NUPLAZID® (pimavanserin) 34mg capsules

Treatment & Service Request Form

acadia connect

Please complete and fax to 1-844-737-2224, email to number-12 email to number-12 email to number-12 enote that email communications sent to Acadia or its third-party service providers may not be encrypted or secured, and safeguards established under the HIPAA Security Rule would not apply to these communications. See Indication and Important Safety Information, including Boxed WARNING, on page 2. Please read accompanying full Prescribing Information, also available at NUPLAZIDhcp.com.

Patient & Caregiver Support

Phone: 1-844-737-2223

		IFURIVIAI	IUN & INSUR	ANCE Plea	ise fax copies	s of the front and back of prescriptio				*Indicates required field.	
*Patient first name						Section required if patient has	insurance		Patient	does not have insurance	
*Patient last	name					*Prescription drug plan					
*Address			*City			*ID number				Phone number	
*State	*ZIP	*DOB (MM/	DD/YYYY)	Gender		Plan number		Gr	oup nu	mber	
*Patient pho	ne number		*Preferred contact:	☐ Patient ☐	Caregiver	PCN		BI	N numb	per	
*Caregiver name						Medicare Beneficiary ID					
*Caregiver phone number						Preferred language, if not English					
Patient email/Caregiver email						Preferred pharmacy name					
*Patient resi	des: At home A	Assisted living	g ☐ Skilled nursin	g facility/nursin	g home						
LONG TE	RM CARE FACILITIES	If "Assisted I	iving" or "Skilled nur	sing facility/nur	sing home" is	selected, please complete the info	rmation belo	w. Skip S	Section	3 if not needed for resident.	
*Facility na	ame					*Facility phone number					
Address				City			Sta	te		ZIP	
Facility co	ntact name					Job title					
Pharmacy name			Pharmacy pl	hone number		NUPLAZI	NUPLAZID® (pimavanse		nserin) Order on File: Yes No		
☐ Check	this box if your resident is	currently cov	vered under Medicar	e Part A; exped	cted discharge	e date:					
2											
∠ DIAG	NOSIS/PRESCRI	BER INFO	ORMATION								
*Confirmation of diagnosis required ☐ Hallucinations and delusions associated with Parkinson's disease psychosis (PDP). ☐ Other diagnosis:				PP).				*Please confirm dose: 34 mg capsule Other:			
*Prescriber 1	irst and last name				*P	rescriber NPI number		State	license	number (If available)	
Practice/Facility name					*Addres						
Primary contact name					*City	City			te *ZIP		
Prescriber email					*Phone number		*Fa	*Fax			
my patient's assist patient means under prescription f to comply wi	Protected Health Informations in determining their insuration applicable law to the dispeor NUPLAZID, with securing the state-specific presor	in ("PHI") to Adance coverage ensing pharma g any insurand ription require	cadia Pharmaceutical e for NUPLAZID that I cy chosen by or for th ce coverage for NUPL ments such as e-pre	s Inc. or its reprehave elected to be patient, to the AZID to which the scribing, state-s	esentatives or prescribe. I di patient's heal he patient is e pecific prescri	agents (collectively "Acadia") as may rect Acadia to convey, on my behalf, a th insurance company, or to other thin ntitled, or other third parties to assist v	be necessar any prescription d parties as re with patient as	y for the pon information in the policy of t	patient's ation de ecessary or redu	legal representative) for the release of participation in a program designed to divered to Acadia for NUPLAZID by any to assist this patient with filling his/heced-cost medication. I understand I are for additional information relating to	
	iber or authorized agent s	innature (No et	tamo allowed)							·Date	
2		-		if NUPLAZID®	(pimavanse	rin) order is on file for long term	care resider	nt.		Duto	
Known drug	allergies:			☐ None	Concurrent	medications (attach list, if more spa	ace is neede	d):		☐ None	
NUPLAZIO)® (pimavanserin) ON	GOING PR	ESCRIPTION If r	narking checkb	ox for ongoin	g prescription already sent to pharm	nacy or prefe	er to e-pr	escribe	, skip prescription fields.	
	provided prescription to									ect confirms appropriate pharmacy	
Refills (#		sig. Take 34	4 mg capsule orally.	once daily	Dispense: 3	0-day supply Other [†]	<u>'</u>			# of days to be dispensed:	
Dispense a	,		3 1			Substitution >>					
written	*Prescriber signati	ırρ		Dat		permitted Prescriber sign	nnature			Date	
EDEE 4A F			mayancorin\ Mat			y per fill (only for patients diagnosed		ations s	d del		
	iption already sent to Ran					ake 34 mg capsule orally, once dail			iu ueius	IOIIS ASSOCIALEU WILL PDP)	
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	nect® may send ree 14-Day Supply	> >									

if extra time is needed.

