Name PID No.	: Mr. NAGARAJAN V : MED111408894	Register On	: 10/	12/2022 1:09 PM	m
SID No.	: 222021108	-	: 10	/12/2022 2:43 PM	
Age / Sex	: 53 Year(s) / Male	Report On	: 12	2/12/2022 8:56 AM	MEDALL
Туре	: OP	Printed On	: 21	/12/2022 6:19 PM	
Ref. Dr	: MediWheel				
<u>Investiga</u>	ation	<u>Observed</u> <u>Value</u>	<u>t</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
TYPING	GROUPING AND Rh	'A' 'Positi	ve'		
	RETATION: Reconfirm the Blood g	roup and Typing b	efore l	blood transfusion	
	e Blood Count With - ESR	1 51 6			
Haemogl (EDTA Blo	obin oodSpectrophotometry)	14.5		g/dL	13.5 - 18.0
Packed C	Cell Volume(PCV)/Haematocrit	43.3		%	42 - 52
RBC Cou (EDTA Blo	ant pod/Impedance Variation)	4.82		mill/cu.mm	4.7 - 6.0
	rpuscular Volume(MCV) oodDerived from Impedance)	89.8		fL	78 - 100
	rpuscular Haemoglobin(MCH) oodDerived from Impedance)	30.1		pg	27 - 32
concentra	rpuscular Haemoglobin ation(MCHC) ood/Derived from Impedance)	33.5		g/dL	32 - 36
RDW-CV		13.0		%	11.5 - 16.0
RDW-SE (EDTA Blo) ood/Derived from Impedance)	41.7		fL	39 - 46
	akocyte Count (TC)	5910		cells/cu.mm	4000 - 11000
Neutroph (EDTA Blo <i>Cytometry</i>)	ood/Impedance Variation & Flow	50.2		%	40 - 75
Lymphoc (EDTA Blo <i>Cytometry</i>)	ood/Impedance Variation & Flow	36.7		%	20 - 45
Eosinoph (EDTA Blo Cytometry)	ood/Impedance Variation & Flow	6.6		%	01 - 06





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The results pertain to sample tested.

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Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> Reference Interval
Monocytes (EDTA Blood/Impedance Variation & Flow Cytometry)	5.4	%	01 - 10
Basophils (EDTA Blood/Impedance Variation & Flow Cytometry)	1.1	%	00 - 02
INTERPRETATION: Tests done on Automated F	ive Part cell counte	er. All abnormal results ar	e reviewed and confirmed microscopically.
Absolute Neutrophil count (EDTA Blood/Impedance Variation & Flow Cytometry)	2.96	10^3 / µl	1.5 - 6.6
Absolute Lymphocyte Count (EDTA Blood/Impedance Variation & Flow Cytometry)	2.17	10^3 / µl	1.5 - 3.5
Absolute Eosinophil Count (AEC) (EDTA Blood/Impedance Variation & Flow Cytometry)	0.39	10^3 / µl	0.04 - 0.44
Absolute Monocyte Count (EDTA Blood/Impedance Variation & Flow Cytometry)	0.32	10^3 / µl	< 1.0
Absolute Basophil count (EDTA Blood/Impedance Variation & Flow Cytometry)	0.07	10^3 / µl	< 0.2
Platelet Count (EDTA Blood/Impedance Variation)	231	10^3 / µl	150 - 450
MPV (EDTA Blood/Derived from Impedance)	10.0	fL	7.9 - 13.7
PCT (EDTA Blood/Automated Blood cell Counter)	0.231	%	0.18 - 0.28
ESR (Erythrocyte Sedimentation Rate) (Blood/Automated - Westergren method)	9	mm/hr	< 20
BUN / Creatinine Ratio	12.56		6.0 - 22.0
Glucose Fasting (FBS) (Plasma - F/ <i>GOD-PAP</i>)	159.3	mg/dL	Normal: < 100 Pre Diabetic: 100 - 125





