



# ECHOCARDIOGRAPHY REPORT

Name:	Mr.BHARATH PAI P	Age: 33	Date:	10/12/2022	
Ref By:	PACKAGE	Sex: M	SRD No:	VL000891	

M MODE

2D ECHO

MV/Area: Normal

AORTA: 21 mm

LA: 40mm

LVIDD: 36 mm

LVIDS: 27 mm

IVSD: 25mm

MV : Normal AV: Normal RWMA: Nil LA: Normal LV: Hypertrophied PV: Normal
TV: Normal
RA: Normal
RV: Normal
IVS:Thickened
A-V Relationship: Normal

IVSD: 25mm IVSS: 28 mm LVPWD: 16 mm LVPWS: 18 mm LVEF: 62 %

IAS: Intact Situs: Solitus V-A Relationship: Normal Syst. V. Drainage: Normal Pericardial Effusion: Nil

Great Vessel Relationship: Normal Pulmonary V Drainage: Normal

**DOPPLER** 

FS: 28%

Pul Velocity: 1.0

MV Velocity:

E: 0.83 m/s

A: 0.63m/s

E/A:1.32

MV Area (PHT):

AO Area:

TV Velocity: 0.6 m/s

AV Velocity: 1.4 m/s

RVSP: 31 mmHg

COLOUR

MR: Mild ASD:
AR: Nil VSD:
TR: Mild PDA
PR:Nil CoA:

Wall motion abnormalities : Nil Pericardium: No pericardial effusion

Vegetation/Thrombus: Nil

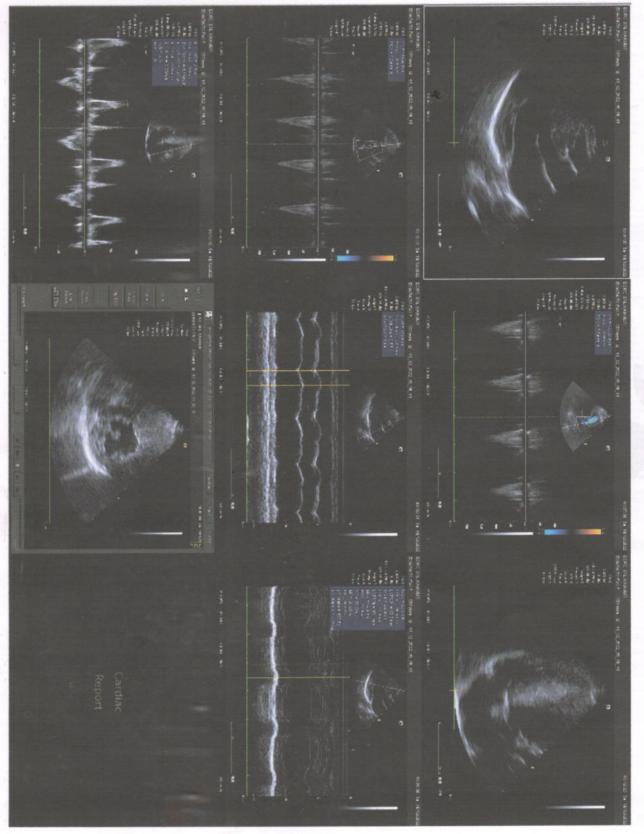
# IMPRESSION:

- HCM
- Mild MR
- Mild TR, Mild PAH
- No RWMA
- Normal LV Systolic function

# DDRC SRL KANNUR

BHARATH PAI P: 10\_12\_2022\_09\_56\_15

20221210



Diagnostic Services

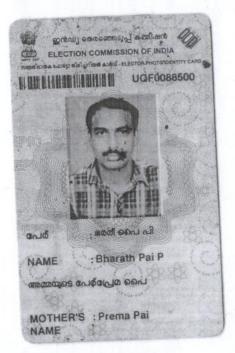
Diagnostic Services

ellour SEX : "nigones / Male ജനനത്തിയതി / വയസ്സ് DATE OF BIRTH/AGE : 12/04/1989 / 25 aadafleome : 17/733, ഗരണഷ്ട്യസാർ; വയലും പി ഒ ADDRESS: 17/733, Ganeshprasad, Vayalalam, P O Thiruvangad 670103 Date: 20/09/2015 ELECTORAL REGISTRATION OFFICER അസംബ്ളി നിയാജകരണ്ഡലം : 13, തലവ്ലേരി ASSEMBLY CONSTITUENCY No. 6 : 13, THALASSERY note move : 77 PART No. NOTE / AITH type and ocamented cases one, close characterings empta-cipation on necessarian enacements over an energy of characteristic energy affairs characteristic over cond-cases over the condition of the condition of the condi- Date of Birth mannessed in this card shall not be treated as a proof of age/D.O.B. for any purpose other than registration in electrical cost. one six out evel 

