

External Visit Id

PO BOX 214-545, S-G09 / S-G10 , Al Garhoud Business Centre , Al Garhoud , Dubai, UAE.

Toll Free: 800 325 222

Email: info@fortecl.ae | Web: www.fortecl.ae

# CONFIDENTIAL LABORATORY REPORT

: 46566

Patient Name : Ms. MINA MESTOURI : 21/04/2025 21:13 Request Date Age / Sex : 24 Y / Female **Collected Date** : 21/04/2025 21:10 DOB : 22/10/2000 : 21/04/2025 22:16 Acceptance Date Referral Doctor Report Date : 22/04/2025 21:47

: CITICARE MEDICAL CENTER L.L.C Referrer : Final Report Report Status

Patient No : FCL020331

SID.No.

FD020231

INVESTIGATION / SPECIMEN	RESULT	UNIT	REFERENCE RANGE	METHOD
HAEMATOLOGY CITICARE PULSEFIT PACKAGE COMPLETE BLOOD COUNT (CBC)				
EDTA Whole Blood HEMOGLOBIN	13.2	g/dL	12 - 15.5	Colorimetric
RBC COUNT	4.72	10^6/uL	3.5 - 5.5	EI
HEMATOCRIT	40.4	%	37 - 54	Calculation
MCV	85.5	fL	80 - 100	Calculation
MCH	28	pg	27 - 34	Calculation
MCHC	32.7	g/dL	32 - 36	Calculation
RDW	13.2	%	11 - 16	Calculation
MPV	13.6	fL	8 - 12	Calculation
PLATELET COUNT	173	10^3 /uL	140 - 450	EI
WBC COUNT	7.47	10^3 /uL	4 - 10	EI
DIFFERENTIAL COUNT (DC)				
EDTA Whole Blood NEUTROPHILS	59.6	%	50 - 70	Flow Cytometry
LYMPHOCYTES	34.5	%	20 - 40	Flow Cytometry
EOSINOPHILS	0.9	%	0.5 - 5	Flow Cytometry
MONOCYTES	4.7	%	3 - 12	Flow Cytometry
BASOPHILS	0.3	%	0 - 1	Flow Cytometry
ABSOLUTE NEUTROPHIL COUNT	4.45	10^3 cells/uL	2 - 7	
ABSOLUTE LYMPHOCYTE COUNT	2.58	10^3 cells/uL	0.8 - 4	
ABSOLUTE MONOCYTE COUNT	0.35	10^3 cells/uL	0.12 - 1.2	
ABSOLUTE EOSINOPHIL COUNT	0.07	10^3 cells/uL	0.02 - 0.5	
ABSOLUTE BASOPHIL COUNT	0.02	10^3 cells/uL	0 - 0.1	

**Prepared By** 

Inayatulla Ibrahim Dhanse **General Lab Technologist** DGAP022039



Approved By

Dr Anjum Nawshehri **Laboratory Director** DHA-00208333-004

Page 1 of 8



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INVESTIGATION / SPECIMEN	RESULT	UNIT	REFERENCE RANGE	METHOD
CLINICAL BIOCHEMISTRY CITICARE PULSEFIT PACKAGE				
GLUCOSE (RANDOM)	80.3	mg/dL	< 200	GOD - POD
Fluoride Plasma GLYCATED HEMOGLOBIN (HbA1C) ^				
EDTA Whole Blood				
HBA1C	5.0	%	Normal:4.6-5.6 Prediabetes: 5.7- 6.4 Diabetes: >6.5	HPLC
eAG (estimated Average Glucose)	97	%		Calculation
BLOOD UREA NITROGEN (SERUM)				
Serum				
UREA	22.5	mg/dL	16.87 - 43.37	Urease-glutamate Dehydrogenase, UV method
BUN	10.5	mg/dL	7 - 18	Calculation
CREATININE	0.51	mg/dL	0.5 - 1.1	Sarcosine Oxidase
Serum				
BUN/CREATININE RATIO	19	NULL	Prerenal: > 20:1 Normal or Postrenal: 10-20:1 Intrarenal: < 10:1	Calculation
Serum				
URIC ACID	3.46	mg/dL	1.9 - 7.5	Uricase - POD
Serum IRON	71.5	ug/dL	37 - 145	Ferrozine
Serum	/1.5	ug/uL	37 - 143	rerrozine
CHOLESTEROL (TOTAL)	165.6	mg/dL	Desirable: < 200 Borderline High: 200 - 239 High: > 239	CHOD-POD
Serum NON-HDL CHOLESTEROL	96	mg/dL	< 140	Calculation
Serum	90	mg/uL	< 1 <del>11</del> 0	Calculation
Scruiii				

