDIRECT	0.35	mg/dl	0.0-1.0	
method : Calculated SGOT(AST)	29	U/L	Upto 40	
Without P5p	46	U/L	Upto 41	
SGPT(ALT) Without P5p ALKALINE PHOSPHATASE	72	IU/L	35 - 140	
Method : PNPP GAMMA GT	42	U/L	8 - 61	
Szasz Method TOTAL PROTEIN	7.9	g/dl	6.4 - 8.7	
method : Biuret	4.6	g/dl	3.5-5.2	
ALBUMIN Method : BCG	3.3	gm/dl	2.5-3.8	
GLOBULIN method: Derived A/G RATIO method: Calculated	1.3		1.0-2.1	

··· End Of Report ···

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	Final R	teport	D former Bange	Sample Type
Malhad	Results	Units	Reference Range	Odinpio 13F
Test Name / Method	CLINICAL PA	ATHOLOGY		STOOL
STOOL ROUTINE method: Manual COLOUR Method: Macroscopic Examination CONSISTENCY Method: Macroscopic Examination MUCUS Method: Macroscopic Examination REACTION Method: Macroscopic Examination PUS CELLS Method: Microscopic Examination RBCs Method: Microscopic Examination RBCs Method: Microscopic Examination EPITHELIAL CELLS Method: Microscopic Examination OVA Method: Microscopic Examination CYSTS Method: Microscopic Examination OTHER FINDING Method: Microscopic Examination		Cells/hpf Cells/hpf Cells/hpf	Brownish Semi Solid Absent Alkaline 0 - 5 Nil Absent Absent Absent	URINE

URINE ROUTINE/ANALYSIS

method: Macroscopic Examination

method : Macroscopic Examination		
PHYSICAL EXAMINATION	PALE YRLLOW	Pale Yellow/Clear
COLOUR Method: Macroscopic examination		ml -
VOLUME Method: Macroscopic examination	SEMI TURBID	Clear
APPEARANCE Method: Macroscopic examination SPECIFIC GRAVITY	1.025	1.005-1.030
Method: Reagent Strip Method (Ion exchange)	(B)	
CHEMICAL EXAMINATION pH Method: Reagent Strip Method (Double	6.0	4.6-8.0
Indicator) PROTEIN Method:Reagent Strip Method	TRACE	Negative
(Protein Error of Indicator/35A 1637)	NIL	% Negative
GLUCOSE Method:Reagent Strip Method (GOD-POD/Benedict's Semiquantitative method) KETONES Method:Reagent Strip Method (Sodium Nitroprusside Test)	NEGATIVE	Negative
	Page 9 of 11	

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LEUCOCYTE ESTERASE

Method:Reagent Strip Method (Diazonium Method/FOUCHET'S METHOD)

BLOOD Method:Reagent Strip Method

(Peroxidase - Like Activity)

Method:Reagent Strip Method (Diazonium

Method)

Method:Microscopic Examination

PUS CELLS

Method:Microscopic Examination

BACTERIA

Method:Microscopic Examination

CRYSTALS

Method:Microscopic Examination

OTHERS

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UROBILINOGEN

Method:Reagent Strip Method (Modified Ehrlich Reaction/Ehrlich Reagent)

NITRITES

MICROSCOPIC EXAMINATION

RBCs

Method:Microscopic Examination

EPITHELIAL CELLS

Method:Microscopic Examination

NEGATIVE

NIL

Negative

<1.0 mg/dL

NEGATIVE

Negative

NEGATIVE

Negative Negative

NEGATIVE

0-2

0-5

/HPF NIL

/HPF

/HPF 2-4

0-3

NOT SEEN

1-2

Not Seen

NOT SEEN

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:
 - Specimen quality insufficient (inadequate collections/spillage in transit) a)
 - Specimen quality unacceptable (haemolysed/clotted/lipemic etc.) b)

 - 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 c) 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
- 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 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.

P?



2D - ECHO CARDIOGRAM & COLOUR DOPPLER REPORT

Patient's Name - K.RAVI KIRAN , Age/Sex :- 33Y/M Date: 22-10-2022 OP No: 68759

M-MODE:

LV: 4.5 X 2.9 cms EF: 62 %

FS: 31%

LA: 2.7

cms

AO: 2.6

cms

IVS: 1.2

cms

PW: 1.0

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: 1.3 M/s, PV Flow: 1.2M/s, RVSP: 22 mmHg IMPRESSION

NO RWMA

NORMAL LV FUNCTION

TRIVIAL MR, TRIVIAL TR, NO PAH

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.

Age/Sex: 33 yrs/M Name: K. Ravi Kiran

Date: 22.10.2022 Ref. By: Dr. BHANU PRAVEEN NAIDU MRCSP(UK)

ULTRASONOGRAPHY OF ABDOMEN

Normal in size and increased echotexture. LIVER:

Focal hypoechoic area measuring approximately 1.2 cm seen

adjacent in GB fossa-likely focal fatty sparing.

PORTAL VEIN: Normal in calibre.

Distended. Wall thickness is normal. GALLBLADDER:

No calculi / peri cholecystic fluid collection.

Normal in calibre. CBD:

Normal in size and texture. PANCREAS:

No focal lesions / ductal dilatation / calcifications.

Normal in size and echotexture. No focal lesions. SPLEEN:

RETROPERITONEUM: Aorta & IVC are normal in calibre.

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

adrenal region.

Normal in size, position and texture. No focal lesions. RIGHT KIDNEY:

No calculi / hydronephrosis.

LEFT KIDNEY: Normal in size, position and texture. No focal lesions.

No calculi / hydronephrosis.

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

PROSTATE : Normal

*No obvious pelvic pathology noted.

*No free fluid noted in peritoneal cavity.

CONCLUSION:

GRADE I FATTY LIVER.

GF FOCAL FATTY SPARING AREA IN LIVER.

SUGGEST CLINICAL CORRELATION.

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