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

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

Hetlioz® Prior Authorization Request Form

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

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:	Zip:

Medication Information (required)			
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 (required)
<p>Continuation of therapy: Is this a continuation of prior therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No If "yes" to the above question, please submit documentation (e.g., medical records, chart notes, pharmacy claims) or provide the dates, duration, and previous regimen used below: (REQUIRED)</p> <hr/>

<p>Select the diagnosis below: <input type="checkbox"/> Non-24-hour sleep-wake disorder <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____</p>
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<p>Clinical Information: Is there documentation the patient has had a sleep study? <input type="checkbox"/> Yes <input type="checkbox"/> No Is there presence of contraindications or absolute drug interactions with existing therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>

<p>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: _____</p> <p>For continuation of existing therapy, also answer the following: Would sudden discontinuation of the dose trigger withdrawal symptoms? <input type="checkbox"/> Yes <input type="checkbox"/> No Would discontinuation of the dose be unsafe for the patient and their condition may worsen or exacerbate? <input type="checkbox"/> Yes <input type="checkbox"/> No Is the prescribing provider attempting to taper or reduce the dose necessary? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>

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 1-855-297-2870.
 This form may be used for non-urgent requests and faxed to 1-844-403-1029.

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.**
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