

Antinarcolepsy: Provigil, Nuvigil, Armodafinil, and Modafanil

IVICI	nber Information		
1.	Last Name:	2. First Name:	
3.	Trillium ID #:	2. First Name: 4. Date of Birth:	5. Gender:
	criber Information		
1.	Prescriber Name:	2. NPI #:	
2.	Requestor Name (Nurse/Office S	Staff): City: Ext	
3.	Mailing Address:	City:	State: Zip:
4.	Phone #:	Ext Fax #:	
	g Information		
		2. Strength: 3. Quantity per 30 Days:	
4.	Length of Therapy (in days): 🗆 up to 30 Days 🗆 60 Days 🗆 90 Days 🗆 120 Days 🗆 180 Days 🗆 365 Days 🗅 Other		
Clin	ical Information		
Fo	r Initial Authorization, please ans	wer questions 1-7	
1.	Is this an initial authorization? Select 'Yes' for an initial authorization. Select 'No' for a reauthorization request.		
	\Box Yes \Box No		
2.	Does the member have a diagnosis of Narcolepsy? \Box Yes \Box No		
	Does the member have a diagnosis of excessive sleepiness associated with Shift Work Sleep Disorder?		
	□ Yes □ No		
4.	Does the member have excessive fatigue associated with Multiple Sclerosis or Myotonic Dystonia? Yes No		
5.	Does the member have a diagnosis of Obstructive Sleep Apnea-/ Hypopnea Syndrome? Yes No		
6.			
	If the member is being prescribed non-preferred modafanil, has the member tried and failed Provigil and		
	Nuvigil? Yes No		
	a. If 'NO', state a clinical reason why the member cannot use the preferred Brand medications:		
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Fo	r Continuation therapy, please ar	nswer questions 1-8	
8.	Has the member experienced a reduction in excessive daytime sleepiness from pre-treatment baseline as		
	measured by a validated scale (e.g., Epworth Sleepiness Scale, Stanford Sleepiness Scale, Karolinska Sleepiness		
		e.g., Epworth Sieepiness Scale, Stanford Siee	epiness seale, Ratolinska sleepiness

Signature of Prescriber:

_____ Date: _____

(Prescriber Signature Mandatory)

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.