MAYO CLINIC
Mayo Medical Laboratories

1-800-533-1710

BHCG

Beta-Human Chorionic Gonadotropin, Quantitative, Serum

Patient ID	Patient Name		Birth Date	Gender	Age
SA00094070	TESTINGRNV, REPORTS NORM		1976-09-30	F	40
Order Number	**Client Order Number**	**Ordering Physician**	**Report Notes**		
SA00094070	SA00094070	CLIENT,CLIENT			
Account Information		**Collected**			
C7028846 DLMP Rochester		17 Jul 2017 08:00			

Beta-HCG, Quantitative, S [1] SDL

<0.6 IU/L

REFERENCE VALUE

Females:
<1.0 (Premenopausal, non-pregnant)
<7.0 (Postmenopausal)

The testing method is an electrochemiluminescence assay manufactured by Roche Diagnostics Inc. and performed on the Modular or Cobas system.

Values obtained with different assay methods or kits may be different and cannot be used interchangeably.

Test results cannot be interpreted as absolute evidence for the presence or absence of malignant disease.

Received: 18 Jul 2017 12:10 **Reported:** 18 Jul 2017 12:12

Laboratory Notes

[1] This test has been modified from the manufacturer's instructions. Its performance characteristics were 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.

Performing Site Legend

Code	Laboratory	Address
SDL	Mayo Clinic Laboratories - Rochester Superior Drive	3050 Superior Drive NW, Rochester MN 55901