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CLIENT CODE: CA00010147 - MEDIWHEEL

CLIENT'S NAME AND ADDRESS :

MEDIWHEEL ARCOFEMI HEALTHCARE LIMITED

F701A, LADO SARAI, NEW DELHI,

SOUTH DELHI, DELHI, SOUTH DELHI 110030

DELHI INDIA 8800465156 DDRC SRL DIAGNOSTICS

KERALA, INDIA Tel: 93334 93334

Email: customercare.ddrc@srl.in

**PATIENT NAME: BHARATH PAI P** PATIENT ID: BHARM1204894053

ACCESSION NO: 4053VL000891 AGE: 33 Years SEX: Male ABHA NO:

RECEIVED: 10/12/2022 08:40 10/12/2022 15:41 DRAWN: REPORTED:

REFERRING DOCTOR: SELF CLIENT PATIENT ID:

**Test Report Status** Results **Biological Reference Interval Units** <u>Final</u>

# MEDIWHEEL HEALTH CHECKUP BELOW 40(M)2DECHO

**OPTHAL** 

**COMPLETED OPTHAL** 

PHYSICAL EXAMINATION

PHYSICAL EXAMINATION COMPLETED









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# MEDIWHEEL HEALTH CHECKUP BELOW 40(M)2DECHO

DIIN	/CREAT	DATIO
DUN	/LKEAI	KAIIU

5.00 - 15.00 **BUN/CREAT RATIO** 14

**CREATININE, SERUM** 

**CREATININE** 0.90 18 - 60 yrs : 0.9 - 1.3 mg/dL

**GLUCOSE, POST-PRANDIAL, PLASMA** 

GLUCOSE, POST-PRANDIAL, PLASMA 104 Diabetes Mellitus : > or = 200. mg/dL

Impaired Glucose tolerance/ Prediabetes: 140 - 199. Hypoglycemia: < 55.

**GLUCOSE, FASTING, PLASMA** 

GLUCOSE, FASTING, PLASMA Diabetes Mellitus : > or = 126. mg/dL 86

Impaired fasting Glucose/ Prediabetes: 101 - 125. Hypoglycemia : < 55.

**GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE** 

**BLOOD** 

Normal : 4.0 - 5.6%. % GLYCOSYLATED HEMOGLOBIN (HBA1C) 5.9

Non-diabetic level : < 5.7%. Diabetic : >6.5%

Glycemic control goal

More stringent goal : < 6.5 %. General goal : < 7%. Less stringent goal : < 8%.

Glycemic targets in CKD :-If eGFR > 60 : < 7%. If eGFR < 60:7-8.5%.

LIPID PROFILE, SERUM

Desirable: < 200 193 mg/dL **CHOLESTEROL** 

Borderline: 200-239 : >or= 240

Normal : < 150 **TRIGLYCERIDES** 69 mg/dL

: 150-199 Hiah

Hypertriglyceridemia: 200-499

Very High: > 499

General range: 40-60 mg/dL HDL CHOLESTEROL 48 Ontimum : < 100 mg/dL DIRECT LDL CHOLESTEROL 125

