







Lab No/ManualNo **Patient** Mrs. PARVINA GHANCHI 5261866/

UHIDNo/IPNO 300428879 CollectionDate 20/10/2023 9:08AM Age/Gender 30 Years/Female **Receiving Date** 20/10/2023 9:44AM **Bed No/Ward** OPD 20/10/2023 4:11PM **Report Date**

Referred By Dr. Rikin Mahendrabhai Shah **Report Status** Final

Test Name		Result	Unit	Bio. Ref. Range	Method	Sample
			Biochemistry			
		ArcoFemi H	ealthcare Ltd Belov	w 40 Female		
Gamma GT		6.30	U/L	6.00 - 42.00	Enzymatic method	Serum
		0.55			laffa Maadia Oosaa	Serum
Creatinine		0.55	mg/dL	0.50 - 0.90	Jaffe Kinetic Compe	ensated
Age		30				
Weight		66				
Gender eGFR	н	0.85 155.83	mL/minute/1. 73 m2	71 - 140		
Uric Acid		2.7	mg/dL	2.4 - 5.7	Uricase / Peroxidas (Colorimetric)	Serum e
Fasting Glucose		97.7	mg/dL	70.0 - 100.0	Hexokinase	Serum
Post prandial Glucose Post prandial Urine Glucose		119.9 SNG	mg/dL	70.0 - 140.0	Hexokinase	Serum
LIVER FUNCTION TEST (LFT) SEF	RUM					Serum
SGPT(ALT)		10.90	U/L	0.00 - 33.00	IFCC without pyrido phosphate	xal
SGOT(AST)		11.90	U/L	0.00 - 32.00	IFCC without pyrido phosphate	xal
Alkaline Phosphatase		88.3	U/L	35.0 - 140.0	PNP-Standardize	
Bilirubin Total		0.26	mg/dL	0.00 - 1.00	Diazo Method	
Bilirubin Direct		0.13	mg/dL	0.00 - 0.20	Diazo Method	

Dr. Kazumi Gondalia M.D (Path)

Reg.No.: G-21729









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Referred By	Dr. Rikin Mahendrat	ohai	i Shah		Report Status	Fina	l	
Bilirubin Indirect			0.13	mg/dL	0.00 - 1.10		Calculate from Total and Direct Billirubin	
Protein Total			6.63	g/dL	6.40 - 8.20		Biuret Method	
Albumin		L	3.74	g/dL	3.97 - 4.95		BCG Endpoint	
Globulin			2.89	g/dL	2.20 - 3.50		Calculated	
A/G Ratio			1.29	Ratio	0.90 - 2.80		Ratio	
								EDTA Blood
HbA1c (Glyco Hb)			5.94	%	4.8 % - 5.9 % No 5.9 % - 7.0 % Go diabetic Control 7.0 % - 10.00 % Diabetic Control >10.0 % Poor dia Control	ood Fair	Immunoturbidimetric	
Mean Plasma Glucos	se		134.0	mg/dL	80.0 - 140.0			
								Serum
Blood Urea		L	11.1	mg/dL	16.6 - 48.5		Urease,Kinetic,GLDH	
BUN*		L	5.2	mg/dL	6.0 - 20.0		Ureas with UV	
								Serum
TOTAL T3*		Н	2.200	ng/mL	0.850 - 2.020		ECLIA.	
TOTAL T4*			12.040	ug/dL	5.130 - 14.060		ECLIA.	
THYROID STIMULA	TING HORMONE		2.280	uIU/mL	0.270 - 4.200		ECLIA.	

End Of Report

Dr. Kazumi Gondalia M.D (Path) Reg.No.: G-21729

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(*) Not in NABL Scope









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	AmagEanri II	Biochemistry	y elow 40 Female		
LIDID DDOELLE (WITH DIDECT LDI	Arcoremi H	eanneare Lia Bo	elow 40 Female		Serum
LIPID PROFILE (WITH DIRECT LDL Sample Type	Fasting				Serum
• • •	•	(-II	l ann thair 400 mar/dl	En Tymotic	
Cholesterol Total	178.60	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	102.80	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 Hlgh	Enzymatic colorin	netric
Cholesterol HDL	45.30	mg/dL	Less than 40 mg/dL Low 60 mg/dL or Above Excellent	Homogenous Enz	rymatic
LDL Cholesterol (Direct)	120.00	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	rymatic
Cholesterol VLDL	20.56	mg/dL	< 30		
LDL/HDL RATIO	2.65		< 3.50	Calculated	
Cholesterol Total / HDL Ratio	3.94		< 4.50		

