



5151 CORPORATE WAY
 JUPITER, FL 33458-3101
 (866)720-8386

Client: ACCESS MEDICAL LABORATORI 1 5151 CORPORATE WAY JUPITER, FL 33458	Patient: SAMPLE 6, TEST Room# _____ DOB. 01/01/1970 Age:47 Sex:F Phone: () - _____ Fasting: N ID#: A1710001391470 Route#: 0
Phys: (561) 745-1233	Page:2

Acc# 001391470	Coll. Date: 10/23/17	Recv. Date: 10/23/17	Print Date: 02/22/19
Chart#	Coll. Time: 00:00 AM	Recv. Time:05:50 PM	Print Time: 13:50
First reported on:	10/24/17	Final report date:	10/24/17

Test Name	Results	Normal Range	Units
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GENERAL CHEMISTRY (Continued)

			in GFR
Stage 3	30 to 59 ml/min		Moderate decrease in GFR
Stage 4	15 to 29 ml/min		Severe decrease in GFR
Stage 5	< 15 ml/min		Kidney failure, or on dialysis

IRON/ANEMIA EVALUATION

FERRITIN	51	13 - 300	ng/ml
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CORONARY RISK

TRIGLYCERIDES	100	<150	mg/dl
CHOLESTEROL, TOTAL	198	<200	mg/dl
HDL CHOLESTEROL	55	>40	mg/dl
LDL CHOLESTEROL, calc.	123 H	<100	mg/dl
CHOL/HDL RATIO	3.6	<4.4	

The higher the Ratio, the higher CHD risk.

CRP, Cardio	31 H	<3.0	mg/L
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****Risk of Cardiovascular Disease****

Low Risk	CRP < 1.0	mg/L
Medium Risk	CRP 1.0 - 3.0	mg/L
High Risk	CRP > 3.0	mg/L

THYROID TESTING

T3, FREE	35 H	1.8 - 4.6	pg/ml
T4, FREE	12 H	0.9 - 1.7	ng/dl
TSH	16 H	0.27 - 4.2	uIU/ml

ENDOCRINE EVALUATION

FSH	35	1.7 - 134.8	mIU/ml
Men		1.5 - 12.4	mIU/ml
Women			
Follicular Phase		3.5 - 12.5	,,
Ovulation Phase		4.7 - 21.5	,,
Luteal Phase		1.7 - 7.7	,,
Postmenopause		25.8 - 134.8	,,

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ENDOCRINE EVALUATION (Continued)

LH 15 1.0 - 95.6 mIU/ml

Men 1.7 - 8.6 mIU/ml
 Women
 Follicular Phase 2.4 - 12.6 ,,
 Ovulation Phase 14.0 - 95.6 ,,
 Luteal Phase 1.0 - 11.4 ,,
 Postmenopause 7.7 - 58.5 ,,

PROGESTERONE 23 ng/mL

****Women Reference Ranges****

Follicular Phase 0.057 - 0.893 ng/mL
 Ovulation Phase 0.121 - 12.0 ng/mL
 Luteal Phase 1.83 - 23.9 ng/mL
 Postmenopause <0.05 - 0.126 ng/mL

****Pregnant****

1st trimester 11.0 - 44.3 ng/mL
 2nd trimester 25.4 - 83.3 ng/mL
 3rd trimester 58.7 - 214 ng/mL

As of 09/23/16 Access Medical Labs will be reporting Progesterone using an updated Progesterone Generation-III reagent from Roche. Progesterone-III reduces cross-reactivity, shows higher specificity toward Progesterone and has been traced to the LC/MS reference method. Although results may vary from the previous method, these changes are taken into account in the updated reference ranges.

ESTRADIOL (E2) 15 pg/ml

****Women Reference Ranges****

Follicular Phase 12.5 - 166 pg/ml
 Ovulation Phase 85.8 - 498 pg/ml
 Luteal Phase 43.8 - 211 pg/ml
 Postmenopause <5.0 - 54.7 pg/ml

Pregnancy

1st trimester 215 - >4300pg/ml

Fulvestrant (Faslodex)Cross Reactivity: Fulvestrant therapy will cause false elevated Estradiol Result with this method. Use LC/MS method if patient is on Fulvestrant to avoid Cross reactivity.

DHEA-SULFATE **35 L** 35.4 - 256.0 ug/dl

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ENDOCRINE EVALUATION (Continued)

TESTOSTERONE, TOTAL	3514 H	6 - 82	ng/dl
SEX HORMONE BIND GLOBULIN	135 H	20 - 130	nmol/L
TESTOSTERONE, FREE	47.56 H	0.2 - 2.6	ng/dl
IGF-1	53 L	119 - 166	ng/mL

Access Medical Laboratories uses Siemens Healthcare Diagnostics as the supplier for IGF-1 Immunoassay Testing System. Siemens Introduced a Restandardization of IGF-1 assay using WHO 1st International Standard (IS), NIBSC Code 02/254.

Starting 01/25/2017 we will be implementing the new restandardized IGF-1 assay for testing Patients. New Reference Ranges will reflect expected lower values for patient IGF-1 results.

END OF REPORT