Above Optimum: 100-139 Borderline High : 130-159 High : 160-189

Very High : >or= 190









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NON HDL CHOLESTEROL	145	High	Desirable-Less than 130 Above Desirable-130-159 Borderline High-160-189 High-190-219 Very High- >or =220	mg/dL
CHOL/HDL RATIO	4.0		3.3 - 4.4 Low Risk 4.5 - 7.0 Average Risk 7.1 - 11.0 Moderate Risk > 11.0 High Risk	
LDL/HDL RATIO	2.6		0.5-3 Desirable/Low risk 3.1-6 Borderline/Moderate risk >6.0 High Risk	
VERY LOW DENSITY LIPOPROTEIN LIVER FUNCTION TEST WITH GGT	13.7		= 30.0</td <td>mg/dL</td>	mg/dL
BILIRUBIN, TOTAL	1.20		General Range : < 1.1	mg/dL
BILIRUBIN, DIRECT	0.24		General Range : < 0.3	mg/dL
BILIRUBIN, INDIRECT	0.96	High	0.00 - 0.60	mg/dL
TOTAL PROTEIN	7.4		Ambulatory: 6.4 - 8.3 Recumbant: 6 - 7.8	g/dL
ALBUMIN	4.9		20-60yrs : 3.5 - 5.2	g/dL
GLOBULIN	2.5		2.0 - 4.0	g/dL
ALBUMIN/GLOBULIN RATIO	1.9		1.0 - 2.0	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	77		Adults: < 40	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT)	165	High	Adults: < 45	U/L
ALKALINE PHOSPHATASE	51		Adult(<60yrs): 40 - 130	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT) TOTAL PROTEIN, SERUM	60		Adult(male): < 60	U/L
TOTAL PROTEIN	7.4		Ambulatory: 6.4 - 8.3 Recumbant: 6 - 7.8	g/dL
URIC ACID, SERUM				
URIC ACID  ABO GROUP & RH TYPE, EDTA WHOLE BLOOD	3.7		Adults: 3.4-7	mg/dL
ABO GROUP	TYPE B			
RH TYPE	POSITIVE			
BLOOD COUNTS,EDTA WHOLE BLOOD	4.5.5		12.0 17.0	/ 11
HEMOGLOBIN	16.2		13.0 - 17.0	g/dL
RED BLOOD CELL COUNT	5.13		4.5 - 5.5	mil/µL









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WHITE BLOOD CELL COUNT	8.80		4.0 - 10.0	thou/µL
PLATELET COUNT	251		150 - 410	thou/µL
RBC AND PLATELET INDICES				
HEMATOCRIT	48.2		40 - 50	%
MEAN CORPUSCULAR VOL	94.0		83 - 101	fL
MEAN CORPUSCULAR HGB.	31.7		27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION	33.7		31.5 - 34.5	g/dL
MENTZER INDEX	18.3			
MEAN PLATELET VOLUME	9.9		6.8 - 10.9	fL
WBC DIFFERENTIAL COUNT				
SEGMENTED NEUTROPHILS	59		40 - 80	%
LYMPHOCYTES	28		20 - 40	%
MONOCYTES	2		2 - 10	%
EOSINOPHILS	10	High	1 - 6	%
BASOPHILS	1		0 - 2	%
ABSOLUTE NEUTROPHIL COUNT	5.19		2.0 - 7.0	thou/µL
ABSOLUTE LYMPHOCYTE COUNT	2.46		1 - 3	thou/µL
ABSOLUTE MONOCYTE COUNT	0.18	Low	0.20 - 1.00	thou/µL
ABSOLUTE EOSINOPHIL COUNT	0.88	High	0.02 - 0.50	thou/µL
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	2.1			
ERYTHROCYTE SEDIMENTATION RATE (ESR),W BLOOD				
SEDIMENTATION RATE (ESR)	5		0 - 14	mm at 1 hr
STOOL: OVA & PARASITE				
COLOUR	BROWN			
CONSISTENCY	SEMI LIQUID			
ODOUR	FAECAL			
MUCUS	ABSENT		NOT DETECTED	
VISIBLE BLOOD	ABSENT		ABSENT	
POLYMORPHONUCLEAR LEUKOCYTES	1-2		0 - 5	/HPF
RED BLOOD CELLS	NOT DETECTED		NOT DETECTED	/HPF
CYSTS	NOT DETECTED		NOT DETECTED	
OVA	NOT DETECTED			

