

Prior Authorization Form



**Note:** Please provide as much information as possible on this form. Missing data may cause processing delays for requested authorization(s). Attach additional sheets to this form if necessary.

Please fax the completed PA form and any additional informational sheets to Nirvanahealth at the following fax number:  
+1(866) 871-8565

Patient Information	Prescriber Information
Patient Name: _____	Prescriber Name: _____
Health Plan Name: _____	Prescriber Address: _____
Patient Insurance Id: _____	_____
Patient Date of Birth: _____	Prescriber Phone: (     ) _____
Patient Phone: _____	Prescriber Fax: (     ) _____
	Prescriber Specialty: _____
	Prescriber DEA: _____
	Prescriber NPI: _____

Medication & Medical Information	
Requested Drug(s) & Strength(s):	<input type="checkbox"/> Lupron Depot 11.25 mg (3 month) intramuscular syringe kit <input type="checkbox"/> Lupron Depot 22.5 mg (3 month) intramuscular suspension <input type="checkbox"/> Lupron Depot 22.5 mg (3 month) intramuscular syringe kit
Requested Quantity Limit Over Time – Amount:	
Requested Quantity Limit Over Time – Days:	
Expected Length of Therapy:	
Directions:	
Diagnosis and Diagnosis Codes (ICD-10 Standard Codes):	
List drugs used previously to treat the same condition:	
Additional clinical information or history. Please include any relevant test results and/or medical record notes:	

### Questionnaire

Q1: I, as the provider or designated representative of the provider, certify and attest that the information provided is complete and accurate and that, upon request, I shall provide any information to RxAdvance that RxAdvance determines is reasonably necessary to verify my responses. (Check only one that apply)

Yes

No

Q2: Is the member currently treated with this medication? (Check only one that apply)

Yes (please list start date of therapy (month/day/year)) \_\_\_\_\_  
(\*Required)

No

Q3: What is the member's diagnosis? (Check only one that apply)

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Prostate Cancer

Endometriosis

Uterine Leiomyomata (UL)

Other (please specify the member's diagnosis and provide clinical rationale for the request)

\_\_\_\_\_ (\*Required)

Q4: Requested medication will be used for one of the following: (Check only one that apply)

Anemia caused by Uterine Leiomyomata

Fibroids prior to surgery of Uterine Leiomyomata

Q5: What is the member's diagnosis? (Check only one that apply)

Prostate Cancer

Endometriosis

Uterine Leiomyomata (UL)

Other (please specify the member's diagnosis and provide clinical rationale for the request)

\_\_\_\_\_ (\*Required)

Q6: Does the member have advanced or metastatic disease? (Check only one that apply)

Yes (please specify the member's diagnosis) \_\_\_\_\_ (\*Required)

No (please provide clinical rationale for the request) \_\_\_\_\_

(\*Required)

Q7: Request is for one of the following strength? (Check only one that apply)

7.5 mg

22.5 mg

30 mg

45 mg

Other (please provide clinical strength) \_\_\_\_\_ (\*Required)

Q8: Has the member had surgical ablation to prevent recurrence? (Check only one that apply)

Yes

No

Q9: Has the member had an inadequate response, intolerance or experienced contraindication(s) to at least one NSAID (e.g., diclofenac, ibuprofen, meloxicam, naproxen)? (Check only one that apply)

Yes (please specify at least one drug name, corresponding contraindication(s) or intolerance experienced and the start and end date(s) of therapy (month/year)) \_\_\_\_\_ (\*Required)

No (please provide clinical rationale for the request) \_\_\_\_\_

(\*Required)

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Q10: Has the member had an inadequate response, intolerance or experienced contraindication(s) to at least one oral contraceptive (e.g., norethindrone-ethinyl estradiol, estradiol and norethindrone)? (Check only one that apply)

Yes (please specify at least one drug name, corresponding contraindication(s) or intolerance experienced and the start and end date(s) of therapy (month/year)) \_\_\_\_\_ (\*Required)

No (please provide clinical rationale for the request) \_\_\_\_\_ (\*Required)

Q11: Request is for one of the following strength? (Check only one that apply)

3.75 mg

11.25 mg

Other (please provide clinical strength) \_\_\_\_\_ (\*Required)

Q12: Request is for one of the following strength? (Check only one that apply)

3.75 mg

11.25 mg

Other (please provide clinical strength) \_\_\_\_\_ (\*Required)

Q13: Is the requested medication being used prior to surgery to reduce the size of fibroids to facilitate a surgical procedure (e.g., myomectomy, hysterectomy)? (Check only one that apply)

Yes

No

Q14: Is medication being used for the treatment of anemia? (Check only one that apply)

Yes

No (please provide clinical rationale for the request) \_\_\_\_\_ (\*Required)

Q15: Is the anemia caused by uterine leiomyomata (fibroids)? (Check only one that apply)

Yes

No (please provide clinical rationale for the request) \_\_\_\_\_ (\*Required)

Q16: Is medication being used prior to surgery? (Check only one that apply)

Yes

No (please provide clinical rationale for the request) \_\_\_\_\_ (\*Required)

**Attestation:** I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, Insurer, Medical Group or its designated representatives may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Signature of Prescriber or Authorized Representative:

Date:

Print Authorized Representative Name: