Quanum
Interactive Insights®

DOB:	Age:	Specimen:	Collected:	Client #:	131 E BROAD ST STE 204
Sex:	Fasting:	Requisition:	Received:		FALLS CHURCH, VA 22046-4520
Phone:		Report Status: FINAL	Reported:	HEALTH ONE - PWN	
Patient ID:				Phone: (703) 534-6766	

FASTING:NO

ESTRADIOL

FINAL

ESTRADIOL (2243-4)	**40 H**	Reference Range: < OR = 39 (pg/mL)

Reference range established on post-pubertal patient population. No pre-pubertal reference range established using this assay. For any patients for whom low Estradiol levels are anticipated (e.g. males, pre-pubertal children and hypogonadal/post-menopausal females), the Quest Diagnostics Nichols Institute Estradiol, Ultrasensitive, LCMSMS assay is recommended (order code 30289).
Please note: patients being treated with the drug fulvestrant (Faslodex(R)) have demonstrated significant interference in immunoassay methods for estradiol measurement. The cross reactivity could lead to falsely elevated estradiol test results leading to an inappropriate clinical assessment of estrogen status. Quest Diagnostics order code 30289-Estradiol, Ultrasensitive LC/MS/MS demonstrates negligible cross reactivity with fulvestrant.

TESTOSTERONE, TOTAL, MALES (ADULT), IA

FINAL

TESTOSTERONE, TOTAL, MALES (ADULT), IA (2986-8)	**657**	Reference Range: 250-827 (ng/dL)

TSH

FINAL

TSH (3016-3)	**0.60**	Reference Range: 0.40-4.50 (mIU/L)

PSA, TOTAL

FINAL

PSA, TOTAL (2857-1)	**0.7**	Reference Range: < OR = 4.0 (ng/mL)

This test was performed using the Siemens chemiluminescent method. Values obtained from different assay methods cannot be used interchangeably. PSA levels, regardless of value, should not be interpreted as absolute evidence of the presence or absence of disease.

Performing Sites