**SUGAR URINE - POST PRANDIAL** 









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SUGAR URINE - POST PRANDIAL	NOT DETECTED	NOT DETECTED	
THYROID PANEL, SERUM			
T3	148.20	80.00 - 200.00	ng/dL
T4	10.06	5.10 - 14.10	μg/dl
TSH 3RD GENERATION	3.920	21-50 yrs : 0.4 - 4.2	μIU/mL
PHYSICAL EXAMINATION, URINE			
COLOR	PALE YELLOW		
APPEARANCE	CLEAR		
CHEMICAL EXAMINATION, URINE			
PH	5	4.7 - 7.5	
SPECIFIC GRAVITY	1.020	1.003 - 1.035	
PROTEIN	NOT DETECTED	NOT DETECTED	
GLUCOSE	NOT DETECTED	NOT DETECTED	
KETONES	NOT DETECTED	NOT DETECTED	
BILIRUBIN	NOT DETECTED	NOT DETECTED	
UROBILINOGEN	NORMAL	NORMAL	
NITRITE	NOT DETECTED	NOT DETECTED	
MICROSCOPIC EXAMINATION, URINE			
RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
WBC	2-3	0-5	/HPF
EPITHELIAL CELLS	DETECTED (OCCASIONAL)	NOT DETECTED	/HPF
CASTS	ABSENT		
CRYSTALS	ABSENT		
BACTERIA	NOT DETECTED	NOT DETECTED	
SUGAR URINE - FASTING			
SUGAR URINE - FASTING	NOT DETECTED	NOT DETECTED	

Interpretation(s)
CREATININE, SERUM-Higher than normal level may be due to:
• Blockage in the urinary tract

- Kidney problems, such as kidney damage or failure, infection, or reduced blood flow
  Loss of body fluid (dehydration)
  Muscle problems, such as breakdown of muscle fibers

- Problems during pregnancy, such as seizures (eclampsia)), or high blood pressure caused by pregnancy (preeclampsia)

Lower than normal level may be due to:









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• Myasthenia Gravis

Muscular dystrophy

GLUCOSE, POST-PRANDIAL, PLASMA-

ADA Guidelines for 2hr post prandial glucose levels is only after ingestion of 75grams of glucose in 300 ml water, over a period of 5 minutes. GLUCOSE, FASTING, PLASMA-

ADA 2012 guidelines for adults as follows: Pre-diabetics: 100 - 125 mg/dL Diabetic: > or = 126 mg/dL

(Ref: Tietz 4th Edition & ADA 2012 Guidelines)

GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD-Used For:

- 1. Evaluating the long-term control of blood glucose concentrations in diabetic patients.
- 2.Diagnosing diabetes.3.Identifying patients at increased risk for diabetes (prediabetes).

The ADA recommends measurement of HbA1c (typically 3-4 times per year for type 1 and poorly controlled type 2 diabetic patients, and 2 times per year for well-controlled type 2 diabetic patients) to determine whether a patients metabolic control has remained continuously within the target range.

- 1.eAG (Estimated average glucose) converts percentage HbA1c to md/dl, to compare blood glucose levels.
- 2. eAG gives an evaluation of blood glucose levels for the last couple of months. 3. eAG is calculated as eAG (mg/dl) = 28.7 \* HbA1c 46.7

**HbA1c Estimation can get affected due to :**I.Shortened Erythrocyte survival : Any condition that shortens erythrocyte survival or decreases mean erythrocyte age (e.g. recovery from acute blood loss, hemolytic

II. Vitamin C & E are reported to falsely lower test results. Fructosamine is recommended in these patients which indicates diabetes control over 15 days.

III. Vitamin C & E are reported to falsely lower test results. (possibly by inhibiting glycation of hemoglobin.

III. Iron deficiency anemia is reported to increase test results. Hypertriglyceridemia, uremia, hyperbilirubinemia, chronic alcoholism, chronic ingestion of salicylates & opiates addiction are reported to interfere with some assay methods, falsely increasing results.

IV.Interference of hemoglobinopathies in HbA1c estimation is seen in

a.Homozygous hemoglobinopathy. Fructosamine is recommended for testing of HbA1c. b.Heterozygous state detected (D10 is corrected for HbS & HbC trait.)

