







Lab No/ManualNo **Patient** Mrs. ARUNA PATANI 5366957/

UHIDNo/IPNO 300443557 CollectionDate 24/02/2024 9:35AM Age/Gender 28 Years/Female **Receiving Date** 24/02/2024 9:57AM **Bed No/Ward** OPD 24/02/2024 3:28PM **Report Date**

Referred By Dr. Casualty Medical Officer **Report Status** Final **Sample Quality** Normal

Test Name		Result	Unit	Bio. Ref. Range	Method	Sample		
Biochemistry ArcoFemi Healthcare Ltd Below 40 Female								
						Serum		
Gamma GT		13.50	U/L	6.00 - 42.00	Enzymatic method			
						Serum		
Creatinine		0.58	mg/dL	0.50 - 0.90	Jaffe Kinetic Compe	ensated		
Age		28						
Weight		56.2						
Gender		0.85						
eGFR	Н	128.12	mL/minute/1. 73 m2	72 - 110				
						Serum		
Uric Acid		4.0	mg/dL	2.4 - 5.7	Uricase / Peroxidase (Colorimetric)	е		
						Serum		
Fasting Glucose		79.9	mg/dL	< 100.0	Hexokinase			

Dr. Kazumi Gondalia M.D (Path)

Reg.No.: G-21729











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As per ADA Guideline

For Fasting Plasma Glucose Normal: Less than 100 mg/dL Prediabetes: 100 mg/dL to 125 mg/dL Diabetes: 126 mg/dL or Higher

For 2 hrs Plasma Glucose after 75 Gms Glucose load

Normal: Less than 140 mg/dL Prediabetes: 140 to 199 mg/dL Diabetes: 200 mg/dL or higher For Random Plasma Glucose

Diabetes is diagnosed at blood glucose greater than or equal to 200 mg/dL

Serum

Post prandial Glucose Hexokinase 93.8 mg/dL < 140.0

Post prandial Urine Glucose S.N.G. = Sample Not Given

As per ADA Guideline For Fasting Plasma Glucose Normal: Less than 100 mg/dL

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LIVER FUNCTION TEST (LFT) SERUM

Serum

IFCC without pyridoxal SGPT(ALT) 10.90 U/L 0.00 - 33.00phosphate SGOT(AST) 13.70 U/L 0.00 - 32.00IFCC without pyridoxal phosphate

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Alkaline Phosphata	ase	106.4	U/L	35.0 - 140.0	PNP-Standardize
Bilirubin Total		0.27	mg/dL	0.00 - 1.00	Diazo Method
Bilirubin Direct		0.10	mg/dL	0.00 - 0.20	Diazo Method
Bilirubin Indirect		0.17	mg/dL	0.00 - 1.10	Calculate from Total and Direct Billirubin
Protein Total		7.65	g/dL	6.40 - 8.20	Biuret Method
Albumin		4.95	g/dL	3.97 - 4.95	BCG Endpoint
Globulin		2.70	g/dL	2.20 - 3.50	Calculated
A/G Ratio		1.83	Ratio	0.90 - 2.80	Ratio
HbA1c (Glyco Hb)		5.46	%	4.8 % - 5.9 % Nor 5.9 % - 7.0 % Goo diabetic Control 7.0 % - 10.00 % F Diabetic Control >10.0 % Poor dial Control	od Fair
Mean Plasma Gluc	ose	117.2	mg/dL	80.0 - 140.0	
					Serum
Blood Urea		19.0	mg/dL	16.6 - 48.5	Urease,Kinetic,GLDH
BUN*		8.9	mg/dL	6.0 - 20.0	Ureas with UV
					Serum
TOTAL T3*		1.210	ng/mL	0.850 - 2.020	ECLIA.
TOTAL T4*		7.170	ug/dL	5.130 - 14.060	ECLIA.
THYROID STIMUL	ATING HORMONE	1.250	uIU/mL	0.270 - 4.200	ECLIA.

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JCI (USA)

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Referred By Dr. Casualty Medical Officer **Report Status** Final

Sample Quality

(*) Not in NABL Scope

End Of Report









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Test Name	Result	Unit	Bio. Ref. Range	Method	Sample
	A E II	Biochemistry			
LIPID PROFILE (WITH DIRECT LDL	ArcoFemi H	lealthcare Ltd Bo	elow 40 Female		Serum
LIFID PROFILE (WITH DIRECT LDE					Serum
Sample Type	Fasting				
Cholesterol Total	147.80	mg/dL	Less than 160 mg/dL Excellent Less than 200 mg/dL Desirabale 200-239 mg/dL Borderline High 240 mg/dl & over high	Enzymatic (CHE/CHO/POD)	
Triglycerides	57.40	mg/dL	Less than 150 mg/dL Normal 150-199 mg/dL Borderline High 200-499 mg/dL High 500 mg/dL or greater very HIgh	GPO-PAP	
HDL Cholesterol	42.50	mg/dL	Less than 40 mg/dL Low 60 mg/dL or Above Excellent	Homogenous Enz	ymatic
LDL Cholesterol (Direct)	96.00	mg/dL	Less than 80 mg/dL Excellent Less than 100 mg/dL Optimal 100-129 mg/dL Near or above optimal 130-159 mg/dL Borderline High 160-189 mg/dL High 190 mg/dL & above Very High	Homogenous Enz	ymatic
VLDL Cholesterol	11.48	mg/dL	< 30	Calculated	
LDL/HDL RATIO	2.26		< 3.50	Calculated	
Cholesterol Total / HDL Ratio	3.48		< 4.50		

