

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

 Patient Name
 : Ms. OLENA
 Request Date
 : 25/04/2025 15:04

 Age / Sex
 : 25 Y / Female
 Collected Date
 : 25/04/2025 15:03

 DOB
 : 13/01/2000
 Acceptance Date
 : 25/04/2025 15:17

 Referral Doctor
 : Report Date
 : 25/04/2025 22:16

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

SID.No. : **FD020699** External Visit Id : **45025**

Patient No : FCL020805

INVESTIGATION / SPECIMEN	RESULT	UNIT	REFERENCE RANGE	METHOD
CLINICAL BIOCHEMISTRY				
CITICARE PULSEFIT PACKAGE				
GLUCOSE (FASTING)	91	mg/dL	70 - 110	GOD - POD
Fluoride Plasma GLYCATED HEMOGLOBIN (HbA1C) ^				
EDTA Whole Blood				
HBA1C	5.4	%	Normal:4.6-5.6 Prediabetes: 5.7- 6.4 Diabetes: >6.5	HPLC
eAG (estimated Average Glucose)	108	%		Calculation
BLOOD UREA NITROGEN (SERUM)				
Serum				
UREA	18.6	mg/dL	16.87 - 43.37	Urease-glutamate Dehydrogenase, UV method
BUN	8.7	mg/dL	7 - 18	Calculation
CREATININE	0.99	mg/dL	0.5 - 1.1	Sarcosine Oxidase
Serum				
BUN/CREATININE RATIO	9.0	NULL	Prerenal: > 20:1 Normal or Postrenal: 10-20:1 Intrarenal: < 10:1	Calculation
Serum				
URIC ACID	5.25	mg/dL	1.9 - 7.5	Uricase - POD
Serum				
IRON	136.2	ug/dL	37 - 145	Ferrozine
Serum				
CHOLESTEROL (TOTAL)	216.1	mg/dL	Desirable: < 200 Borderline High: 200 - 239 High: > 239	CHOD-POD
Serum				
NON-HDL CHOLESTEROL	127	mg/dL	< 140	Calculation
Serum				

Prepared By

Inayatulla Ibrahim Dhanse General Lab Technologist DGAP022039



Approved By

Dr Anjum Nawshehri Laboratory Director DHA-00208333-004

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CLINICAL BIOCHEMISTRY				
CITICARE PULSEFIT PACKAGE				
TRIGLYCERIDES	37.2	mg/dl	Normal: < 150 Borderline High: 150 - 199 High: 200 - 499 Very High: > 499	GPO-POD
Serum			, -	
HDL CHOLESTEROL	89.4	mg/dL	Optimum>60 Borderline : 50-59 High risk : <50	Direct
Serum			3	
LDL CHOLESTEROL	123.6	mg/dl	Optimal: < 100 Near/Above Optimal: 100 - 129 Borderline High: 130 - 159 High: 160 - 189 Very High: > 189	Direct
Serum				
VLDL CHOLESTEROL	7.44	mg/dL	0 - 30	Calculation
Serum LDL / HDL RATIO	1.4		< 3.5	Calculation
Serum				

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CLINICAL BIOCHEMISTRY				
CITICARE PULSEFIT PACKAGE				
TOTAL CHOLESTEROL / HDL RATIO	2.4		< 4.5	Calculation
Serum				

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CLINICAL BIOCHEMISTRY				
CITICARE PULSEFIT PACKAGE BILIRUBIN (TOTAL)	1.66	mg/dL	Up to 1.2	DSA
Serum BILIRUBIN (DIRECT)	0.3	mg/dL	0.00 - 0.4	DSA
Serum INDIRECT BILIRUBIN	1.36	mg/dL	< or = 0.90	Calculation
Serum TOTAL PROTEIN	8.0	g/dL	6 - 8	Biuret
Serum ALBUMIN (SERUM)	4.4	g/dL	3.5 - 4.8	Bromcresol green (BCG)
Serum GLOBULIN	3.6	g/dL	2.0 - 3.5	Calculation
Serum ALBUMIN / GLOBULIN RATIO	1.22	NULL	0.8 - 2.0	Calculation
Serum ALT / SGPT	10.8	U/L	10 - 40	IFCC
Serum ALP (ALKALINE PHOSPHATASE)	64.4	U/L	30 - 115	IFCC
Serum GGT (GAMMA GLUTAMYL TRANSFERASE)	20.1	U/L	7 - 50	IFCC
Serum AST / SGOT	20.7	U/L	10 - 40	IFCC
Serum				

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ENDOCRINOLOGY				
CITICARE PULSEFIT PACKAGE TRIIODOTHYRONINE, TOTAL (T3)	1.19	ng/mL	0.8 - 1.9	CLIA
Serum THYROXINE, TOTAL (T4)	7.3	ug/dL	5 - 13	CLIA
Serum THYROID STIMULATING HORMONE (TSH)	0.884	μIU/mL	0.35 - 4.75	CLIA
Serum FOLLICLE STIMULATING HORMONE (FSH)	7.011	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
Serum LUTEINISING HORMONE (LH)	6.917	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	227 774	TI I / I		CLTA
PROLACTIN Serum	227.771	uIU/mL	51 - 580	CLIA
TESTOSTERONE (TOTAL)	0.25	ng/mL	0.1 - 0.9	CLIA
Serum TESTOSTERONE (FREE)	0.00357	ng/mL	0.0005 - 0.002	Calculation
Serum SHBG (SEX HORMONE BINDING GLOBULIN)	67.26	nmol/L	18.2 - 135.5	CLIA
Serum VITAMIN B12	478.369	pg/mL	164 - 905	CLIA
Serum VITAMIN D, 25-OH (TOTAL)	50	ng/mL	Deficient: < 20 Insufficient: 20 - 30 Sufficient: 30 - 100 Upper Safety Limit:: > 100	CLIA
Serum				

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ENDOCRINOLOGY CITICARE PULSEFIT PACKAGE BETA HCG 2.0 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	INVESTIGATION / SPECIMEN	RESULT	UNIT	REFERENCE RANGE	METHOD
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	BETA HCG	2.0	mIU/mL	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 -	CLIA

Serum

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ENDOCRINOLOGY				
CITICARE PULSEFIT PACKAGE				
ESTRADIOL (E2)	67.7	pg/mL	Follicular Phase: 30 - 150 Mid Cycle Peak: 60 - 480 Luteal Phase: 45 - 250 Post Menopausal: < 60.0	CLIA
Serum				
PROGESTERONE	0.089	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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