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5261866/

DEPARTMENT OF LABORATORY SERVICES

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 20/10/2023 11:02AM

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 20/10/2023 12:00PM

Referred By Dr. Rikin Mahendrabhai Shah Report Status Final

Test Name	Result	Unit	Bio. Ref. Range	Method	Sample
		linical Patho			
	ArcoFemi He	althcare Ltd F	Below 40 Female		
URINE ROUTINE EXAMINATION					Urine
Physical Examination:					
Quantity	20 ml			Visual method	
Colour	Pale Yellow			Visual method	
Appearence:	Clear			Visual method	
Reaction	8.0		4.5 - 8.0	Reflectance ph	otometer
Sp. Gravity	1.015		1.015 - 1.030	Reflectance phoreaction	otometer/Enzymatic
Chemical Examination:				Reflectance photometer/Ma	nual
U.Albumin	Nil				
U.Gluocse	Nil				
U.Acetone	Absent				
BS/BP	Absent				
Microscopic Examination				Microscopy	
Pus Cell	Occasional		/H.P.F.		
Red Blood Cell	Nil		/H.P.F.		
Epithelial cell	1-2		/H.P.F.		
Cast	Absent				
Crystals	Absent				
Amorphous	Absent				
Monilia	Absent				
Other:	Absent				
	End Of Re	eport			

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Test Name		Result	Unit	Bio. Ref. Range	Method Sample
			Haematology		
		ArcoFemi H	lealthcare Ltd Bel	ow 40 Female	
CBC WITH ESR					EDTA Blood
Haemoglobin	L	11.5	g/dL	12.5 - 16.0	SLS Method
Hematocrit/PCV	L	36.2	%	37.0 - 47.0	H.focusing Method
RBC COUNT		4.23	mill/Cmm	4.20 - 5.40	H.focusing impedance
MCV		85.6	fl	83.0 - 101.0	Calculated
MCH		27.2	pg	27.0 - 31.0	Calculated
MCHC	L	31.8	g/dL	32.0 - 36.0	Calculated
RDW-CV	Н	15.6	%	11.5 - 14.0	Calculated
Platelet count		209000	/cumm	150000 - 450000	H.focusing impedance
Mean Platelet Volume(MPV)*		10	fl	8 - 12	Calculated
Total Leucocyte Count (TLC)		7140.00	/cumm	4000.00 - 10500.00	Flow Cytometry
Differential Leucocyte Count					Flowcytometry/Microscopic
Neutrophils		66	%	40.0 - 70.0	
Lymphocytes		28	%	22 - 45	
Eosinophils		02	%	1.0 - 4.0	
Monocytes		04	%	1.0 - 6.0	
Basophils		00	%	0.0 - 1.0	
Immature Granulocytes		00	%	0 - 2	
Absolute Leucocyte Count					
Absolute Neutrophil Count*		4712.4	/cumm	1800 - 7700	
Absolute Lymphocyte count*		1999.2	/cumm	1000 - 4800	
Absolute Eosinophil Count (AEC)		142.8	/cumm	0.0 - 450.0	
Absolute Monocyte Count*		285.6	/cumm	0 - 800	
Peripheral Smear Study		RBCs shows Anisocytosis(+). Platelets are adequate in number. Malarial Parasites are not seen. No Premature cells are seen.			
Erythrocyte Sedimentation Rate (ESR)	Н	31	mm/hr	0 - 12	Modified westergren Method

Japan

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CytoPathology

ArcoFemi Healthcare Ltd Below 40 Female

Cytopathology Pathology Report

Specimen

Cervical PAP smear.

Clinical Diagnosis

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.

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Method

Sample

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Unit Immuno-Haematology

ArcoFemi Healthcare Ltd Below 40 Female

EDTA Blood **BLOOD GROUPING**

"O" **ABO Group Tube Agglutination Method**

Rh Type Negative

End Of Report

Result

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