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Page 2 of 8

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INVESTIGATION / SPECIMEN	RESULT	UNIT	REFERENCE RANGE	METHOD
CLINICAL BIOCHEMISTRY				
CITICARE PULSEFIT PACKAGE				
TRIGLYCERIDES	106.6	mg/dl	Normal: < 150 Borderline High: 150 - 199 High: 200 - 499 Very High: > 499	GPO-POD
Serum				
HDL CHOLESTEROL	69.9	mg/dL	Optimum>60 Borderline : 50-59 High risk : <50	Direct
Serum			•	
LDL CHOLESTEROL	89.6	mg/dl	Optimal: < 100 Near/Above Optimal: 100 - 129 Borderline High: 130 - 159 High: 160 - 189 Very High: > 189	Direct
Serum				
VLDL CHOLESTEROL	21.32	mg/dL	0 - 30	Calculation
Serum				
LDL / HDL RATIO	1.3		< 3.5	Calculation
Serum				

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Page 3 of 8



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INVESTIGATION / SPECIMEN	RESULT	UNIT	REFERENCE RANGE	METHOD
CLINICAL BIOCHEMISTRY				
CITICARE PULSEFIT PACKAGE				
TOTAL CHOLESTEROL / HDL RATIO	2.4		< 4.5	Calculation
Serum				

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Page 4 of 8



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INVESTIGATION / SPECIMEN	RESULT	UNIT	REFERENCE RANGE	METHOD
CLINICAL BIOCHEMISTRY				
CITICARE PULSEFIT PACKAGE BILIRUBIN (TOTAL)	0.34	mg/dL	Up to 1.2	DSA
Serum BILIRUBIN (DIRECT)	0.11	mg/dL	0.00 - 0.4	DSA
Serum INDIRECT BILIRUBIN	0.23	mg/dL	< or = 0.90	Calculation
Serum TOTAL PROTEIN	6.2	g/dL	6 - 8	Biuret
Serum ALBUMIN (SERUM)	4.1	g/dL	3.5 - 4.8	Bromcresol green (BCG)
Serum GLOBULIN	2.1	g/dL	2.0 - 3.5	Calculation
Serum ALBUMIN / GLOBULIN RATIO	1.95	NULL	0.8 - 2.0	Calculation
Serum ALT / SGPT	18	U/L	10 - 40	IFCC
Serum ALP (ALKALINE PHOSPHATASE)	82.7	U/L	30 - 115	IFCC
Serum GGT (GAMMA GLUTAMYL TRANSFERASE)	20.4	U/L	7 - 50	IFCC
Serum AST / SGOT	19.8	U/L	10 - 40	IFCC
Serum				