Diabetic: >= 126

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Type : OP	Printed On : 2	1/12/2022 6:19 PM	
Ref. Dr : MediWheel			
Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	Biological Reference Interval
INTERPRETATION: Factors such as type, blood glucose level.	quantity and time of food	l intake, Physical activity	v, Psychological stress, and drugs can influence
Glucose Postprandial (PPBS) (Plasma - PP/GOD-PAP)	173.2	mg/dL	70 - 140
INTERPRETATION: Factors such as type, quantity and time of foc Fasting blood glucose level may be higher the resistance, Exercise or Stress, Dawn Phenom	an Postprandial glucose,	because of physiological	surge in Postprandial Insulin secretion, Insulin
Urine Glucose(PP-2 hours) (Urine - PP)	Positive(++)		Negative
Blood Urea Nitrogen (BUN) (Serum/Urease UV / derived)	13.7	mg/dL	7.0 - 21
Creatinine (Serum/ <i>Modified Jaffe</i>)	1.09	mg/dL	0.9 - 1.3
INTERPRETATION: Elevated Creatinine v ingestion of cooked meat, consuming Protein such as cefoxitin, cefazolin, ACE inhibitors, a etc.	/ Creatine supplements, I	Diabetic Ketoacidosis, pr	
Uric Acid (Serum/ <i>Enzymatic</i>)	5.4	mg/dL	3.5 - 7.2
Liver Function Test			
Bilirubin(Total) (Serum/DCA with ATCS)	0.59	mg/dL	0.1 - 1.2
Bilirubin(Direct) (Serum/Diazotized Sulfanilic Acid)	0.15	mg/dL	0.0 - 0.3
Bilirubin(Indirect) (Serum/Derived)	0.44	mg/dL	0.1 - 1.0
SGOT/AST (Aspartate Aminotransferase) (Serum/ <i>Modified IFCC</i>)	31.7	U/L	5 - 40
SGPT/ALT (Alanine Aminotransferase (Serum/ <i>Modified IFCC</i>)	e) 45.2	U/L	5 - 41
GGT(Gamma Glutamyl Transpeptidas (Serum/IFCC / Kinetic)	e) 49.0	U/L	< 55
Dr. N.V.VARDHINI Ph.D Consultant Geneticist			DR CURUPRITAJ PATHOLOGIST Reg No : 13-48036
VERIFIED BY			APPROVED BY
The results pertain to sample tested		D	e 3 of 8

The results pertain to sample tested.

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Ref. Dr	: MediWheel		

Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
Alkaline Phosphatase (SAP) (Serum/ <i>Modified IFCC</i>)	45.1	U/L	56 - 119
Total Protein (Serum/Biuret)	7.53	gm/dl	6.0 - 8.0
Albumin (Serum/Bromocresol green)	4.52	gm/dl	3.5 - 5.2
Globulin (Serum/Derived)	3.01	gm/dL	2.3 - 3.6
A : G RATIO (Serum/Derived)	1.50		1.1 - 2.2
<u>Lipid Profile</u>			
Cholesterol Total (Serum/CHOD-PAP with ATCS)	204.5	mg/dL	Optimal: < 200 Borderline: 200 - 239 High Risk: >= 240

Remark: kindly correlate clinically, suggested repeat testing with a fresh sample, if clinically indicated.

Triglycerides (Serum/GPO-PAP with ATCS)	245.9	mg/dL	Optimal: < 150 Borderline: 150 - 199
			High: 200 - 499
			Very High: >= 500

INTERPRETATION: The reference ranges are based on fasting condition. Triglyceride levels change drastically in response to food, increasing as much as 5 to 10 times the fasting levels, just a few hours after eating. Fasting triglyceride levels show considerable diurnal variation too. There is evidence recommending triglycerides estimation in non-fasting condition for evaluating the risk of heart disease and screening for metabolic syndrome, as non-fasting sample is more representative of the `usual_circulating level of triglycerides during most part of the day.

Remark: kindly correlate clinically, suggested repeat testing with a fresh sample, if clinically indicated.

HDL Cholesterol	38.3	mg/dL	Optimal(Negative Risk Factor): >=
(Serum/Immunoinhibition)			60
			Borderline: 40 - 59
			High Risk: < 40

Remark: kindly correlate clinically, suggested repeat testing with a fresh sample, if clinically indicated.





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Ref. Dr	: MediWheel		

Investigation **Observed** <u>Unit</u> **Biological** Value Reference Interval Optimal: < 100LDL Cholesterol 117 mg/dL Above Optimal: 100 - 129 (Serum/Calculated) Borderline: 130 - 159 High: 160 - 189 Very High: >=190 < 30 49.2 mg/dL VLDL Cholesterol (Serum/Calculated) Optimal: < 130 Non HDL Cholesterol 166.2 mg/dL Above Optimal: 130 - 159 (Serum/Calculated) Borderline High: 160 - 189 High: 190 - 219 Very High: ≥ 220

INTERPRETATION: 1.Non-HDL Cholesterol is now proven to be a better cardiovascular risk marker than LDL Cholesterol. 2.It is the sum of all potentially atherogenic proteins including LDL, IDL, VLDL and chylomicrons and it is the "new bad cholesterol" and is a co-primary target for cholesterol lowering therapy.

