

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.

Attitual Floatist States	EMPLOYEE DETAILS
PARTICULARS	
	MS. SRAVANI BANDI
NAME	172896
EC NO.	SINGLE WINDOW OPERATOR A
DESIGNATION	GANNAVARAM
PLACE OF WORK	
BIRTHDATE	18-06-1990
PROPOSED DATE OF HEALTH	22-10-2022
CHECKUP BOOKING REFERENCE NO.	22D172896100027670E

This letter of approval / recommendation is valid if submitted along with copy of the Bank of Baroda employee in 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 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 Healuncare Limited))



नाम | बंडी श्रावणी

Name: Bandi Sravani धारक के हस्ताक्षर | Signature of Holder

B. Stavani

्राधिकारी जाशिकारी

कूट क. E.C. No.: 172896



परचान पत्र रहीते, चोदी या सरस्य होने पर धारक इसके लिए जिम्मेदार होगा.

पहचान पत्र सोने की मुचनी गुरना पुलिस दर्व मुल कार्यासय वा नैक की नजरीकी शासा को दें.

filled av - Fruffillen od søere आंधनिक मुख्या अधिकारी \$2000 - 400 024.

THE THE Blood Group : B + VA

Holder will be held responsible against loss, theft or damage.

police and parent/nearest bank office.

Zonal Security Officer onterior नुश्चा आधाराम् Zonal Security Officer रैंक आँक वर्गाटा, आंपनिक कार्यालय tiank of flareda - Zonal Office जो. नं , ३-६-२८६, कारीम पंत्रिक, हेटदगुडा, Door No.3-6-285, Kareen Montil, Hyderguda, Hyderabou - 500 029.

सम्पर्क टेलिफोन नं. Phone 940-23287213 (0)

Identification Marks : A Mole On The Left Hand Right Finger
A Mole on the Left Cheek

To
Bandi Sravani
ಐಂಪ ੍ਰਾਡਜ਼ੋਰੇ
D/O Anjaneyulu
514M
KALTEX ROAD
RAILWAY COLONY
RAILWAY STATION ROAD
Vijayawada (Urban)
Vijayawada,Krishna
Andhra Pradesh - 520001
9052465653



UF003874232IN

387423



మీ ఆధార్ సంఖ్య / Your Aadhaar No. :

8858 1848 5524

ఆధార్ - సామాన్యుని హక్కు



భారత ప్రభుత్వం GOVERNMENT OF INDIA



ಬಂಪಿ (ತಾವಣಿ Bandi Sravani

పుట్టిన సంచర్భకం/Year of Birth : 1990 ప్రై / Female

8858 1848 5524



5 ఆధార్ - సామాన్యుని హక్కు







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

Name

: Mrs. SRAVANI BANDI OP MR 68758

Visit No.

: V200012079

Age/Gender

: 32 Y/Female

Referred by

: Dr DR SOUMYA MEDARAMETLA

External Visit ID

Patient No.

: P100009619

Registered On

: 22/10/2022 10:30

Collected On

: 22/10/2022 10:29

Reported On

: 22/10/2022 13:07

Final Report

	7 77 7 77 7	The state of the s		
Test Name / Method	Results	Units	Reference Range	Sample Type
	HAEMAT	TOLOGY		
ERYTHROCYTE SEDIMENTATION	10	mm/hr	0 - 20	Whole Blood

RATE-ESR
Manual-Modified Westergren

BLOOD GROUP & RH TYPING

method : Slide Agglutination/Reverse And Forward

"B" 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

VERIFIED BY: G SURESH

MEERJA RAFI

M.Sc,M.Phil,DCR

Dr.MUSTHAQ AHMED

M.Sc, PHD

ELIVA IVALI

SREE VANI BADDIPUTI

MBBS, MD.

Reg.No: 66636

Page 1 of 11

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Test Name / Method	Results	Units	Reference Range	Sample Type
	HAEMAT	OLOGY		
Complete Blood Count				
HAEMOGLOBIN Photometry- SLS Method	12.4	gms/dl	12.0- 15.0	Whole Blood
OTAL COUNT/WBC Automated -Electrical Impedence/Manual	9700	cells/cumm	4000- 11000	
Automated -Flow Cytometry/Manual	U		11	
IFFERENTIAL COUNT (DC)				
IEUTROPHILS	69	%	40-75	
YMPHOCYTES	24	%	20-40	
OSINOPHILS	03	%	0-6	
MONOCYTES	04	%	1-10	
ASOPHILS	00	%	0-1	
RED BLOOD COUNT - RBC method :Electrical Impedance	4.88	million/cumm	4- 5.5	
ACKED CELL VOLUME- PCV method : Calculated	39.8	%	34- 48	
MEAN CORPUSCULAR VOLUME-MCV method : Calculated	81.6	fL	80- 96	
MEAN CORPUSCULAR HAEMGLOBIN- MCH method: Calculated	25.5	pg	27- 32	
MEAN CORPUSCULAR HAEMOGLOBIN CONCENTRATIONMCHC method: Calculated	31.2	gm/dl	30- 35	
RDW Automated-Electrical Impedence	12.7	%	11.0 - 16.0	
PLATELET COUNT Automated -Electrical Impedence	2.20	Lakhs/cmm	1.5 - 4.1	
	··· End Of	Report ***		
PROCESSED BY MOGHAL HAJAVALI				
VERIFIER RV. A RURERU				

VERIFIED BY : G SURESH

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MEDGENOME



TRIDENT DIAGNOSTICS

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	Final F	Report		
Test Name / Method	Results	Units	Reference Range	Sample Type
	CLINICAL BIO	CHEMISTRY		
FASTING BLOOD SUGAR method : Hexokinase	99	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	4	%	Nil	URINE
GLYCOSYLATED HEMOGLOBIN (HbA1c) *method : Turbidimetric Inhibition Immunoassay				Whole Blood
GLYCOSYLATED HEMOGLOBIN (HbA1c)	4.9	%	= 5.6 % - Normal<br 5.7 - 6.4 % -Prediab >/= 6.5 % - Diabetes	
Estimated Average Glucose(eAG)	94	mg/dl		

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 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

mg/dl

6 - 20

SERUM

BUN/Creatinine Ratio

method : Calculated

%

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SERUM CREATININE

method : Jaffe Kinetic

0.76

mg/dl

0.5 - 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

3.0

mg/dl

2.4 - 5.7

" End Of Report "

PROCESSED BY

: MOGHAL HAJAVALI

VERIFIED BY : G SURESH

........

Dr.MUSTHAQ AHMED

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	I IIIdi I	topoit		
Test Name / Method	Results	Units	Reference Range	Sample Type
	CLINICAL BIO	CHEMISTRY		
Lipid Profile				
CHOLESTEROL TOTAL Method: CHOD-POD	168	mg/dl	200-239: Borderline >240: Elevated <200: Normal	SERUM
TRIGLYCERIDES Method: GPO/POD	82	mg/dl	<150: Normal 151-200: Borderline 201-499:High >500:Very High	
HDL CHOLESTEROL Direct Method	37	mg/dl	>55 NoRisk 35-55 Moderate Risk <35 High Risk	
LDL CHOLESTEROL Direct Method	119	mg/dl	<100: Optimal 101-129: Near/Above Optimal 130-159: Borderline 160-189: High >190: Very High	
VLDL CHOLESTEROL method : Calculated	12	mg/dl	7.0-40.0	
CHOL/HDL RATIO method : Calculated	4.5		0.0-4.5	
LDL/HDL RATIO method : Calculated	3.2		0.0-3.5	

*** End Of Report ***

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: MOGHAL HAJAVALI

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Test Name / Method	Results	Units	Reference Range	Sample Type
	CLINICAL BIO	CHEMISTRY		
THYROID PROFILE				
TRIIODO THYRONINE-T3 TOTAL Method: ECLIA	1.16	ng/ml	0.80 - 2.0	SERUM
THYROXINE -T4 TOTAL Method: ECLIA	12.28	ug/dl	5.1 - 14.1	
THYROID STIMULATING HORMONE - TSH (Ultra Sensitive) Method: ECLIA	2.39	mIU/mI	0.40 - 4.20 First Trimester:0.1-2.5* Second & Third Trimester:0.2-3.0* *American Thyroid Association trimesterspecific	

*** End Of Report ***

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	rinain	epoit		
Test Name / Method	Results	Units	Reference Range	Sample Type
	CLINICAL BIO	CHEMISTRY		
Liver Function Test				
TOTAL BILIRUBIN method : Diazonium	0.9	mg/dl	0.0-1.2	SERUM
BILIRUBIN DIRECT method : Diazonium	0.32	mg/dl	0 - 0.3	
BILIRUBIN INDIRECT method : Calculated	0.58	mg/dl	0.0-1.0	
SGOT(AST) Without P5p	17	U/L	Upto 32	
GGPT(ALT) Without P5p	15	U/L	Upto 33	
ALKALINE PHOSPHATASE Method: PNPP	62	IU/L	35 - 140	
SAMMA GT Szasz Method	11	U/L	5 - 36	
TOTAL PROTEIN method: Biuret	7.5	g/dl	6.4 - 8.7	
ALBUMIN Method : BCG	4.4	g/dl	3.5-5.2	
GLOBULIN method : Derived	3.1	gm/dl	2.5-3.8	
A/G RATIO method : Calculated	1.4		1.0-2.1	