C.HbF > 25% on alternate paltform (Boronate affinity chromatography) is recommended for testing of HbA1c.Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy

LIPID PROFILE, SERUM-Serum cholesterol is a blood test that can provide valuable information for the risk of coronary artery disease This test can help determine your risk

of the build up of plaques in your arteries that can lead to narrowed or blocked arteries throughout your body (atherosclerosis). High cholesterol levels usually don't cause any signs or symptoms, so a cholesterol test is an important tool. High cholesterol levels often are a significant risk factor for heart disease and important for diagnosis of hyperlipoproteinemia, atherosclerosis, hepatic and thyroid diseases.

Serum Triglyceride are a type of fat in the blood. When you eat, your body converts any calories it doesn"""""""t need into triglycerides, which are stored in fat cells. High diabetes with elevated blood sugar levels. Analysis has proven useful in the diagnosis and treatment of patients with diabetes mellitus, nephrosis, liver obstruction, other diseases involving lipid metabolism, and various endocrine disorders. In conjunction with high density lipoprotein and total serum cholesterol, a triglyceride determination provides valuable information for the assessment of coronary heart disease risk. It is done in fasting state.

High-density lipoprotein (HDL) cholesterol. This is sometimes called the ""good"" cholesterol because it helps carry away LDL cholesterol, thus keeping arteries open and blood flowing more freely. HDL cholesterol is inversely related to the risk for cardiovascular disease. It increases following regular exercise, moderate alcohol consumption and with oral estrogen therapy. Decreased levels are associated with obesity, stress, cigarette smoking and diabetes mellitus.

SERUM LDL The small dense LDL test can be used to determine cardiovascular risk in individuals with metabolic syndrome or established/progressing coronary artery disease, individuals with triglyceride levels between 70 and 140 mg/dL, as well as individuals with a diet high in trans-fat or carbohydrates. Elevated sdLDL levels are associated with metabolic syndrome and an 'atherogenic lipoprotein profile', and are a strong, independent predictor of cardiovascular disease. Elevated levels of LDL arise from multiple sources. A major factor is sedentary lifestyle with a diet high in saturated fat. Insulin-resistance and pre-diabetes have also been implicated, as has genetic predisposition. Measurement of sdLDL allows the clinician to get a more comprehensive picture of lipid risk factors and tailor treatment accordingly. Reducing LDL levels will reduce the risk of CVD and MI.

Non HDL Cholesterol - Adult treatment panel ATP III suggested the addition of Non-HDL Cholesterol as an indicator of all atherogenic lipoproteins (mainly LDL and VLDL). NICE guidelines recommend Non-HDL Cholesterol measurement before initiating lipid lowering therapy. It has also been shown to be a better marker of risk in both primary and secondary prevention studies.

# Recommendations:

Results of Lipids should always be interpreted in conjunction with the patient's medical history, clinical presentation and other findings.

NON FASTING LIPID PROFILE includes Total Cholesterol, HDL Cholesterol and calculated non-HDL Cholesterol. It does not include trialycerides and may be best used in

patients for whom fasting is difficult.

TOTAL PROTEIN, SERUM-Serum total protein, also known as total protein, is a biochemical test for measuring the total amount of protein in serum. Protein in the plasma is made up of albumin and globulin

Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstrom's disease



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Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome,Protein-losing enteropathy etc. URIC ACID, SERUM-

Causes of Increased levels

Dietary

- High Protein Intake
- Prolonged Fasting,Rapid weight loss.

Gout

Lesch nyhan syndrome.

Type 2 DM.

Metabolic syndrome.

## Causes of decreased levels

- Low Zinc Intake
- OCP's
- Multiple Sclerosis

Nutritional tips to manage increased Uric acid levels

- · Drink plenty of fluids
- · Limit animal proteins
- · High Fibre foods
- Vit C Intake

• Antioxidant rich foods ABO GROUP & RH TYPE, EDTA WHOLE BLOOD-

Blood group is identified by antigens and antibodies present in the blood. Antigens are protein molecules found on the surface of red blood cells. Antibodies are found in plasma. To determine blood group, red cells are mixed with different antibody solutions to give A,B,O or AB.

Disclaimer: "Please note, as the results of previous ABO and Rh group (Blood Group) for pregnant women are not available, please check with the patient records for availability of the same.

The test is performed by both forward as well as reverse grouping methods.

BLOOD COUNTS, EDTA WHOLE BLOOD-

The cell morphology is well preserved for 24hrs. However after 24-48 hrs a progressive increase in MCV and HCT is observed leading to a decrease in MCHC. A direct smear is recommended for an accurate differential count and for examination of RBC morphology.

