Prior Authorization Request Form for **desmopressin nasal spray** (Noctiva)



JOHNS HOPKINS HEALTHCARE

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FAX Completed Form and Applicable Progress Notes to: (410) 424-4037

USFHP Pharmacy Prior Authorization Form

To be completed by Requesting provider		
Drug Name:	Strength:	
Dosage/Frequency (SIG):	Duration of Therapy:	

Questions? Contact the Pharmacy Dept at: (888) 819-1043, option 4

Clinical Documentation must accompany form in order for a determination to be made.

Step	Please complete patient and physicial	nt):					
1	Patient Name: Physician Name:		ame:				
	Address:	Addr	ess:				
	Sponsor ID #	Phon					
<u> </u>	Date of Birth:	Secure Fa	ax #:				
Step	Please complete the clinical asses	Please complete the clinical assessment:					
2	1. Has the patient received this medication benefit in the last 6 months? Please cho not previously have a TRICARE approved Patient/Provider must answer questions a and medications each time (Questions 17,	oose "No" if the patient did PA for Noctiva. bout medical conditions	☐ Yes Proceed to question 16	☐ No Proceed to question 2			
	2. Is Noctiva being prescribed by an urolo endocrinologist, or a nephrologist?	gist, a geriatrician, an	☐ Yes Proceed to question 3	□ No STOP Coverage not approved			
	3. Has nocturnal polyuria been confirmed collection?	with a 24-hour urine	☐ Yes Proceed to question 4	□ No STOP Coverage not approved			
	4. Has the patient had 2 or more nocturnal least 6 months?	Il voids per night for at	☐ Yes Proceed to question 5	□ No STOP Coverage not approved			
	5. Is the patient GREATER than or EQUAL (Only the low dose is allowed for patients		☐ Yes Proceed to question 6	☐ No STOP Coverage not approved			
	6. Is the patient GREATER than or EQUAL (Only the low dose is allowed for patients		☐ Yes Proceed to question 7	☐ No Proceed to question 8			
	7. Provider acknowledges that patients or greater risk of hyponatremia and has a this significant safety concern.		☐ Yes Proceed to question 8	☐ No STOP Coverage not approved			
	8. Is the provider aware that Noctiva has a risk of hyponatremia?	a black box warning for	☐ Yes Proceed to question 9	□ No STOP Coverage not approved			

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inter restr medi	the patient tried non-pharmacologic techniques or lifestyle ventions to manage the nocturia (e.g., nighttime fluid iction, avoidance of caffeine and alcohol, earlier timing of cations, leg elevation and/or use of compression kings)?	☐ Yes Proceed to question 10	☐ No STOP Coverage not approved
	the patient tried oral desmopressin acetate tablets VP tablets, generics)?	☐ Yes Proceed to question 11	□ No STOP Coverage not approved
	iummEq/mL Date Not approved if sodium level is not provided	☐ Yes Proceed to question 12	☐ No STOP Coverage not approved Not approved if sodium level is not provided
	s the patient have a normal sodium level (135-145 meq/L) to initiation of therapy?	☐ Yes Proceed to question 13	□ No STOP Coverage not approved
thera	the patient's sodium level be rechecked after one week of py, and another sodium level is rechecked after 1 month erapy?	☐ Yes Proceed to question 14	□ No STOP Coverage not approved
14. Doe	s the patient have acute or chronic rhinitis?	☐ Yes STOP Coverage not approved	□ No Proceed to question 15
15. Doe	s the patient have atrophy of nasal mucosa?	☐ Yes STOP Coverage not approved	☐ No Proceed to question 17
16. Has	the patient shown a reduction in nocturia episodes?	☐ Yes Proceed to question 17	□ No STOP Coverage not approved
17. Doe	s the patient have any of the following conditions;	□ Yes	□ No
•	renal impairment (eGFR less than 50 mL/min)	STOP	Proceed to question 18
•	polydipsia nocturnal enuresis	Coverage not approved	
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	The state of the state		
•	Suspicion of bladder outlet obstruction (BOO) or urine flow,		
•	surgical treatment, including transurethral resection, for BOO or benign prostatic hyperplasia within the past 6 months		

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	18. Does the patient have any of the following conditions;	□ Yes	□ No
	urinary retention or a post-void residual volume in	STOP	Proceed to question 19
	excess of 250 mL as confirmed by bladder ultrasound performed after suspicion of urinary retention	d Coverage not approved	
	 current or a history of urologic malignancies (eg; urothelium, prostate, or kidney cancer) 		
	 genitourinary tract pathology (eg; infection or stone the bladder and urethra causing symptoms), 	in	
	 neurogenic detrusor activity (detrusor overactivity) 		
	 suspicion or evidence of cardiac failure 		
	 history of obstructive sleep apnea 		
	 hepatic and/or biliary diseases 		
	 previous desmopressin treatment for nocturia 		
	 treatment with another investigational product within 3 months prior to initiating therapy 	1	
	 known alcohol or substance abuse OR work or lifestyle that may have interfered with regular nighttime sleep 		
	19. Is the patient currently taking any of the following medication	ns:	□ No
	loop diuretics, thiazide diuretics, systemic or inhaled corticosteroids, lithium, alpha1-adrenoceptor antagonists, 5-	STOP	Sign and date below
	alpha reductase inhibitors (5-ARIs), anticholinergics,	Coverage not approved	
	antispasmodics, sedative/hypnotic agents, NSAIDs, selective	•	
	serotonin reuptake inhibitors (SSRIs), serotonin- norepinephrine reuptake inhibitors (SNRIs), antidepressants,		
	anti-epileptics, opioids, or sodium glucose co-transporter 2 inhibitors (SGLT2s)?		
Step	I certify the above is true to the best of my knowledge	e. Please sign and date	<u> </u>
3			
	Prescriber Signature	Date	
			[12 September 2019]
For Inter	nal Use Only		
Appro	ved:	Duration of Approval:	month(s)
Denied	d:	Authorized By:	
Incom	plete/Other:	PA#:	
Date Fax	ed to MD:	Date Decision Rendered:	