







Patient	Mr. RAVIRAJ I PARMAR	Lab No/ManualNo	5388961/
UHIDNo/IPNO	300446756	CollectionDate	19/03/2024 2:52PM
Age/Gender	38 Years/Male	Receiving Date	19/03/2024 2:52PM
Bed No/Ward	OPD	Report Date	19/03/2024 7:37PM
Referred By	Dr. Casualty Medical Officer	Report Status	Final
		Sample Quality	

Test Name	Result	Unit	Bio. Ref. Range	e Method	Sample
	ArcoFemi I	Biochemistry Healthcare Ltd Bo			
					Serum
Gamma GT	14.20	U/L	10.00 - 71.00	Enzymatic method	
					Serum
Creatinine	0.73	mg/dL	0.70 - 1.20	Jaffe Kinetic Compe	ensated
					Serum
Uric Acid	5.7	mg/dL	3.4 - 7.0	Uricase / Peroxidas (Colorimetric)	e
					Serum
Fasting Glucose As per ADA Guideline Normal : Less than 100 mg/dL Prediabetes : 100 mg/dL to 125 mg/dL Diabetes : 126 mg/dL or Higher	86.4	mg/dL	< 100.0	Hexokinase	
					Serum
Post prandial Glucose Post prandial Urine Glucose	61.3 S.N.G. = San	mg/dL nple Not Given	< 140.0	Hexokinase	
				Actor	<u>~</u>
			М.	. Kazumi Gondalia D (Path) g.No.: G-21729	

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L-Low H-High CH -Critical High CL - Critical Low

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As per ADA Guideline Normal : Less than 100 mg/dL Prediabetes : 100 mg/dL to 125 mg/dL Diabetes : 126 mg/dL or Higher

***LIVER FUNCTION TEST (LFT) SERUM**

SGPT(ALT)		18.00	U/L	0.00 - 41.00	IFCC without pyridoxal phosphate	
SGOT(AST)		18.60	U/L	0.00 - 40.00	IFCC without pyridoxal phosphate	
Alkaline Phosphatase		103.5	U/L	40.0 - 129.0	PNP-Standardize	
Bilirubin Total		0.89	mg/dL	0.00 - 1.10	Diazo Method	
Bilirubin Direct	н	0.32	mg/dL	0.00 - 0.20	Diazo Method	
Bilirubin Indirect		0.57	mg/dL	0.00 - 1.10	Calculate from Total and Direct Billirubin	I
Protein Total		7.55	g/dL	6.40 - 8.20	Biuret Method	
Albumin		4.94	g/dL	3.97 - 4.95	BCG Endpoint	
Globulin		2.61	g/dL	2.20 - 3.50	Calculated	
A/G Ratio		1.89	Ratio	0.90 - 2.80	Ratio	
						EDTA Blood
						LDTA Blood
HbA1c (Glyco Hb)		5.10	%	4.8 % - 5.9 % Normal 5.9 % - 7.0 % Good diabetic Control 7.0 % - 10.00 % Fair Diabetic Control >10.0 % Poor diabetic Control	Immunoturbidimetric	
Mean Plasma Glucose		106.0	mg/dL	80.0 - 140.0		

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Serum

Serum

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Blood Urea		16.6	mg/dL	16.6 - 48.5	Urease,Kinetic,GLDH		
BUN*		7.8	mg/dL	6.0 - 20.0	Ureas with UV		
						Serum	
TOTAL T3*		1.400	ng/mL	0.850 - 2.020	ECLIA.		
TOTAL T4*		7.310	ug/dL	5.130 - 14.060	ECLIA.		
THYROID STIMUL	ATING HORMONE	3.960	ulU/mL	0.270 - 4.200	ECLIA.		

