



OptumRx has partnered with CoverMyMeds to receive prior authorization requests, saving you time and often delivering real-time determinations.

Visit go.covermymeds.com/OptumRx to begin using this free service.

Please note: All information below is required to process this request.

Mon-Fri: 5am to 10pm Pacific / Sat: 6am to 3pm Pacific

Lupron Depot® & Lupron Depot-Ped® Prior Authorization Request Form (Page 1 of 2)

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:

Medication Information <small>(required)</small>			
Medication Name:		Strength:	Dosage Form:
<input type="checkbox"/> Check if requesting brand		Directions for Use:	
<input type="checkbox"/> Check if request is for continuation of therapy			

Clinical Information <small>(required)</small>
<p>Select the diagnosis below:</p> <p><input type="checkbox"/> Central precocious puberty (idiopathic or neurogenic)</p> <p><input type="checkbox"/> Endometriosis</p> <p><input type="checkbox"/> Prostate cancer</p> <p><input type="checkbox"/> Uterine leiomyomata (fibroids)</p> <p><input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____</p>
<p>For central precocious puberty (idiopathic or neurogenic), answer the following:</p> <p>Did the onset of secondary sexual characteristics occur in the patient at less than 8 years of age if female or less than 9 years of age if male? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Has the patient had a pubertal response to a GnRH stimulation test? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Does the patient have a bone age that has advanced at least one year beyond the chronological age? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Does the patient have a peak luteinizing hormone (LH) level above pre-pubertal range? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Does the patient have a random LH level in the pubertal range? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Is the requested medication prescribed by or in consultation with a pediatric endocrinologist? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Reauthorization:</p> <p>Is there documentation of bone age monitoring (e.g., radiographic imaging)? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Does the patient have LH levels that have been suppressed to pre-pubertal levels? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Is the requested medication prescribed by or in consultation with a pediatric endocrinologist? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>For endometriosis, answer the following:</p> <p>Has the patient had surgical ablation to prevent recurrence? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Has the patient had a trial and failure, contraindication, or intolerance to a non-steroidal anti-inflammatory drug (NSAID)? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Has the patient had a trial and failure, contraindication, or intolerance to an oral contraceptive? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Is this request for continuation of prior therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Reauthorization:</p> <p>Have the patient's symptoms recurred after one course of therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Will Lupron Depot be used in combination with norethindrone 5mg daily, other "add-back" sex-hormones, or other bone-sparing agents? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>

This document and others if attached contain information that is privileged, confidential and/or may contain protected health information (PHI). The Provider named above is required to safeguard PHI by applicable law. The information in this document is for the sole use of OptumRx. Proper consent to disclose PHI between these parties has been obtained. If you received this document by mistake, please know that sharing, copying, distributing or using information in this document is against the law. **If you are not the intended recipient, please notify the sender immediately.**

Office use only: LupronDepot-LupronDepotPed_CMS_2019Jan-W



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Lupron Depot® & Lupron Depot-Ped®

Prior Authorization Request Form (Page 2 of 2)

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For prostate cancer, answer the following:

- Does the patient have advanced prostate cancer? Yes No
Does the patient have metastatic prostate cancer? Yes No
Is Lupron Depot being used for palliative care? Yes No
Is this request for continuation of prior therapy? Yes No
Has the patient used Lupron Depot within the past 120 days? Yes No

For uterine leiomyomata (fibroids), answer the following:

- Is Lupron Depot being used prior to surgery to reduce the size of fibroids to facilitate a surgical procedure (e.g., myomectomy, hysterectomy)? Yes No
Is Lupron Depot being used for the treatment of anemia? Yes No
Is the anemia caused by uterine leiomyomata (fibroids)? Yes No
Is Lupron Depot being used prior to surgery? Yes No
Is this request for continuation of prior therapy? Yes No

Quantity Limit:

What is the quantity requested per MONTH? _____

What is the reason for exceeding the plan limitations?

- Titration or loading-dose purposes
Patient is on a dose-alternating schedule (e.g., one tablet in the morning and two tablets at night, one to two tablets at bedtime)
Requested strength/dose is not commercially available
There is a medically necessary justification why the patient cannot use a higher commercially available strength to achieve the same dosage and remain within the same dosing frequency. Please specify:
Other:

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note:

This request may be denied unless all required information is received.
If the patient is not able to meet the above standard prior authorization requirements, please call 1-800-711-4555.
For urgent or expedited requests please call 1-800-711-4555.
This form may be used for non-urgent requests and faxed to 1-800-527-0531.