*Prescriber signature

I hereby authorize and direct my health care providers (including physicians providers of long-term care, and pharmacies) and health insurance companies and each of their respective representatives. employees, staff, and agents (collectively "Providers") to disclose my Protected Health Information ("PHI") to Acadia Pharmaceuticals Inc. and its representatives and agents (collectively "Acadia") for obtaining Acadia Connect support services. I understand that this PHI may include, but is not limited to, my name, address, phone number, and other contact information; information relating to my medical condition, treatment, care management, and health insurance; as well as information provided on this form and any prescription. I understand that pharmacies may receive remuneration (payment) from Acadia for providing patient support services and disclosing associated PHI to Acadia pursuant to this Form.

I authorize Acadia to use and further disclose the PHI it receives as a result of this Form for:

- Providing reimbursement support associated with the filling of my prescription, including verification of my insurance benefits and assistance in securing coverage to which I am entitled.
- Facilitating the provision of patient assistance, reduced-cost medication, co-pay assistance, and/or other product-related services offered by Acadia, patient advocacy organizations, or other third parties.
- Sending me communications related to the Acadia Connect support services.
- Administrative purposes related to the above services.
- Following de-identification, use for research purposes.

I authorize Acadia to contact me using the contact information I have provided this Form for the above purposes. I also authorize Acadia to report back to my Providers any PHI about me that Acadia may create or receive.

I understand that once my PHI is disclosed to Acadia pursuant to this Form, it may be no longer be protected by the Health Insurance Portability and Accountability Act (HIPAA) and may be subject to re-disclosure.

I understand that I may refuse to sign this Form and my refusal will not affect the treatment I receive from my Providers, nor will it affect my enrollment or eligibility for health insurance benefits to which I am otherwise entitled. I also understand that I may cancel (revoke) this authorization at any time by mailing a letter requesting such cancellation to the address below; however, this cancellation will not apply to any PHI already used or disclosed in reliance on this Form before notice of the cancellation is received by my Providers.

I understand that this authorization is valid for a period of 10 years or for a shorter period dictated by applicable state law. I understand that I will be provided with a signed copy of this authorization by the Provider who collects it from me.

Further information concerning Acadia's privacy practices can be found at https://www.acadia-pharm.com/privacy. If you are a resident of California, a description of the personal information collected by Acadia and your rights under the California Consumer Privacy Act can be found at this address.

Address to Opt Out of Communications or to Cancel This Form: Acadia Connect, PO Box 15713, Pittsburgh, PA 15244

>>	Patient signature	Date
>>	Personal representative (if applicable) signature	Date

AUTHORIZATION TO DISCLOSE INFORMATION TO INDIVIDUALS INVOLVED IN MY CARE (optional)

I further authorize Acadia Pharmaceuticals Inc. to discuss the coordination of my care with the following family member(s) and/or caregiver(s): Authorized representative Name (please print) _ Relationship to patient _ Patient signature/legal guardian signature _ Date _

NUPLAZID is indicated for the treatment of hallucinations and delusions associated with Parkinson's disease psychosis.

Important Safety Information

WARNING: INCREASED MORTALITY IN ELDERLY PATIENTS WITH DEMENTIA-RELATED PSYCHOSIS

- Elderly patients with dementia-related psychosis treated with antipsychotic drugs are at an increased risk of death.
- NUPLAZID is not approved for the treatment of patients with dementia who experience psychosis unless their hallucinations and delusions are related to Parkinson's disease.
- Contraindication: NUPLAZID is contraindicated in patients with a history of a hypersensitivity reaction to pimavanserin or any of its components. Rash, urticaria, and reactions consistent with angioedema (e.g., tongue swelling, circumoral edema, throat tightness, and dyspnea) have been reported.
- Warnings and Precautions: QT Interval Prolongation
- NUPLAZID prolongs the QT interval. The use of NUPLAZID should be avoided in patients with known QT prolongation or in combination with other drugs known to prolong QT interval (e.g., Class 1A antiarrhythmics, Class 3 antiarrhythmics, certain antipsychotics or antibiotics).

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- NUPLAZID should also be avoided in patients with a history of cardiac arrhythmias, as well as other circumstances that may increase the risk of the occurrence of torsade de pointes and/or sudden death, including symptomatic bradycardia, hypokalemia or hypomagnesemia, and presence of congenital prolongation of the QT interval.
- Adverse Reactions: The adverse reactions (≥2% for NUPLAZID and greater than placebo) were peripheral edema (7% vs 2%), nausea (7% vs 4%), confusional state (6% vs 3%), hallucination (5% vs 3%), constipation (4% vs 3%), and gait disturbance (2% vs <1%).

• Drug Interactions:

- Coadministration with strong CYP3A4 inhibitors increases NUPLAZID exposure.
- Reduce NUPLAZID dose to 10 mg taken orally as one tablet once daily. Coadministration with strong or moderate CYP3A4 inducers reduces NUPLAZID exposure. Avoid concomitant use of strong or moderate CYP3A4 inducers with NUPLAZID.

Dosage and Administration

Recommended dose: 34 mg capsule taken orally once daily, without titration, with or without food.

NUPLAZID is available as 34 mg capsules and 10 mg tablets.

Please read the accompanying full <u>Prescribing Information</u>, including **Boxed WARNING**, also available at <u>NUPLAZIDhcp.com</u>.



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