RBC AND PLATELET INDICESMentzer index (MCV/RBC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anaemia(>13) from Beta thalassaemia trait (<13) in patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait. WBC DIFFERENTIAL COUNT-

The optimal threshold of 3.3 for NLR showed a prognostic possibility of clinical symptoms to change from mild to severe in COVID positive patients. When age = 49.5 years old and NLR = 3.3, 46.1% COVID-19 patients with mild disease might become severe. By contrast, when age < 49.5 years old and NLR < 3.3, COVID-19 patients tend to

(Reference to - The diagnostic and predictive role of NLR, d-NLR and PLR in COVID-19 patients; A.-P. Yang, et al.; International Immunopharmacology 84 (2020) 106504 This ratio element is a calculated parameter and out of NABL scope.

ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD-TEST DESCRIPTION: Erythrocyte sedimentation rate (ESR) is a test that indirectly measures the degree of inflammation present in the body. The test actually measures the rate of fall (sedimentation) of erythrocytes in a sample of blood that has been placed into a tall, thin, vertical tube. Results are reported as the millimetres of clear fluid (plasma) that are present at the top portion of the tube after one hour. Nowadays fully automated instruments are available to measure ESR.

ESR is not diagnostic; it is a non-specific test that may be elevated in a number of different conditions. It provides general information about the presence of an inflammatory condition.CRP is superior to ESR because it is more sensitive and reflects a more rapid change.

# TEST INTERPRETATION

**Increase** in: Infections, Vasculities, Inflammatory arthritis, Renal disease, Anemia, Malignancies and plasma cell dyscrasias, Acute allergy Tissue injury, Pregnancy, Estrogen medication, Aging.

Finding a very accelerated ESR(>100 mm/hour) in patients with ill-defined symptoms directs the physician to search for a systemic disease (Paraproteinemias, Disseminated malignancies, connective tissue disease, severe infections such as bacterial endocarditis)

In pregnancy BRI in first trimester is 0-48 mm/hr(62 if anemic) and in second trimester (0-70 mm /hr(95 if anemic). ESR returns to normal 4th week post partum.

Decreased in: Polycythermia vera, Sickle cell anemia

False elevated ESR : Increased fibrinogen, Drugs(Vitamin A, Dextran etc), Hypercholesterolemia False Decreased: Poikilocytosis, (SickleCells, spherocytes), Microcytosis, Low fibrinogen, Very high WBC counts, Drugs (Quinine, salicylates)

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition; 2. Paediatric reference intervals, AACC Press, 7th edition, Edited by S. Soldin; 3. The reference for









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the adult reference range is "Practical Haematology by Dacie and Lewis,10th edition. SUGAR URINE - POST PRANDIAL-METHOD: DIPSTICK/BENEDICT'S TEST SUGAR URINE - FASTING-METHOD: DIPSTICK/BENEDICT'S TEST





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DELHI INDIA 8800465156 DDRC SRL DIAGNOSTICS

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Email: customercare.ddrc@srl.in

**PATIENT NAME: BHARATH PAI P** PATIENT ID: BHARM1204894053

ACCESSION NO: 4053VL000891 AGE: 33 Years SEX: Male ABHA NO:

RECEIVED: 10/12/2022 08:40 10/12/2022 15:41 DRAWN: REPORTED:

**REFERRING DOCTOR: SELF** CLIENT PATIENT ID:

**Test Report Status** Results Units <u>Final</u>

# MEDIWHEEL HEALTH CHECKUP BELOW 40(M)2DECHO

**ECG WITH REPORT** 

**RFPORT** 

COMPLETED

2D - ECHO WITH COLOR DOPPLER

COMPLETED

**USG ABDOMEN AND PELVIS** 

COMPLETED

**CHEST X-RAY WITH REPORT** 

**REPORT** 

COMPLETED

\*\*End Of Report\*\* Please visit www.srlworld.com for related Test Information for this accession

JINSHA KRISHNAN

LAB TECHNOLOGIST

**DR.INDUSARATH S CONSULTANT PATHOLOGIST** 

**SREENA A** LAB TECHNOLOGIST

**Msc Medical Biochemistry** 





Scan to View Report



# MEDICAL EXAMINATION REPORT (MER)

If the examinee is suffering from an acute life threatening situation, you may be obliged to disclose the result of the medical examination to the examinee.