Total Cholesterol/HDL Cholesterol Ratio (Serum/ <i>Calculated</i>)	5.3		Optimal: < 3.3 Low Risk: 3.4 - 4.4 Average Risk: 4.5 - 7.1 Moderate Risk: 7.2 - 11.0 High Risk: > 11.0
Triglyceride/HDL Cholesterol Ratio (TG/HDL) (Serum/ <i>Calculated</i>)	6.4		Optimal: < 2.5 Mild to moderate risk: 2.5 - 5.0 High Risk: > 5.0
LDL/HDL Cholesterol Ratio (Serum/Calculated)	3.1		Optimal: 0.5 - 3.0 Borderline: 3.1 - 6.0 High Risk: > 6.0
<u>Glycosylated Haemoglobin (HbA1c)</u>			
HbA1C (Whole Blood/ <i>HPLC</i>)	7.2	%	Normal: 4.5 - 5.6 Prediabetes: 5.7 - 6.4 Diabetic: >= 6.5

INTERPRETATION: If Diabetes - Good control : 6.1 - 7.0 %, Fair control : 7.1 - 8.0 %, Poor control >= 8.1 %





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Ref. Dr	:	MediWheel					
<u>Investiga</u>	atio	<u>n</u>	<u>Observe</u> <u>Value</u>	d	_	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
		ndly correlate clinically, suggeste			ith		ally indicated.
Estimate (Whole Blo		Average Glucose	159.94	Ļ		mg/dL	
HbA1c pro control as Conditions hypertrigly Conditions	ovi con s th yce s th	mpared to blood and urinary gluco hat prolong RBC life span like Iror ridemia,hyperbilirubinemia,Drugs	se determinations a deficiency anem a, Alcohol, Lead P e or chronic blood	ia Poi I le	, V iso oss	/itamin B12 & Folate defi ning, Asplenia can give fa s, hemolytic anemia, Hem	
	-	ecific antigen - Total(PSA) metric method)	0.27			ng/mL	Normal: 0.0 - 4.0 Inflammatory & Non Malignant conditions of Prostate & genitourinary system: 4.01 - 10.0 Suspicious of Malignant disease of Prostate: > 10.0
<u>THYRO</u> T3 (Triio (Serum/ <i>Ch</i>	<u>D</u>	TATION: REMARK : PSA alone <u>PROFILE / TFT</u> othyronine) - Total iluminescent Immunometric Assay	should not be use	ed	as	an absolute indicator of r ng/ml	nalignancy. 0.4 - 1.81
(CLIA)) INTERPH Comment Total T3 v Metabolica	: ari	ation can be seen in other conditio	n like pregnancy,	dı	rug	gs, nephrosis etc. In such (cases, Free T3 is recommended as it is
		ne) - Total iluminescent Immunometric Assay	5.12			µg/dl	4.2 - 12.0
INTERPE Comment Total T4 v Metabolica	: ari	ation can be seen in other conditio	n like pregnancy,	dı	ruş	gs, nephrosis etc. In such	cases, Free T4 is recommended as it is
	•	oid Stimulating Hormone) iluminescent Immunometric Assay	4.21			µIU/mL	0.35 - 5.50
Dr. N	1.1	V.Vardhini Ph.D ultant Geneticist					DR GURUPRIYA J PATHOLOGIST Reg No : 13-48036
VE	ER	IFIED BY					APPROVED BY

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Ref. Dr	: MediWheel			

InvestigationObserved
ValueUnitBiological
Reference IntervalINTERPRETATION:Reference range for cord blood - upto 201 st trimester: 0.1-2.52 nd trimester 0.2-3.03 rd trimester : 0.3-3.0(Indian Thyroid Society Guidelines)

Comment :

1.TSH reference range during pregnancy depends on Iodine intake, TPO status, Serum HCG concentration, race, Ethnicity and BMI. 2.TSH Levels are subject to circadian variation, reaching peak levels between 2-4am and at a minimum between 6-10PM.The variation can be of the order of 50%,hence time of the day has influence on the measured serum TSH concentrations.

 $3. Values \& amplt \\ 0.03 \ \mu IU/mL need to be clinically correlated due to presence of rare TSH variant in some individuals.$

Others

(Urine)

NIL

INTERPRETATION: Note: Done with Automated Urine Analyser & Automated urine sedimentation analyser. All abnormal reports are reviewed and confirmed microscopically.