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Page 5 of 8



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SID.No. : **FD020231** 

Patient No : FCL020331

INVESTIGATION / SPECIMEN	RESULT	UNIT	REFERENCE RANGE	METHOD
ENDOCRINOLOGY				
CITICARE PULSEFIT PACKAGE				
TRIIODOTHYRONINE, TOTAL (T3)	1.2	ng/mL	0.8 - 1.9	CLIA
Serum				
THYROXINE, TOTAL (T4)	7.1	ug/dL	5 - 13	CLIA
Serum				
THYROID STIMULATING HORMONE (TSH)	0.973	μIU/mL	0.35 - 4.75	CLIA
Serum				
FOLLICLE STIMULATING HORMONE (FSH) Serum	7.507	mIU/mL	Follicular phase: 2.50 - 11.40 Midcycle Peak: 3.3 - 21.7 Luteal Phase: 1.2 - 7.0 Postmenopausal: 18.8 - 132.0	CLIA
LUTEINISING HORMONE (LH)	9.739	mIU/mL	Follicular phase: 1.2 - 12.7 Midcycle Peak: 15.2 - 90.0 Luteal Phase: 0.5 - 14.6 Postmenopausal: 15.6 - 72.0	CLIA
Serum				
PROLACTIN	567.91	uIU/mL	51 - 580	CLIA
Serum				
TESTOSTERONE (TOTAL)	0.17	ng/mL	0.1 - 0.9	CLIA
Serum				
TESTOSTERONE (FREE)	0.003	ng/mL	0.07-0.3	Calculation
Serum SHBG (SEX HORMONE BINDING GLOBULIN)	36.09	nmol/L	18.2 - 135.5	CLIA
Serum				
VITAMIN B12	9033	pg/mL	180 - 916	CLIA
Serum				
omments: Test repeated twice & confirmed. Co relate		ng/ml	Deficient : < 20	CLIA
VITAMIN D, 25-OH (TOTAL)	34	ng/mL	Insufficient: < 20 Insufficient: 20 - < 30 Sufficient: 30 - 100 Upper Safety Limit:: > 100	CLIA
O FLIDO				

Serum

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Page 6 of 8

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INVESTIGATION / SPECIMEN	RESULT	UNIT	REFERENCE RANGE	METHOD
ENDOCRINOLOGY CITICARE PULSEFIT PACKAGE				
BETA HCG	<2	mIU/mL	Non-Pregnant Females: <2.9 Week after LMP 1-10 weeks 195 - 229796 11-15 weeks 22305 - 233036 16-22 weeks 7902 - 49645 23-40 weeks 1580 - 48608	CLIA

Serum

**Prepared By** 

Juhana Pullichola Microbiology Technologist 17945483



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Page 7 of 8

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INVESTIGATION / SPECIMEN	RESULT	UNIT	REFERENCE RANGE	METHOD
ENDOCRINOLOGY				
CITICARE PULSEFIT PACKAGE				
ESTRADIOL (E2)	43.1	pg/mL	Follicular Phase: 30 - 150 Mid Cycle Peak: 60 - 480 Luteal Phase: 45 - 250 Post Menopausal: < 60.0	CLIA
Serum				
PROGESTERONE	0.255	ng/mL	Follicular Phase: 0.00 - 1.40 Luteal Phase: 3.34 - 25.56 Mid luteal phase: 4.44 - 28.03 Post Menopausal: 0.00 - 0.73 First trimester: 11.22 - 90.0 Second trimester: 25.55 - 89.40 Third trimester: 48.40 - 422.50	CLIA

#### Serum

Interpretation Notes: A progesterone test may be used: To help recognize and manage some causes of infertility. Since progesterone levels vary throughout the menstrual cycle, multiple (serial) measurements can be used for this purpose. To determine whether or not a woman has ovulated, when ovulation occurred, or to monitor the success of induced ovulation learly pregnancy to help diagnose an ectopic or failing pregnancy, along with human chorionic gonadotropin (hCG) testing To monitor a high-risk pregnancy to help evaluate placenta and fetal healthlif a woman is receiving progesterone injections to help support her early pregnancy, to help determine the effectiveness of the replacement treatmentAlong with other tests such as an FSH, LH, hCG, thyroid tests, clotting tests, and a complete blood count (CBC) to help determine the cause of abnormal uterine bleeding in non-pregnant women

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Page 8 of 8

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