*** End Of Report ***

PROCESSED BY

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est Name / Method	Results	Units	Reference Range	Sample Type
	CLINICAL PA	THOLOGY		
TOOL ROUTINE method : Manual				STOOL
OLOUR Method:Macroscopic Examination			Brownish	
CONSISTENCY Method:Macroscopic xamination	A.		Semi Solid	
IUCUS Method:Macroscopic Examination			Absent	
EACTION Method:Macroscopic Examination			Alkaline	
US CELLS Method:Microscopic Examination		Cells/hpf	0 - 5	
BCs Method:Microscopic Examination		Cells/hpf	Nil	
PITHELIAL CELLS Method:Microscopic xamination		Cells/hpf	Absent	
VA Method:Microscopic Examination	. // m.		Absent	
YSTS Method:Microscopic Examination			Absent	
THER FINDING Method:Microscopic xamination	131			
RINE ROUTINE/ANALYSIS method : Macroscopic Examination HYSICAL EXAMINATION				URINE
OLOUR Macroscopic examination	PALE YELLOW		Pale Yellow/Clear	
OLUME lethod: Macroscopic examination	20	ml	-	
PPEARANCE lethod: Macroscopic examination	SEMI TURBID		Clear	
PECIFIC GRAVITY lethod: Reagent Strip Method (Ion exchange)	1.020		1.005-1.030	
HEMICAL EXAMINATION				
H Method: Reagent Strip Method (Double idicator)	6.0		4.6-8.0	
ROTEIN Method:Reagent Strip Method Protein Error of indicator/SSA Test)	TRACE		Negative	
LUCOSE Method:Reagent Strip Method (GOD- OD/Benedict's Semiquantitative method)	NIL	%	Negative	
ETONES Method:Reagent Strip Method Sodium Nitroprusside Test)	NEGATIVE		Negative	

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LEUCOCYTE	ESTERASE
-----------	----------

UROBILINOGEN

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

BILIRUBIN

Method:Reagent Strip Method (Diazonium

Method/FOUCHET'S METHOD)

BLOOD Method:Reagent Strip Method

(Peroxidase - Like Activity)

NITRITES

Method:Reagent Strip Method (Diazonium

Method)

MICROSCOPIC EXAMINATION

RBCs

Method:Microscopic Examination

EPITHELIAL CELLS

Method:Microscopic Examination

PUS CELLS

Method:Microscopic Examination

BACTERIA

Method:Microscopic Examination

CRYSTALS

Method:Microscopic Examination

Method:Microscopic Examination

OTHERS

NEGATIVE

NIL

Negative

<1.0 mg/dL

NEGATIVE

NEGATIVE

Negative

Negative

NEGATIVE

Negative

/HPF NIL

/HPF

/HPF

0-2 0-5

1-2

0-3

NOT SEEN

6-8

NOT SEEN

Not Seen

Not Seen

NOT SEEN

Not Seen

*** End Of Report ***

PROCESSED BY

: MOGHAL HAJAVALI

VERIFIED BY : G SURESH

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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)
 - Incorrect specimen type. C)
 - 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 d) 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 fest 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 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 Bengaiuru shall have exclusive jurisdiction in all disputes/claims concerning the tests and the results of the tests.

P? ß 100B F 60~ 0.15-100 Hz 94 V5rate> 99 74 Unconfirmed Diagnosis Chest: 10.0 mm/mV - OTHERWISE NORMAL ECG -V3 V2 Б Limb: 10 mm/mV . Sinus tachycardia..... Speed: 25 mm/sec aVL aVF aVR 12 Lead; Standard Placement 71 50 139 100 319 414 101 --AXIS--Device: Rate PR QRSD QT QTC PQRS III H H



2D - ECHO CARDIOGRAM & COLOUR DOPPLER REPORT

Patient's Name - B.SRAVANI , Age/Sex :- 32Y/F Date: 22-10-2022 OP No: 68758

M-MODE:

LV: 4.4 X 2.9 cms EF: 62 %

FS:31%

LA: 2.7

cms

AO: 2.6

cms

IVS: 1.0

cms

PW: 0.9

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: 1.2 M/s, PV Flow: 0.9M/s, RVSP: 18 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.

Name: B. Sravani

Age/Sex: 32yrs /F

Ref. By: Dr. Dr. N. BHANU PRAVEEN NAIDU MRCP (UK)

Date: 22.10.2022

ULTRASONOGRAPHY OF ABDOMEN

LIVER:

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:

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:

Normal in size, position and texture. No focal lesions.

No calculi / hydronephrosis.

LEFT KIDNEY:

Normal in size, position and texture. No focal lesions.

No calculi / hydronephrosis.

URINARY BLADDER: Minimally Distended.

UTERUS

: Suboptimal.

OVARIES

: Suboptimal.

*No free fluid noted in peritoneal cavity.

CONCLUSION:

NO ABNORMALITY NOTED TO THE VISUALLY NOTED.

Advised review scan with full bladder if clinically indicated / Needed to Review pelvic pathologically

Dr. M. BHAVANI SHANKA

MBBS,MD,DNB (Radio Diagnosic

Consultant, Endovascular and Interventional Radio Carva
MBBS, MD, DNB(Radio Diagnosis), Five

Regd.No: 91805 ultant Endovascular and