



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

Orilissa™ Prior Authorization Request Form (Page 1 of 2)

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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>							
Select the diagnosis below:							
<input type="checkbox"/> Moderate to severe pain associated with endometriosis							
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____							
Clinical information:							
Has the patient had surgical ablation to prevent recurrence? <input type="checkbox"/> Yes <input type="checkbox"/> No							
Is the patient premenopausal? <input type="checkbox"/> Yes <input type="checkbox"/> No							
Was Orilissa prescribed by or in consultation with an Obstetrics/Gynecologist (OB/GYN) OR reproductive endocrinologist? <input type="checkbox"/> Yes <input type="checkbox"/> No							
Medication history:							
Select if the patient has had a history of inadequate pain control response to the following (must indicate length of trial):							
<input type="checkbox"/> Combination (estrogen/progesterone) oral contraceptive		Duration of trial: <input type="checkbox"/> ≥ 3 months		<input type="checkbox"/> ≥ 6 months			
<input type="checkbox"/> Danazol		Duration of trial: <input type="checkbox"/> ≥ 3 months		<input type="checkbox"/> ≥ 6 months			
<input type="checkbox"/> Progestins		Duration of trial: <input type="checkbox"/> ≥ 3 months		<input type="checkbox"/> ≥ 6 months			
<input type="checkbox"/> Two analgesics (e.g., ibuprofen, meloxicam, naproxen)		Duration of trial: <input type="checkbox"/> ≥ 3 months		<input type="checkbox"/> ≥ 6 months			
Select if the patient has had a history of intolerance or contraindication to the following:							
<input type="checkbox"/> Combination (estrogen/progesterone) oral contraceptive							
<input type="checkbox"/> Danazol							
<input type="checkbox"/> Progestins							
<input type="checkbox"/> Two analgesics (e.g., ibuprofen, meloxicam, naproxen)							
Reauthorization:							
For reauthorization requests, answer the following:							
Has the patient had improvement in pain associated with endometriosis (e.g., improvement in dysmenorrhea and non-menstrual pelvic pain)? <input type="checkbox"/> Yes <input type="checkbox"/> No							
Has impact to bone mineral density been considered? <input type="checkbox"/> Yes <input type="checkbox"/> No							
Has the treatment duration of Orilissa exceeded a total of 24 months? <input type="checkbox"/> Yes <input type="checkbox"/> No							
Quantity limit requests:							
What is the quantity requested per DAY? _____							
What is the reason for exceeding the plan limitations?							
<input type="checkbox"/> Titration or loading dose purposes							
<input type="checkbox"/> 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)							
<input type="checkbox"/> Requested strength/dose is not commercially available							
<input type="checkbox"/> Other: _____							

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: Orilissa_Comm_2019Feb-W



Orilissa™ Prior Authorization Request Form (Page 2 of 2)
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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.
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.