MHS PHARMACY BENEFIT NARCOLEPSY AGENTS PRIOR AUTHORIZATION REQUEST FORM

MHS

550 N. Meridian St. Suite 101 Indianapolis, IN, 46204-1208 Phone: (877) 647-4848 Fax: (866) 399-0929



Today's Date / / Note: This form must be completed by	the prescribing provid	der.	
Patient's Medicaid #	ons must be complete		the request will be returned** e of Birth / / / / / / / / / / / / / / / / / / /
Patient's Name			scriber's Name
Prescriber's IN License #			cialty
Prescriber's NPI #			scriber's Signature
Return Fax #			arn Phone #
Check box if requesting retro-active PA		Date retro	e(s) of service requested for o-active eligibility (if applicable):
eligibility timelines) with dates of service service 30 calendar days or less and go Requested Medication		ays of	submission separately from current PA requests (dates of Dosing
PA Requirements for Nuvigil (arm	odafinil):		
The member is 18 years of age or o	•	the fo	ollowing diagnoses:
 □ Bipolar depression in conjun • List any other medianstabilizers): 			edical intervention(s) itilized for bipolar depression (e.g., mood
☐ Narcolepsy with excessive da	aytime sleepiness		
with appropriate medical inte	rvention(s)		dual excessive daytime sleepiness in conjunction tilized for obstructive sleep apnea (e.g., PAP,
□ Shift work sleep disorder			

PA Requireme	nts for Provigil (modafinil):		
Select ONE of t			
) The member is 6 years of age or older and has one of the following diagnoses:		
	□ Attention deficit hyperactivity disorder (ADHD)		
2)	□ Narcolepsy with excessive daytime sleepiness The member is 18 years of age or older and has one of the following diagnoses:		
	☐ Depression-related fatigue in conjunction with appropriate medical intervention(s)		
	 List any other medical intervention(s) being utilized for depression (e.g., antidepressants): 		
	□ Idiopathic hypersomnia		
	 □ Obstructive sleep apnea/hypopnea syndrome with residual excessive daytime sleepiness in conjunction with appropriate medical intervention(s) ■ List any other medical intervention(s) being utilized for obstructive sleep apnea (e.g., PAP, OPT, etc.)? 		
	□ Shift work sleep disorder		
	□ Sleep deprivation		
	☐ Steinert myotonic dystrophy syndrome		
	 Unipolar or bipolar depression in conjunction with appropriate medical intervention(s) List any other medical intervention(s) being utilized for unipolar/bipolar depression (e.g., antidepressants/mood stabilizers): 		
PA Requireme	nts for Sunosi (solriamfetrol):		
The member is	18 years of age or older and has one of the following diagnoses:		
□ Narcolep	psy with excessive daytime sleepiness		
	ive sleep apnea/hypopnea syndrome with residual excessive daytime sleepiness in conjunction priate medical intervention(s)		
•	List any other medical intervention(s) being utilized for obstructive sleep apnea (e.g., PAP, OPT, etc.)?		
•	Has the member had a previous trial and failure with any of the following in the past year:		
	□ Modafinil Dates of use:		
	□ Armodafinil Dates of use:		
	If no, please provide any other medical justification for use:		

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PA Requirements for Wakix (pitolisant):							
Selec	t ONE of t	he followir	ıg:				
	☐ The member is 6 years of age or older and has a diagnosis of narcolepsy with cataplexy or excessive daytime sleepiness						
		e with resi			a diagnosis of obstructive sleep apnea/hypopnea ness in conjunction with appropriate medical		
	•	List any o OPT, etc.	y other medical intervention(s) being utilized for obstructive sleep apnea (e.g., PAP, tc.)?				
	•	Has the n	nember had	a previous trial an	d failure with any of the following in the past year:		
		□ Moda	finil	Dates of use:			
		□ Armo	odafinil	Dates of use:			
		If no, plea	ase documer	nt any other medic	cal justification for use:		
		-			-		
PA R	equireme	nts for Xy	rem (sodiur	n oxybate):			
Initial	Authoriz						
			efollowing: her is 7 vear	s of age or older:	and has narcolensy with catanleyy or excessive		
	 The member is 7 years of age or older and has narcolepsy with cataplexy or excessive daytime sleepiness diagnosis □ Yes □ No 						
	Please provide requested dose per day:						
			•	•	nclude date of collection):		
	2)		ember is 18 years of age or older and has fibromyalgia diagnosis □ Yes □ No s the member had a previous trial and failure with ONE of the following?				
			1 Amitriptyli	ne	Dates of use:		
			SSRIs		Medication name and dates of use:		
			SNRIs		Medication name and dates of use:		
			l Anticonvul gabapentin, բ		Medication name and dates of use:		
			NSAIDs a	nd APAP	Dates of use:		
	If member has not trialed all of the above agents, please provide medical justification as to why that agent or an agent in that class was not trialed.						
		Pleas	e provide re	quested dose per	day:		

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Reauthoriza	
	Please provide the following information showing an attempt to decrease dose or trial and
	failure of an alternative therapy within the past year:
	Date of dose reduction attempt: Original dose member was prescribed:
	Original dose member was prescribed:
	Dose member was reduced to:
	Outcomes of dose reduction:
	 Trial and failure of an alternative therapy (name of medication, date of trial, and an
	explanation as to how the member failed):
	Please provide documentation showing continued benefit from the medication (i.e., reduction in frequency of cataplexy, reduction in symptoms of excessive daytime sleepiness, etc.) without significant adverse events (documentation must include most recent chart notes)
PA Require	nents for Xywav (calcium/magnesium/potassium/sodium oxybates solution):
Initial Author	
	ct ONE of the following:
	The member is 7 years of age or older and has narcolepsy with cataplexy or excessive
	daytime sleepiness diagnosis □ Yes □ No
	Please provide requested dose per day: Disase provide requested dose per day:
	Please provide member's weight (include date of collection):
	2) The member is 18 years of age or older has idiopathic hypersomnia □ Yes □ No
	Please provide requested dose per day:
Reauthoriza	
	 Please provide the following information showing an attempt to decrease dose or trial and failure of an alternative therapy within the past year: Date of dose reduction attempt:
	Original dose member was prescribed:
	Dose member was reduced to:
	Outcomes of dose reduction:
	Trial and failure of an alternative therapy (name of medication, date of trial, and an
	explanation as to how the member failed):
	
	Please provide documentation showing continued benefit from the medication (i.e., reduction in requency of cataplexy, reduction in symptoms of excessive daytime sleepiness, etc.) without significant adverse events (documentation must include most recent chart notes)

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