(*) Not in NABL Scope

End Of Report

Lepon

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		Sample Quality	

Test Name	Result	Unit	Bio. Ref. Range	Method	Sample
	ArcoFemi I	Biochemistry Healthcare Ltd I			
*LIPID PROFILE (WITH DIRECT LDL					Serum
Sample Type	Fasting				
Cholesterol Total	168.50	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	62.70	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	GPO-PAP	
HDL Cholesterol	40.40	mg/dL	Less than 40 mg/dL Low 60 mg/dL or Above Excellent	Homogenous Enzy	ymatic
LDL Cholesterol (Direct)	122.50	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 Enzy	/matic
VLDL Cholesterol	12.54	mg/dL	< 30	Calculated	
LDL/HDL RATIO	3.03		< 3.50	Calculated	
Cholesterol Total / HDL Ratio	4.17		< 4.50		
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		Sample Quality	

(*) Not in NABL Scope

**End Of Report**

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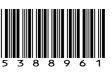
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Bed No/Ward	OPD	Report Date	19/03/2024 3:32PM
Referred By	Dr. Casualty Medical Officer	Report Status	Final
		Sample Quality	

Test Name	Result	Unit	Bio. Ref. Range	Method	Sample
	Clinic	al Pathol	ogy		
	ArcoFemi Health	care Ltd ]	Below 40 Male		
<b>*URINE ROUTINE EXAMINATION</b>					Urine
Physical Examination:					
Quantity	20 ml			Visual method	
Colour	Pale Yellow			Visual method	
Appearence:	Clear			Visual method	
Reaction	6.5		4.5 - 8.0	Reflectance ph	otometer
Sp. Gravity	1.010		1.015 - 1.030	Reflectance ph reaction	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				
(*) Not in NABL Scope	**End Of Report	**			

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## DEPARTMENT OF LABORATORY SERVICES

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Age/Gender	38 Years/Male	Receiving Date	19/03/2024 2:52PM
Bed No/Ward	OPD	Report Date	19/03/2024 5:28PM
Referred By	Dr. Casualty Medical Officer	Report Status	Final
		Sample Quality	

Test Name		Result	Unit	Bio. Ref. Range	Method	Sample
			Haematology			
		ArcoFemi I	Healthcare Ltd Be	elow 40 Male		
<u>*CBC WITH ESR</u>						EDTA Blood
Haemoglobin		16.2	g/dL	13.5 - 18.0	SLS Method	
Hematocrit/PCV		47.6	g/u∟ %	42.0 - 52.0	H.focusing Met	hod
			mill/Cmm		0	
RBC COUNT MCV		4.93 96.6		4.70 - 6.00 83.0 - 101.0	H.focusing imp Calculated	edance
			fl		Calculated	
MCH	н	32.9	pg	27.0 - 31.0		
MCHC		34.0	g/dL	32.0 - 36.0	Calculated	
RDW-CV		13.8	%	11.5 - 14.0	Calculated	
Platelet count		285000	/cumm	150000 - 410000	H.focusing imp	edance
Mean Platelet Volume(MPV)*		9.6	fl	8 - 12	Calculated	
Total Leucocyte Count (TLC)		6840.00	/cumm	4000.00 - 10500.00	Flow Cytometry	
Differential Leucocyte Count					Flowcytometry/	Microscopic
Neutrophils		46	%	40 - 70		
Lymphocytes		43	%	22 - 45		
Eosinophils	н	07	%	1 - 4		
Monocytes		04	%	1 - 6		
Basophils		00	%	0 - 1		
Absolute Leucocyte Count						
Absolute Neutrophil Count*		3146.4	/cumm	1800 - 7700		
Absolute Lymphocyte count*		2941.2	/cumm	1000 - 4800		
Absolute Eosinophil Count (AEC)	н	478.8	/cumm	0.0 - 450.0		
Absolute Monocyte Count*		273.6	/cumm	0 - 800		
Peripheral Smear Study		Platelets are Malarial Para	rmochromic & No adequate in numl sites are not seer e cells are seen.	per.		

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stopped flow kinetic

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Erythrocyte Sedin	nentation Rate (ESR)	05 mr	n/hr 0 - 10	Photometric capillary	

(*) Not in NABL Scope

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		Sample Quality	

Test Name	Result	Unit	Bio. Ref. Range	Method	Sample
		<b>muno-Haema</b> Healthcare Ltd	tology Below 40 Male		
<u>*BLOOD GROUPING</u>					EDTA Blood
ABO Group	"B"		Tube Agglutination Method		
Rh Type	Positive				
(*) Not in NABL Scope	**End Of I	Report**			

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