

Prior Authorization Form



Note: Please provide as much information as possible on this form. Missing data may cause processing delays for requested authorization(s). Attach additional sheets to this form if necessary.

Please fax the completed PA form and any additional informational sheets to Nirvanahealth at the following fax number: +1(866) 871-8565

Patient Information	Prescriber Information
Patient Name: _____	Prescriber Name: _____
Health Plan Name: _____	Prescriber Address: _____
Patient Insurance Id: _____	_____
Patient Date of Birth: _____	Prescriber Phone: () _____
Patient Phone: _____	Prescriber Fax: () _____
	Prescriber Specialty: _____
	Prescriber DEA: _____
	Prescriber NPI: _____

Medication & Medical Information	
Requested Drug(s) & Strength(s):	<input type="checkbox"/> modafinil 100 mg tablet <input type="checkbox"/> modafinil 200 mg tablet
Requested Daily Quantity Limit – Amount:	
Requested Daily Quantity Limit – Days:	
Expected Length of Therapy:	
Directions:	
Diagnosis and Diagnosis Codes (ICD-10 Standard Codes):	
List drugs used previously to treat the same condition:	
Additional clinical information or history. Please include any relevant test results and/or medical record notes:	

Questionnaire

Q1: I, as the provider or designated representative of the provider, certify and attest that the information provided is complete and accurate and that, upon request, I shall provide any information to RxAdvance that RxAdvance determines is reasonably necessary to verify my responses. (Check only one that apply)

Yes

No

Q2: Is the member currently treated with this medication? (Check only one that apply)

Yes (please list start date of therapy (month/day/year)) _____
(*Required)

No

Q3: What is the member's diagnosis? (Check only one that apply)

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- Obstructive sleep apnea
- Shift-work disorder
- Narcolepsy
- Multiple sclerosis fatigue
- Treatment-resistant depression
- Idiopathic hypersomnia
- Other (please specify the member's diagnosis and provide clinical rationale for the request)
_____ (*Required)

Q4: Is the member responding positively to the therapy? (Check only one that apply)

- Yes (please provide documentation supporting positive clinical response)
_____ (*Required)
- No (please provide medical justification for continuation of therapy)
_____ (*Required)

Q5: Is the member responding positively to the therapy? (Check only one that apply)

- Yes (please provide documentation supporting positive clinical response)
_____ (*Required)
- No (please provide medical justification for continuation of therapy)
_____ (*Required)

Q6: Is the member experiencing relief of fatigue with modafinil therapy? (Check only one that apply)

- Yes
- No (please provide medical justification for continuation of therapy)
_____ (*Required)

Q7: Is the member responding positively to the therapy? (Check only one that apply)

- Yes (please provide documentation supporting positive clinical response)
_____ (*Required)
- No (please provide medical justification for continuation of therapy)
_____ (*Required)

Q8: Is the requested medication being used as adjunctive therapy? (Check only one that apply)

- Yes
- No (please provide medical justification for continuation of therapy)
_____ (*Required)

Q9: What is the member's diagnosis? (Check only one that apply)

- Obstructive sleep apnea
- Shift-work disorder
- Narcolepsy

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Multiple sclerosis fatigue

Treatment-resistant depression

Idiopathic hypersomnia

Other (please specify the member's diagnosis and provide clinical rationale for the request)

_____ (*Required)

Q10: Has the member's diagnosis of narcolepsy been confirmed by a sleep study? (Check only one that apply)

Yes (please provide the results and the date of study) _____

(*Required)

No

Q11: Is there prescriber justification confirming why a sleep study is not feasible? (Check only one that apply)

Yes (please provide the documentation supporting prescriber justification)

_____ (*Required)

No (please provide clinical rationale for the request) _____

(*Required)

Q12: Is the member experiencing fatigue? (Check only one that apply)

Yes

No (please provide clinical rationale for the request) _____

(*Required)

Q13: Member's treatment-resistant depression is associated with which of the following diagnoses: (Check only one that apply)

Diagnosis of major depressive disorder (MDD)

Diagnosis of bipolar depression

Other (please provide clinical rationale for the request) _____

(*Required)

Q14: Has the member had an intolerance, inadequate response or contraindication to at least 2 antidepressants from different classes (e.g., SSRIs, SNRIs, bupropion)? (Check only one that apply)

Yes (please specify the drug name(s), type of response(s), and start and end date(s) of therapy)

_____ (*Required)

No _____ (*Required)

Q15: Is the requested medication being used as adjunctive therapy? (Check only one that apply)

Yes

No (please provide medical justification for continuation of therapy)

_____ (*Required)

Q16: Has the member's diagnosis of idiopathic hypersomnia been confirmed by a sleep study? (Check only one that apply)

Yes (please provide the results and the date of study) _____

(*Required)

No

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Q17: Is there prescriber justification confirming why a sleep study is not feasible? (Check only one that apply)

Yes (please provide the documentation supporting prescriber justification)

_____ (*Required)

No (please provide clinical rationale for the request) _____

(*Required)

Q18: Is the member experiencing 15 or more obstructive respiratory events per hour of sleep confirmed by a sleep study? (Check only one that apply)

Yes (please provide the results and the date of study) _____

(*Required)

No

Q19: Is there prescriber justification confirming why a sleep study is not feasible? (Check only one that apply)

Yes (please provide the documentation supporting prescriber justification)

_____ (*Required)

No

Q20: Is the member experiencing 5 or more obstructive respiratory events per hour of sleep confirmed by a sleep study? (Check only one that apply)

Yes (please provide the results and the date of study) _____

(*Required)

No

Q21: Is there prescriber justification confirming why a sleep study is not feasible? (Check only one that apply)

Yes (please provide the documentation supporting prescriber justification)

_____ (*Required)

No

Q22: Is the member is experiencing one of the following symptoms: unintentional sleep episodes during wakefulness, daytime sleepiness, unrefreshing sleep, fatigue, insomnia, waking up breath holding/gasping/choking, loud snoring, or breathing interruptions during sleep? (Check only one that apply)

Yes (please specify the type of symptom(s) and date of onset)

_____ (*Required)

No (please provide clinical rationale for the request) _____

(*Required)

Q23: Has the member experienced symptoms of excessive sleepiness or insomnia associated with a work period (usually night work) that occurs during the normal sleep period for at least 3 months? (Check only one that apply)

Yes (please specify the type of symptom(s) and date of onset)

_____ (*Required)

No (please specify the clinical rationale for the request) _____

(*Required)

Q24: Does the member's sleep study demonstrate loss of a normal sleep-wake pattern (i.e., disturbed chronobiologic rhythmicity)? (Check only one that apply)

Yes (please provide the results and the date of study) _____

(*Required)

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No (please specify the clinical rationale for the request) _____
(*Required)

Q25: Has it been confirmed that there are no other medical conditions or medications that are causing the symptoms of excessive sleepiness or insomnia? (Check only one that apply)

Yes

No (please specify the medical condition(s) and/or medication(s))
_____ (*Required)

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, Insurer, Medical Group or its designated representatives may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Signature of Prescriber or Authorized Representative:

Date:

Print Authorized Representative Name: