

Patient Information	Specimen Informat	ion	Client Information
DOB: AGE: Gender: Phone: Patient ID:	Specimen: Requisition: Lab Ref #: Collected: Received: Reported:		
COMMENTS: FASTING: YES			
Test Name ESTRADIOL Reference range est population. No pre- established using whom low Estradiol pre-pubertal child females), the Quest Estradiol, Ultrase (order code 30289) Please note: patie fulvestrant (Faslo interference in im measurement. The con- elevated estradiol inappropriate clint Quest Diagnostics Ultrasensitive LC/ reactivity with fur PSA, TOTAL The total PSA value standardized againt result will be app to the equimolar-st Coulter). Comparist interpreted with to This test was perference	nts being treated with dex(R)) have demonstrat munoassay methods for e ross reactivity could 1 test results leading t ical assessment of estr order code 30289-Estrad MS/MS demonstrates neg1 lvestrant. 1.0 e from this assay syste st the WHO standard. Th roximately 20% lower wh tandardized total PSA (on of serial PSA result his fact in mind.	ge ients for (e.g. males, t-menopausal nstitute s recommended the drug ed significant stradiol ead to falsely o an ogen status. iol, igible cross m is e test en compared Beckman s should be	<pre>Reference Range < OR = 39 pg/mL < OR = 4.0 ng/mL</pre>
different assay me interchangeably. F value, should not evidence of the pr TESTOSTERONE, FREE (DIALYSIS) AND TOTAL, F TESTOSTERONE, TOTAL, F Men with clinica testosterone value	699 11y significant hypogon s repeatedly in the ran it from testosterone tr	f ute sease. adal symptoms a ge of the 200-3	00 ng/dL
Metab 1973;36:1132 hypogonadal sympto range of the 200-3 treatment after ad For additional in http://education.c	n Invest 1974:53:819-82 -1142. Men with clinica ms and testosterone val 00 ng/dL or less, may b equate risk and benefit formation, please refer uestdiagnostics.com/faq informational/ educati	lly significant ues repeatedly enefit from tes s counseling. to to 7/FAQ165 (This l	in the tosterone ink is



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Test Name

In Range Out Of Range Reference Range

Lab

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics Nichols Institute Valencia. It has not been cleared or approved by the US Food and Drug Administration. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

PERFORMING SITE: