UnitedHealthcare Community Plan

Orilissa
Prior Authorization Request Form

Please complete this **entire** form and fax it to: **866-940-7328**. If you have questions, please call **800-310-6826.**
This form may contain multiple pages. Please complete all pages to avoid a delay in our decision.
Allow at least 24 hours for review.

Section A – Member Information

First Name:	Last Name:	Member ID:
Address:		
City:	State:	ZIP Code:
Phone:	DOB:	Allergies:
Primary Insurance Information (if any):		

Is the requested medication: □ New or □ Continuation of Therapy? If continuation, list start date: _____

Is this patient currently hospitalized? □ Yes □ No If recently discharged, list discharge date: _____

Section B - Provider Information

First Name:		Last Name:		M.D./D.O.
Address:		City:	State:	ZIP code:
Phone:	Fax:	NPI #:	Specialty:	
Office Contact Name / Fax attention to:				

Section C - Medical Information

Medication:	**Strength:**
Directions for use:	**Quantity:**
Diagnosis (Please be specific & provide as much information as possible)**:**	**ICD-10 CODE:**

Is this member pregnant? □ Yes □ No If yes, what is this member's due date? _____

Section D – Previous Medication Trials

Medication Name	Strength	Directions	Dates of Therapy	Reason for failure / discontinuation

Section E – Additional information and Explanation of why preferred medications would not meet the patient's needs: Please refer to the patient's PDL for a list of preferred alternatives

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Member First name:	Member Last name:	Member DOB:

Clinical and Drug Specific Information	
ALL REQUESTS	
□ **Yes** □ **No**	**Does the patient have a diagnosis of moderate to severe pain associated with endometriosis?**
□ **Yes** □ **No**	**Is the patient premenopausal?**
□ **Yes** □ **No**	**Does the patient have a history of trial and failure (e.g., inadequate pain relief), contraindication, or intolerance after a three month trial of any analgesics (e.g., ibuprofen, meloxicam, naproxen)?** *(If yes, complete Section D above)*
□ **Yes** □ **No**	**Does the patient have a history of trial and failure, contraindication, or intolerance after a three month trial to one of the following?** *(If yes, check which applies and complete Section D above)* □ Hormonal contraceptives □ Progestins [e.g., norethindrone (generic Aygestin)]
□ **Yes** □ **No**	**Is Orilissa prescribed by, or in consultation with, one of the following?** *(If yes, check which applies)* □ Obstetrics/Gynecologist (OB/GYN) □ Reproductive endocrinologist
ORILISSA 200 MG	
□ **Yes** □ **No**	**Has the patient received 6 months of treatment with Orilissa?** *If yes, list rationale:*
CONTINUATION OF THERAPY	
□ **Yes** □ **No**	**Does the patient have a documented positive clinical response to Orilissa therapy?** *If yes, list response:*
□ **Yes** □ **No**	**Has the impact to bone mineral density been considered?**
□ **Yes** □ **No**	**Has the patient received 24 months of treatment with Orilissa?**

Provider Signature: _____ **Date:** _____