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Oriahnn® Prior Authorization Request Form (Page 1 of 2) DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY HAVE BARCODES.

This form may be faxed to 844-403-1029.

Member Information (required)			Provider Information (required)				
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: ZI		IP:	
	ı	ledication Info	rmation (required)				
Medication Information (required) Medication Name: Strength: Dosage Form:						<u> </u>	
Wedledion Name.			Directions for Use:			<u> </u>	
			Directions for osc.				
		Clinical Inform	nation (required)				
1. Has the patient tried and had an inadequate response to therapy with hormonal contraceptives?						☐ Yes ☐ No	
2. Does the patient have a documented intolerance, FDA labeled contraindication, or hypersensitivity to hormonal contraceptives?						☐ Yes ☐ No	
3. Does the patient have osteoporosis?						☐ Yes ☐ No	
4. Has the patient used Oriahnn previously?						☐ Yes ☐ No	
5. If yes to question 4, document how long the patient has already been on therapy with Oriahnn:							
6. Does the patient have a history of low-trauma fracture or other risk for osteoporosis or bone loss?						☐ Yes ☐ No	
7. If yes to question 6, does the patient meet both of the following?						☐ Yes ☐ No	
 Prescriber has assessed the patient's bone mineral density and states that the patient can continue therapy 							
Patient has not had a fragility fracture since starting therapy with Oriahnn?							
8. Does the patient have heavy menstrual bleeding associated with uterine leiomyomas (fibroids)?						☐ Yes ☐ No	
9. Is the patient premenopausal?						☐ Yes ☐ No	
10. Has the patient received 24 or more months of therapy with Oriahnn?						☐ Yes ☐ No	
Quantity Limit:							
1. Is the quantity (dose) requested for documented titration purposes at the initiation of therapy (authorization for a 90 day titration period)?						☐ Yes ☐ No	
2. Can the prescribed dose be achieved using a lesser quantity of a higher strength?						☐ Yes ☐ No	
3. Does the requested quantity (dose) exceed the maximum FDA labeled dose (when specified), or the safest studied dose per the manufacturer's product insert?							
4. If yes to question 3, will the prescriber submit documentation in support of therapy with a higher dose for the intended diagnosis? Submitted documentation may include medical records OR fax form which reflects medical record documentation that shows the length of time the requested dose has been used, and what other medications and doses have been tried and failed.						☐ Yes ☐ No	

This document – and others if attached – contains information that is privileged, confidential and/or may contain protected health information (PHI). The provider named above is required by applicable law to safeguard PHI. 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: Oriahnn_2021Jan

Oriahnn® Prior Authorization Request Form (Page 2 of 2)

Information	on this form is accurate as of this date.	
Prescriber's Signature:		Date:
Are there any this review?	y other comments, diagnoses, symptoms, medications tried or fail	led, and/or any other information the physician feels is important to
Please note:	This request may be denied unless all required information is r	received.
	For more information about the prior outborization process, places	contact up at 955 911 2219

For more information about the prior authorization process, please contact us at 855-811-2218. Monday – Friday: 8 a.m. to 1 a.m. Eastern, and Saturday: 9 a.m. to 6 p.m. Eastern