Name: BLAYK,BONZE ANNE ROSE DOB: 05/01/1956 Attend Dr: Adam Law MD
Acct: A00088103734 Unit: M000597460 AGE: 62 Location: LAB
Reg: 08/03/18 SEX: F Status: REG REF

Test	Day	Date	Time	Result	Reference	Units
D Dimer Quant	1	AUG 3	1717	< 200(A)	(Less Than 230)	ng/mL

(A) **Please note:
The following may produce a false positive D Dimer test:
- Rheumatoid factor greater than 60 IU/ml
- Plasma hemoglobin greater than 0.05 gm/dl
- Bilirubin greater than 50 mg/dl
- Lipids greater than 1000 mg/dl
- FDP greater than 20 ug/ml

Test	Day	Date	Time	Result	Reference	Units
FSH	1	AUG 3	1717	16.8(B)		mIU/mL

(B) Normally menstruating females
- Follicular phase 3 - 9
- Mid-cycle peak 4 - 23
- Luteal phase 1 - 6
Postmenopausal females 16 - 114

Test	Day	Date	Time	Result	Reference	Units
LH	1	AUG 3	1717	7.6(C)		mcIU/mL

(C) Normally menstruating females
- Follicular Phase 1 - 18
- Mid-Cycle Peak 24 - 105
- Luteal Phase 0.6 - 20
Postmenopausal females 15 - 62

Test	Day	Date	Time	Result	Reference	Units
Prolactin	1	AUG 3	1717	4.4	(1.0-25.0)	ng/mL
Estradiol	1	AUG 3	1717	<40(D)		pg/mL

(D) Estradiols <40 pg/mL are sent to a reference lab for low range testing.
See also (E)

(E) Postmenopausal Females < 20
Ovulating females: by day in cycle relative to LH Peak

Follicular phase	- 12	10-50
	- 4	60-200
Mid-cycle	- 1	120-375
Luteal phase	+ 2	50-155
	+ 6	60-260
	+ 12	15-115

** CONTINUED ON NEXT PAGE **

Patient: BLAYK,BONZE ANNE ROSE A00088103734 (Continued)

Test	Day	Date	Time	Result	Reference	Units
Estradiol, MS	1	AUG 3	1717	19(F)		

(F) ----------REFERENCE VALUE----------
Premenopausal: 15-350 (E2 levels vary widely through the menstrual cycle.)
Postmenopausal: <10
----------ADDITIONAL INFORMATION----------
This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.
Test Performed by:
Mayo Clinic Laboratories - Rochester Superior Drive
3050 Superior Drive NW, Rochester, MN 55901
See also (*G)
(*G) Mayo Medical Laboratories
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Test	Day	Date	Time	Result	Reference	Units
Progesterone	1	AUG 3	1717	0.1(H)		ng/mL

(H) ~Female reference ranges for Progesterone:
Follicular phase.....0.3 - 1.5 ng/ml
Mid-luteal phase.....5.2 - 18.5 ng/ml
Postmenopausal.....< 0.8 ng/ml
Pregnant
1st trimester.....4.7 - 50.0 ng/ml
2nd trimester.....19.4 - 45.3 ng/ml

Test	Day	Date	Time	Result	Reference	Units
Free Test ng/dl	1	AUG 3	1717	**10.3(I) H**	(0.06-0.87)	ng/dL

(I) ----------ADDITIONAL INFORMATION----------
Testing performed by Equilibrium Dialysis.
This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.
See also (*J)
(*J) Mayo Medical Laboratories
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Patient: BLAYK,BONZE ANNE ROSE A00088103734 (Continued)

Test	Day	Date	Time	Result		Reference	Units
Testosterone	1	AUG 3	1717	395(K)	H	(8-60)	ng/dL

(K) ----------ADDITIONAL INFORMATION----------
Testing performed by Liquid Chromatography-Tandem Mass Spectrometry (LC-MS/MS).
This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.
Test Performed by:
Mayo Clinic Laboratories - Rochester Superior Drive
3050 Superior Drive NW, Rochester, MN 55901
See also (*L)
(*L) Mayo Medical Laboratories
.

** END OF REPORT **
* ML = Testing performed at Main Lab