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OPD **Bed No/Ward**

Referred By Dr. Casualty Medical Officer Lab No/ManualNo 5366957/

CollectionDate 24/02/2024 9:35AM

Receiving Date 24/02/2024 11:32AM 24/02/2024 12:26PM **Report Date**

Report Status Final Sample Quality Normal

Test Name Result Unit Bio. Ref. Range Method Sample

Clinical Pathology

ArcoFemi Healthcare Ltd Below 40 Female

URINE ROUTINE EXAMINATION Urine

Physical Examination:

Visual method Quantity 30 ml Colour Pale Yellow Visual method Appearence: Clear Visual method

5 Reaction

Reflectance photometer Sp. Gravity 1.005 1.015 - 1.030 Reflectance photometer/Enzymatic

reaction

Chemical Examination: Reflectance photometer/Manual

U.Albumin Nil **U.Gluocse** Nil **U.Acetone** Nil BS/BP Absent

Microscopic Examination Microscopy

/H.P.F. Pus Cell Occasional Nil /H.P.F. Red Blood Cell Epithelial cell 1-2 /H.P.F.

Cast Absent Absent Crystals **Amorphous** Absent Monilia Absent Other: Absent

End Of Report

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 11:31AM

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Sample Quality

Test Name		Result	Unit	Bio. Ref. Range	Method	Sample
			Haematology			
		ArcoFemi H	lealthcare Ltd Bel	ow 40 Female		
CBC WITH ESR						EDTA Blood
Haemoglobin		14.5	g/dL	12.5 - 16.0	SLS Method	
Hematocrit/PCV		45.0	%	37.0 - 47.0	H.focusing Met	hod
RBC COUNT		5.36	mill/Cmm	4.20 - 5.40	H.focusing imp	
MCV		84.0	fl	83.0 - 101.0	Calculated	
MCH		27.1	pg	27.0 - 31.0	Calculated	
MCHC		32.2	g/dL	32.0 - 36.0	Calculated	
RDW-CV		13.4	%	11.5 - 14.0	Calculated	
Platelet count	Н	470000	/cumm	150000 - 410000	H.focusing impo	edance
Mean Platelet Volume(MPV)*		9.2	fl	8 - 12	Calculated	
Total Leucocyte Count (TLC)		6460.00	/cumm	4000.00 - 10500.00	Flow Cytometry	,
Differential Leucocyte Count					Flowcytometry/	Microscopic
Neutrophils		45	%	40.0 - 70.0		
Lymphocytes	Н	46	%	22 - 45		
Eosinophils		03	%	1.0 - 4.0		
Monocytes		06	%	1.0 - 6.0		
Basophils		00	%	0.0 - 1.0		
Immature Granulocytes		00	%	0 - 2		
Absolute Leucocyte Count						
Absolute Neutrophil Count*		2907	/cumm	1800 - 7700		
Absolute Lymphocyte count*		2971.6	/cumm	1000 - 4800		
Absolute Eosinophil Count (AEC)		193.8	/cumm	0.0 - 450.0		
Absolute Monocyte Count*		387.6	/cumm	0 - 800		
Peripheral Smear Study				ormocytic.Platelets are add n.No Premature cells are s		
Erythrocyte Sedimentation Rate (ESR)		12	mm/hr	0 - 12	Photometric ca stopped flow ki	

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 2:39PM

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 Report Date
 24/02/2024
 4:09PM

Referred By Dr. Casualty Medical Officer **Report Status** Final

Sample Quality

CytoPathology

ArcoFemi Healthcare Ltd Below 40 Female

Cytopathology Pathology Report

Specimen

Cervical PAP smear.

Clinical Diagnosis

LMP: 01/02/2024, P1G1

Gross Description

Two fixed unstained slides received, PAP stain done.

Microscopic Description

Smears are satisfactory for evaluation.

Many superficial, intermediate cells and few parabasal cells seen.

Mild inflammation with predominance of neutrophils seen.

Moderate lactobacilli are seen. No parasites/ fungi.

No evidence of intraepithelial lesion or malignancy.

Diagnosis

Cervical smear - Mild inflammation and no evidence of intraepithelial lesion or malignancy.

Note- The pap test is a screening procedure to aid in the detection of cervical cancer and its precursors. Because false negative results may occur, regular PAP tests are recommended.

(*) Not in NABL Scope

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CollectionDate

24/02/2024 12:16PM

Report Status Sample Quality

Test Name Result Unit Bio. Ref. Range Method Sample

Immuno-Haematology

ArcoFemi Healthcare Ltd Below 40 Female

EDTA Blood **BLOOD GROUPING**

ABO Group "B" **Tube Agglutination Method**

Rh Type Negative

End Of Report

Dr. Jitendra Narendrabhai Nayak

Reg.No:G-14786 Consultant Pathologist