MAYO CLINIC
Mayo Medical Laboratories

1-800-533-1710

BHCG

Beta-Human Chorionic Gonadotropin, Quantitative, Serum

Patient ID	Patient Name		Birth Date	Gender	Age
SA00048389	SAMPLEREPORT, BHCG		1966-06-10	F	46
Order Number	**Client Order Number**	**Ordering Physician**	**Report Notes**		
SA00048389	SA00048389	Client, Client			
Account Information		**Collected**			
C7028846 DLMP Rochester		09 Aug 2012 00:00			

Beta-HCG, Quantitative, S SDL

8.0 IU/L

HCG concentrations above the reference interval can occur if the patient is hypogonadal. Inadequate negative feedback to the pituitary, due to low sex hormone levels, may result in elevated HCG. It is recommended that serum luteinizing hormone (LH) or follicle stimulating hormone (FSH) be determined to assess this possibility.

REFERENCE VALUE

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

ADDITIONAL INFORMATION

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: 10 Aug 2012 12:30 **Reported:** 10 Aug 2012 12:30

Performing Site Legend

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