Stool Analysis - ROUTINE

Colour (Stool)	Brown	Brown
Blood (Stool)	Absent	Absent
<u>Urine Analysis - Routine</u>		
COLOUR (Urine)	Pale yellow	Yellow to Amber
APPEARANCE (Urine)	Clear	Clear
Protein (Urine/Protein error of indicator)	Negative	Negative
Glucose (Urine/GOD - POD)	Positive(+++)	Negative
Pus Cells (Urine/Automated - Flow cytometry)	Occasional /hpf	NIL
Epithelial Cells (Urine/Automated ⁻ Flow cytometry)	Occasional /hpf	NIL





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Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	Biological Reference Interval
RBCs (Urine/Automated - Flow cytometry)	NIL	/hpf	NIL
Casts (Urine/Automated - Flow cytometry)	NIL	/hpf	NIL
Crystals (Urine/Automated - Flow cytometry)	NIL	/hpf	NIL
Mucus (Stool)	Absent		Absent
Reaction (Stool)	Acidic		Acidic
Consistency (Stool)	Semi Solid		Semi Solid
Ova (Stool)	NIL		NIL
Others (Stool)	NIL		NIL
Cysts (Stool)	NIL		NIL
Trophozoites (Stool)	NIL		NIL
RBCs (Stool)	NIL	/hpf	Nil
Pus Cells (Stool)	1 - 2	/hpf	NIL
Macrophages (Stool)	NIL		NIL
Epithelial Cells (Stool)	NIL	/hpf	NIL



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-- End of Report --

The results pertain to sample tested.

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Name	Mr.NAGARAJAN V	ID	MED111408894
Age & Gender	53/MALE	Visit Date	10/12/2022
Ref Doctor Name	MediWheel		

ACOUSTIC WINDOW : GOOD

DEPARTMENT OF CARDIOLOGY

TRANSTHORACIC RESTING ECHO CARDIOGRAPHY REPORT

ECHO INDICATION: Assessment <u>M MODE & 2-D PARAMETERS</u>:

LV STUDY			
IVS(d) cm	1.0		
IVS(s) cm	1.1		
LPW(d) cm	1.0		
LPW(s) cm	1.0		
LVID(d) cm	4.4		
LVID(s) cm	3.1		
EDV ml	87		
ESV ml	30		
SV ml	56		
EF %	64		
FS %	29		
Parameters	Patient		
	Value		
LA cm	3.4		
AO cm	3.3		

DOPPLER PARAMETERS

Valves	Velocity max(m/sec mm/Hg)
AV	0.9
PV	1.0
MV (E)	0.4
((A)	0.5

REPORT DISCLAIMER

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- 2. The results reported here in are subject to interpretation by qualified medical professionals only.
- 3.Customer identities are accepted provided by the customer or their representative.
- 4.information about the customer's condition at the time of sample collection such as fasting, food consumption, medication, etc are accepted as provided by the customer or representative and shall not be investigated for its truthfulness.
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- 6.Test results should be interpreted in context of clinical and other findings if any.In case of any clarification /doubt , the refrering doctor/patient can contact the respective section head of the laboratory.
- 7.Results of the test are influenced by the various factors such as sensitivity, specificity of the procedures of the tests, quality of the samples and drug interactions etc.,
- 8.If the test results are found not to be correlating clinically can contact the lab in charge for clarification or retesting where practicable within 24 hours from the time of issue of results.
- 9.Liability is limited to the extend of amount billed.
- 10.Reports are subject to interpretation in their entirety partial or selective interpretation may lead to false opinion.
- 11.Disputes, if any, with regard to the report findings are subject to the exclusive jurisdiction of the competent courts chennai only.



Name	Mr.NAGARAJAN V	ID	MED111408894
Age & Gender	53/MALE	Visit Date	10/12/2022
Ref Doctor Name	MediWheel		

FINDINGS:

- v Normal left ventricle systolic function (LVEF 64 %).
- ∨ No regional wall motion abnormality.
- **v** Grade I LV diastolic dysfunction.
- v Normal chambers dimension.
- v MR TRIVIAL.
- **v** Normal right ventricle systolic function.
- **v** Normal pericardium / Intact septae.
- v No clot/aneurysm.

IMPRESSION:

- NORMAL LV SYSTOLIC FUNCTION.
- NO REGIONAL WALL MOTION ABNORMALITY.
- GRADE I LV DIASTOLIC DYSFUNCTION.
- TRIVIAL MR.

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Age & Gender	53/MALE	Visit Date	10/12/2022
Ref Doctor Name	MediWheel		

B. SUDHA RANI (BSPA) CARDIOLOGY

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