

Test Definition: FGNDR

Gonadotropin Releasing Hormone (Gn-RH,
Luteinizing Hormone-Releasing Hormone
LH-RH)

Overview

Method Name

Direct Radioimmunoassay

NY State Available

No

Specimen

Specimen Type

Serum

Specimen Required

Patient Preparation: Patient should not be on any steroid, ACTH, gonadotropin, or estrogen medications, if possible, for at 48 hours prior to collection of specimen.

Collection Container/Tube:

Preferred: Red top

Acceptable: Serum gel

Submission Container/Tube: Plastic vial

Specimen Volume: 3 mL Serum

Collection Instructions:

1. Centrifuge as soon as possible and aliquot 3 mL serum into a plastic vial.
2. Freeze specimen immediately after separation.
3. Send frozen.

Specimen Minimum Volume

Serum: 1 mL

Reject Due To

| | |
|-----------------|--------|
| Gross hemolysis | Reject |
| Gross lipemia | Reject |
| Gross icterus | Reject |

Specimen Stability Information

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| Specimen Type | Temperature | Time | Special Container |
|---------------|--------------------|----------|-------------------|
| Serum | Frozen (preferred) | 180 days | |
| | Refrigerated | 7 days | |

Clinical & Interpretive

Clinical Information

Refer to www.interscienceinstitute.com/individual-assays/

Reference Values

Male: 4.0-8.0 pg/mL

Female: 2.0-10.0 pg/mL

Performance

PDF Report

Referral

Day(s) Performed

Monday through Friday

Report Available

9 to 11 days

Performing Laboratory Location

Inter Science Institute

Fees & Codes

Fees

- Authorized users can sign in to [Test Prices](#) for detailed fee information.
- Clients without access to Test Prices can contact [Customer Service](#) 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact [Customer Service](#).

Test Classification

This test has not been cleared or approved by the US Food and Drug Administration.

This test was developed and its performance characteristics determined by Inter Science Institute. Values obtained with different methods, laboratories, or kits cannot be used interchangeably with the results on this report. The results

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cannot be interpreted as absolute evidence of the presence or absence of malignant disease.

CPT Code Information

83727

LOINC® Information

| Test ID | Test Order Name | Order LOINC® Value |
|---------|--------------------------------|--------------------|
| FGNDR | Gonadotropin Releasing Hormone | Not Provided |

| Result ID | Test Result Name | Result LOINC® Value |
|-----------|--------------------------------|---------------------|
| FGNDR | Gonadotropin Releasing Hormone | 13660-6 |