2. Mark of Ide 3. Age/Date of 4. Photo ID Cl	entification : (M f Birth : 3	lole/Scar/an	BHAR, by other (specify -04 - 198 ction Card/PAN	location Gende	r: FAM	
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# **DDRC SRL** Diagnostics Private Limited

Corp. Office: DDRC SRL Tower, G- 131, Panampilly Nagar, Ernakulam - 682 036 Ph No. 0484-2318223, 2318222, e-mail: info@ddrcsrl.com, web: www.ddrcsrl.com

Any disorders of Urinary System?



 Any disorder of the Eyes, Ears, Nose, Throat or Mouth & Skin



# FOR FEMALE CANDIDATES ONLY

a. Is there any history of diseases of breast/genital organs?

b. Is there any history of abnormal PAP Smear/Mammogram/USG of Pelvis or any other tests? (If yes attach reports)

e. For Parous Women, were there any complication during pregnancy such as gestational diabetes,

d. Do you have any history of miscarriage/

abortion or MTP

hypertension etc

c. Do you suspect any disease of Uterus, Cervix or Ovaries?

f. Are you now pregnant? If yes, how many months?

# CONFIDENTAIL COMMENTS FROM MEDICAL EXAMINER

➤ Was the examinee co-operative?

> Is there anything about the examine's health, lifestyle that might affect him/her in the near future with regard his/her job?

Are there any points on which you suggest further information be obtained?

> Based on your clinical impression, please provide your suggestions and recommendations below;

> Do you think he/she is MEDICALLY FIT or UNFIT for employment.

medically

# MEDICAL EXAMINER'S DECLARATION

I hereby confirm that I have examined the above individual after verification of his/her identity and the findings stated above are true and correct to the best of my knowledge.

Name & Signature of the Medical Examiner

Seal of Medical Examiner

Dr. INDUSARATH.S, MBBS,MD,DNB Regd. No: 41964 DDRC SRL, KANNUR

Name & Seal of DDRC SRL Branch

Date: 10.12

Date & Time

# DDRC SRL Diagnostics Private Limited

TANNUR



# OPTHALMOLOGY REPORT

# TO WHOM-SO-EVER IT MAY CONCERN

This is to certify that I have examined Mr. BHARATH PAI, 33 years Male on 10.12.2022 and his visual standards are as follows:

sans antes	OD	os
UNCORRECTED DISTANCE VISUAL ACUITY	6/12(P)	6/12
UNCORRECTED NEAR VISUAL ACUITY	N6	N6
BEST CORRECTED VISUAL ACUITY	6/6	6/6
COLOUR VISION	NORMAL	NORMAL

NOTE: NO HISTORY OF SPECS HISTORY OF CARDIAC PROBLEM ON RX

VIMEGA .V OPTOMETRIST Date: ......

DATE: 10.12.2022



Name	Mr. BHARATH PAI.P	Age/Sex	33/Male
Ref: By:	MEDI WHEEL	Date	10.12.2022

Thanks for referral

# CHEST X-RAY - PA VIEW

Trachea is central. Carina and principal bronchi are normal.

Cardio-thoracic ratio is within normal limits.

Both lungs show normal Broncho-vascular markings. No definite focal opacities noted.

No volume loss in either hemithorax.

No definite mediastinal widening or other abnormalities noted.

CP angles, diaphragm and soft tissue shadows - not remarkable.

Mild deviation of upper thoracic spine is noted towards left of midline.

# IMPRESSION:

- Mild scoliosis of upper thoracic spine towards left.
- · No other abnormality detected.

DR. P. NIYAZI NASIR, MBBS, DMRD

(Because of technical and technological limitation complete diagnosis cannot be assured on imaging sonography. Clinical correlation, consultation if required repeat imaging required in the event of controversies. This document is not for legal purposes).

Dr. P. NIYAZI NASIR. MBBS, DMRD REG. No. 41419 CONSULTANT RADIOLOGIST DDRC SRL DIAGNOSTIC (P) LTD. KANNUR

