

Infertility – Prior Authorization Request

Send completed form to: Case Review Unit CVS/caremark Fax: 888-249-6155

CVS/caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS/caremark toll-free 888-249-6155.** If you have questions regarding the prior authorization, please contact CVS/caremark at **866-814-5506**.

Patient Name:	Date:
Patient's ID:	Patient's Date of Birth:
Physician's Name:	
Specialty:	NPI#:
Physician Office Telephone:	Physician Office Fax:
Quantity:	Frequency:
Route of Administration:	Expected Length of Therapy:

Follistim AQ is the preferred product for your patient's health plan when prescribing Bravelle and/or Gonal-F

1. Which drug(s) are being prescribed for the member above throughout the course of treatment?
Preferred: Follistim AQ **Non-preferred:** Bravelle Gonal-F
 Pregnyl Repronex Cetrotide hCG Menopur
 Leuprolide Acetate Novarel Ganirelix Ovidrel
 Other _____
2. What is the ICD code? _____
3. Would the prescriber like to request an override of the step therapy requirement? Yes No If No, skip to #6
4. Has the member received the medication through a pharmacy or medical benefit within the past 180 days? Yes No
ACTION REQUIRED: Please provide documentation to substantiate the member had a prescription paid for within the past 180 days (i.e., PBM medication history, pharmacy receipt, EOB etc.)
5. Is the medication effective in treating the member's condition? Yes No Continue to #6 and complete this form in its entirety

Complete the following questions if Bravelle and/or Gonal-F are being prescribed.

6. Is the physician willing to switch to the preferred product, Follistim AQ? Yes No
7. Has the patient received treatment with either Bravelle or Gonal-F product within the last 120 days? Yes No
 If Yes, continue to next section.
8. Does the patient have a contraindication to Follistim AQ or any of its drug components? Yes No
 If Yes, continue to next section.
9. Is the patient intolerant to or had confirmed adverse event with Follistim AQ? Yes No
 If Yes, continue to next section.

Complete the following section based on the patient's gender.

Section A: Female Patient

10. What is the diagnosis? Infertility Other _____

11. Indicate therapy status: New therapy Continuation of therapy

12. Indicate prior treatment cycles:

Previous Bravelle cycles (IUI of IVF): _____

Previous Gonal-F cycles (IUI of IVF): _____

Previous Follistim AQ cycles (IUI of IVF): _____

13. Indicate current treatment plan: IUI IVF FET Stimulation/Times Relations

14. Provide procedure authorization number and dates.

Number MUST be obtained from CareFirst Medical Management Dept. prior to submission of this prior authorization request.

AUTH# _____ Approval Date: ____/____/____ to ____/____/____

Section B: Male Patient

15. What is the diagnosis?

Prepubertal cryptorchidism

Testicular hypogonadism

Hypogonadotropic hypogonadism

Other _____

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS/caremark or the benefit plan sponsor.

X _____
Prescriber or Authorized Signature **Date: (mm/dd/yy)**

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