

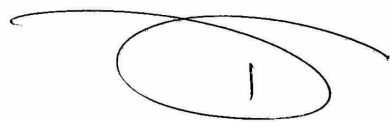
Name : SHIMAA ALSAYED
MRN : 860727-0
Gender : Female
Age / DOB : 33 Y / 14-06-1989
Client Name : IRHAM MEDICAL CENTER
Nationality : Egypt

Lab ID : 1470234
Sample No : 22110800021-
Emirates / Passport ID :
Reg Date : 13-11-2022 23:12:49
Collection Date : 13-11-2022 21:10:00
Reporting Date : 14-11-2022 09:54:05

Hormones Report

Test	Result	Unit	Reference Range	Methodology
TSH, Serum	3.157	uIU/mL	Non-pregnant females: 0.38 - 5.33 Pregnant, 1st Trimester: 0.05 - 3.70 Pregnant, 2nd Trimester: 0.31 - 4.35 Pregnant, 3rd Trimester: 0.41 - 5.18	CLIA

End of Report



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 Lab Director
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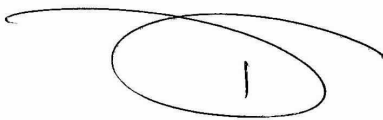


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Immunology Report

Test	Result	Unit	Reference Range	Methodology
HIV 1&2 Antibody and P24 Antigen, Serum	Non Reactive	S/CO	Non Reactive	CLIA
<i>The test is a chemiluminescent immunoassay for the qualitative detection of HIV-1 p24 antigen and antibodies to HIV-1 (groups M and O) and HIV-2 in human serum and plasma using the Access Immunoassay Systems. The Access HIV combo assay is intended to be used as an aid in the diagnosis of HIV-1 or HIV-2 infection and as screening test for blood and plasma donors.</i>				
Hepatitis B Surface Antigen, Serum	Non Reactive	S/CO	Non Reactive	CLIA
HCV antibody, Serum	Non Reactive	S/CO	Non Reactive	CLIA
Rubella Ab, IgG, Serum	97.16 Reactive	IU/mL	<10 IU/mL : Non-Reactive >=15 IU/mL : Reactive >10 to <15 IU/mL : Equivocal	CLIA
CMV Antibodies, IgG	391.3 Reactive	AU/ML	Non-Reactive: <11 Equivocal: 11 - 15 Reactive: >15	CLIA
Syphilis Rapid Plasma Reagin; VDRL / RPR , Qualitative	Non Reactive		Non Reactive	Agglutination
CMV Antibody IgM, Serum	Negative		Negative	ELISA
Remarks: <i>The Test is intended as an aid in the qualitative determination of IgM class antibodies against Cytomegalovirus in human serum. All the values obtained require a careful interpretation that must consider other indicators relative to the patient. Absence of CMV specific IgM does not exclude the possibility of CMV infection. It has been reported that 10-30% of infants may fail to develop CMV IgM antibody response despite congenital infection with CMV. The test can not be used alone for a clinical diagnosis and the should be evaluated together with the patient history and other clinical diagnostic evaluation.</i>				

End of Report



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Name : SHIMAA ALSAYED

MRN : 860727-0

Gender : Female

Age / DOB : 33 Y / 14-06-1989

Client Name : IRHAM MEDICAL CENTER

Nationality : Egypt

Lab ID : 1470234

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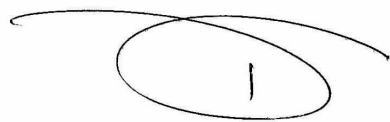
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Chemistry Report

Test	Result	Unit	Reference Range	Methodology
Ferritin, Serum	5.4	L ng/mL	11.0 - 307.0	CLIA
Glucose, Random, Serum	82	mg/dL	Up to 200	Enzymatic Colorimetric

End of Report



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Name :	SHIMAA ALSAYED	Lab ID :	1470234
MRN :	860727-0	Sample No :	22110800021-
Gender :	Female	Emirates / Passport ID :	
Age / DOB :	33 Y / 14-06-1989	Reg Date :	13-11-2022 23:12:49
Client Name :	IRHAM MEDICAL CENTER	Collection Date :	13-11-2022 21:10:00
Nationality :	Egypt	Reporting Date :	14-11-2022 09:54:05

Urine Chex

Test	Result	Unit	Reference Range	Methodology
<u>Urine Macroscopic Examination</u>				
Pus Cells	2-3	/HPF	Up to 3.0	
R.B.Cs	0-2	/HPF	Up to 2.0	
Epithelial Cells	12-14	/HPF	Up to 10.0	
Bacteria	Seen (Few)	/HPF	Not seen/Few	
Casts	Not seen	/HPF	Not seen	
Others	Mucus threads (Few)	/HPF	Not seen	
Crystal	Not seen	/HPF	Not seen	
<u>Urine Chemistry Examination</u>				
Specific Gravity	1.010		1.005	1.030
PH	6.0		5.0 -	7.5
Glucose	Negative		Negative	
Ketones	Negative		Negative	
Protein	Negative		Negative	
Blood	Negative		Negative	
Bilirubin	Negative		Negative	
Urobilinogen	Normal		Normal	
Nitrite	Negative		Negative	
Leucocytes	Negative		Negative	
<u>Urine Microscopic Examination</u>				
Colour	Yellow		Yellow	Visual
Appearance	Trace Turbidity		Clear	Visual

End of Report



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Sample Type: Urine



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Final Report



Name : SHIMAA ALSAYED

Lab ID : 1470234

MRN : 860727-0

Sample No : 22110800021-

Gender : Female

Emirates / Passport ID :

Age / DOB : 33 Y / 14-06-1989

Reg Date : 13-11-2022 23:12:49

Client Name : IRHAM MEDICAL CENTER

Collection Date : 13-11-2022 21:10:00

Nationality : Egypt

Reporting Date : 14-11-2022 09:54:05

Complete Blood Count (CBC), Whole Blood

Test	Result	Unit	Reference Range	Methodology
WBC	5.5	10 ³ /uL	3.8 - 11.0	Impedance
RBC	4.1	10 ⁶ /uL	3.8 - 5.1	Impedance
Hemoglobin	11.3 L	g/dL	11.6 - 15.1	Photometric
Hematocrit	33.9 L	%	34.6 - 44.1	Calculated
MCV	81.9	fL	80.0 - 98.0	Calculated
MCH	27.4	pg	26.6 - 33.5	Calculated
MCHC	33.5	g/dL	32.9 - 35.4	Calculated
RDW	19.7 H	%	12.6 - 15.6	Calculated
Platelet	174	10 ³ /uL	150 - 360	Impedance

Leukocytes Differential Count

Differential Percentage				Absolute Count			
	Value	Unit	Normal Range	Value	Unit	Normal Range	
Lymphocytes	35.79	%	18.51 - 48.98	1.95	10 ³ /uL	1.16 - 3.78	
Monocytes	5.23	%	4.72 - 11.35	0.29	10 ³ /uL	0.28 - 0.83	
Neutrophils	58.44	%	41.35 - 72.27	3.19	10 ³ /uL	2.03 - 7.67	
Eosinophils	0.19 L	%	0.71 - 6.09	0.01 L	10 ³ /uL	0.04 - 0.42	
Basophils	0.35	%	0.05 - 0.47	0.02	10 ³ /uL	Up to 0.03	

Sample Type / ID : EDTA / 22110300054

End of Report



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Sample Type: EDTA

Final Report