Quest Diagnostics

SPECIMEN INFORMATION SPECIMEN: LV340301F REQUISITION: 00954919

Lab ref no:

ESTRADIOL

COLLECTED: 2019/02/14 12:54 RECEIVED: 2019/02/14 12:55 REPORTED: 2019/02/19 14:43

PATIENT INFORMATION

John, Smith

DOB: September 11, 1984

AGE: 20 GENDER: Male FASTING: Unknown

Clinical Info:

16

REPORT STATUS: FINAL

ORDERING PHYSICIAN CLIENT INFORMATION

Jane, Doe

pg/mL

2019-02-19 14:43:00 -0800

01

Lab Testing API 280 Madison Avenue Room 912, 9th Floor New York, NY 10016

Test Name Result Flag Reference Range Lab ESTRADIOL

NORMAL

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.

Performing Laboratory Information:

01: Quest Diagnostics-Wood Dale, 1355 Mittel Blvd, Wood Dale IL, phone: , Medical Director: MD Anthony V Thomas