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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

Addyi® Prior Authorization Request Form (Page 1 of 2)

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| Patient Information (required) | | | Provider Information (required) | | |
|--|--------|------|---------------------------------|--------------|------|
| Patient 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) | | | | | |
| Select the diagnosis below: | | | | | |
| <input type="checkbox"/> Acquired, generalized hypoactive sexual desire disorder (HSDD) | | | | | |
| <input type="checkbox"/> Female sexual interest/arousal disorder | | | | | |
| <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____ | | | | | |
| Clinical information: | | | | | |
| Have the symptoms of HSDD or female sexual interest/arousal disorder persisted for at least 6 months? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Is low sexual desire due to any of the following: A co-existing medical or psychiatric condition, problems within the relationship, or the effects of a medication or other drug substance? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Is the patient premenopausal? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Is the prescriber certified/enrolled in the Addyi REMS Program? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Does the patient have a known history of alcohol abuse? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| For patients with a known history of alcohol abuse, has the patient abstained from alcohol abuse for the past 6 months? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Will the patient abstain from alcohol use during treatment with Addyi? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Does the patient have hepatic impairment (i.e., a Child-Pugh score of 6 points or greater)? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Is the patient concomitantly on a moderate or strong CYP3A4 inhibitor (e.g., ciprofloxacin, clarithromycin, diltiazem, fluconazole, itraconazole, ketoconazole, ritonavir, verapamil)? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Prescriber attestation: | | | | | |
| Does the prescriber attest to ALL of the following? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| <ul style="list-style-type: none"> The potential benefits of Addyi therapy outweigh the risks Both the prescriber and patient have completed the Addyi REMS Program Patient-Provider Agreement Form The information provided is true and accurate to the best of their knowledge and they understand that the plan may perform a routine audit and request the medical information necessary to verify the accuracy of the information provided | | | | | |
| Reauthorization: | | | | | |
| If this is a reauthorization request, answer the following: | | | | | |
| Is there documentation of positive clinical response to Addyi therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Does the patient continue to be premenopausal? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Has the patient continued to abstain from alcohol use during treatment with Addyi? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Does the patient have hepatic impairment (i.e., a Child-Pugh score of 6 points or greater)? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |
| Is the patient concomitantly on a moderate or strong CYP3A4 inhibitor (e.g., ciprofloxacin, clarithromycin, diltiazem, fluconazole, itraconazole, ketoconazole, ritonavir, verapamil)? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | | |

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Office use only: Addyi_Comm_2018Mar-W



Addyi[®] Prior Authorization Request Form (Page 2 of 2)

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Quantity limit requests:

What is the quantity requested per DAY? _____

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
- 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.
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.