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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

Nuvigil® (armodafinil) & Provigil® (modafinil) Prior Authorization Request Form (Page 1 of 2)

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Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
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 <small>(required)</small>					
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 <small>(required)</small>					
<p>Select the diagnosis below and complete the corresponding questions for that diagnosis:</p> <input type="checkbox"/> Bipolar depression [for Provigil (modafinil) only] <input type="checkbox"/> Fatigue due to multiple sclerosis [for Provigil (modafinil) only] <input type="checkbox"/> Major depressive disorder [for Provigil (modafinil) only] <input type="checkbox"/> Narcolepsy <input type="checkbox"/> Obstructive sleep apnea/hypopnea syndrome (OSAHS) <input type="checkbox"/> Shift work sleep disorder (SWSD) <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
<p>Fatigue due to multiple sclerosis [Provigil (modafinil) only]: Is the requested medication being used in combination with standard educational therapies (e.g., psychoeducation, behavioral programs, scheduled naps, additional non-pharmacological therapies, etc.)? <input type="checkbox"/> Yes <input type="checkbox"/> No If this is a reauthorization request, answer the following: Is the patient experiencing relief of fatigue with the requested medication? <input type="checkbox"/> Yes <input type="checkbox"/> No Is the requested medication still being used in combination with standard educational therapies (e.g., psychoeducation, behavioral programs, scheduled naps, additional non-pharmacological therapies, etc.)? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>					
<p>Major depressive disorder or bipolar depression [Provigil (modafinil) only]: Does the patient have a history of failure, contraindication, or intolerance to at least two antidepressants from different classes (e.g., SSRI, SNRI, bupropion)? <input type="checkbox"/> Yes <input type="checkbox"/> No Is the requested medication being used as adjunctive therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No If this is a reauthorization request, answer the following: Is there documentation of positive clinical response to the requested medication? <input type="checkbox"/> Yes <input type="checkbox"/> No Is the requested medication being used as adjunctive therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>					
<p>Narcolepsy: Has the diagnosis of narcolepsy been confirmed by a sleep study? <input type="checkbox"/> Yes <input type="checkbox"/> No If a sleep study has not been completed, please justify why a sleep study was not feasible: _____</p> <p>If this is a reauthorization request, answer the following: Is there documentation of positive clinical response to the requested medication? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>					

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.**

Office use only: Nuvigil-Armodafinil-Provigil-Modafinil_Comm_2019Jan1-W

Nuvigil[®] (armodafinil) & Provigil[®] (modafinil) Prior Authorization Request Form (Page 2 of 2)

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Obstructive sleep apnea/hypopnea syndrome (OSAHS):

Has the diagnosis of OSAHS been confirmed by a sleep study? Yes No

If a sleep study has not been completed, please justify why a sleep study would not be feasible:

Was the diagnosis of OSAHS defined by 15 or more obstructive respiratory events per hour of sleep)? Yes No

Was the diagnosis of OSAHS defined by 5 or more obstructive respiratory events per hour of sleep **AND** one of these 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)? Yes No

Have standard treatments for the underlying obstruction (e.g., continuous positive airway pressure [CPAP], bi-level positive airway pressure [BPAP], etc.) been used for 3 months or longer? Yes No

Is the patient fully compliant on standard treatments for the underlying obstruction? Yes No

If this is a reauthorization request, answer the following:

Does the patient continue to be fully compliant on concurrent standard treatments (e.g., CPAP, BPAP, etc.) for the underlying obstruction? Yes No

Is the patient experiencing relief of symptomatic hypersomnolence with use of the requested medication? Yes No

Shift work sleep disorder (SWSD):

Select If SWSD has been confirmed by one of the following:

Symptoms of excessive sleepiness or insomnia, for at least 3 months, which is temporally associated with a work period (usually night work) that occurs during the habitual sleep phase

Sleep study demonstrating loss of a normal sleep-wake pattern (i.e., disturbed chronobiologic rhythmicity)

Does the patient's sleep disturbance cause clinically significant distress or significant impairment in occupational functioning? Yes No

Has it been confirmed that no other medical, mental health, or sleep disorder accounts for the symptoms producing insomnia or excessive sleepiness? Yes No

If this is a reauthorization request, answer the following:

Is there documentation of positive clinical response to the requested medication? Yes No

Does the patient still require treatment for SWSD (i.e., sleep disturbance continues to cause significant distress or impairment in occupational functioning)? Yes No

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: _____

Also answer the following:

Does the patient have a history of inadequate response to Provigil (modafinil) 200mg/day? Yes No

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.