



Patient Information	Specimen Information	Client Information
DOB: AGE: : Gender: Phone: Patient ID: Health ID:	Specimen: Requisition: Lab Ref #: Collected: Received: Reported:	REQUEST A TEST - PWN HEALTH 7027 MILL RD STE 201 BRECKSVILLE, OH 44141-1852

Test Name	In Range	Out Of Range	Reference Range	Lab
ANTI-MULLERIAN HORMONE ; (AMH), FEMALE	2.5		0.36-10.07 ng/mL	SLI
FSH	17.8		mIU/mL	Z99
	Reference Range			
	Follicular Phase		2.5-10.2	
	Mid-cycle Peak		3.1-17.7	
	Luteal Phase		1.5- 9.1	
	Postmenopausal		23.0-116.3	
LH	11.1		mIU/mL	Z99
			Reference Range	
			Follicular Phase	1.9-12.5
			Mid-Cycle Peak	8.7-76.3
			Luteal Phase	0.5-16.9
			Postmenopausal	10.0-54.7
ESTRADIOL	116		pg/mL	Z99
			Reference Range	
			Follicular Phase:	19-144
			Mid-Cycle:	64-357
			Luteal Phase:	56-214
			Postmenopausal:	< or = 31

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.