OPTIMA HEALTH PLAN

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid)</u> on this request. All other information may be filled in by office staff; <u>fax to 1-800-750-9692</u>. No additional phone calls will be necessary if all information <u>(including phone and fax #s)</u> on this form is correct. <u>If the information provided is not complete, correct, or legible, the authorization process may be delayed.</u>

Drug Requested: Nocdurna® (desmopressin) sublingual tablets

DRUG INFORMATION: Authorization may be delayed if incomplete.				
Drug Name/Form/Strength:				
	Schedule: Length of Therapy:			
Diagn	is: ICD Code, if applicable:	ICD Code, if applicable:		
suppo	ICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be d or request may be denied.			
	Member is 18 years of age or older			
	rescribed by or in consultation with a urologist, geriatrician, or endocrinologist			
	Member is awakening at least two times per night to void while using alternative desmopressin therapuch as desmopressin oral tablets (trial may be waived for members > 65 years of age)	y,		
	Member has a diagnosis of nocturnal polyuria, as confirmed by a 24-hour urine collection, before reatment initiation and the member meets ONE of the following:			
	Nocturnal urine volume exceeds 20% of the total 24-hour urine volume in members < 65 years of age	•		
	Nocturnal urine volume exceeds 33% of the total 24-hour urine volume in members 65 years of ag or older	ge		
	Member has tried non-pharmacologic techniques or lifestyle interventions to manage the nocturia (e.g ighttime fluid restriction, avoidance of caffeine and alcohol, earlier timing of medications, leg elevated nd/or use of compression stockings)			
	Member is <u>NOT</u> using the requested medication along with a loop diuretic (e.g., furosemide) or ystemic/inhaled corticosteroids			
	Member does <u>NOT</u> have any of the following: current or history of hyponatremia, syndrome of nappropriate antidiuretic hormone (SIADH), congestive heart failure (all classes), polydipsia, or ncontrolled hypertension			
	Member does NOT have renal impairment (eGFR below 50 mL/min/1.73 m2)			
	Member has serum sodium concentrations within the normal range of 135-145 mmol/L			
	rovider has ruled out all possible resolvable underlying causes of nocturia and identified the correct nderlying pathophysiologic cause of nocturia (such as bladder dysfunction, excessive nocturnal uring roduction including but not limited to obstructive sleep apnea, neurodegenerative disease, diabetes nellitus and insipidus, electrolyte deficiencies or excess, current medications, chronic kidney disease.			

Reauthorization: 12 months. Check below all that apply. All criteria must be met for approval. To				
support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be				
provided or request may be denied.				
☐ Member continues to meet all initial authorization criteria				

- ☐ Member has experienced a decrease in the number of nocturnal voids from baseline (prior to starting therapy with requested medication)
- ☐ Member has serum sodium concentrations within the normal range of 135-145 mmol/L
- ☐ Member continues to be monitored for hyponatremia, uncontrolled hypertension, renal impairment

Medication being provided by Specialty Pharmacy - PropriumRx

Not all drugs may be covered under every Plan.

If a drug is non-formulary on a Plan, documentation of medical necessity will be required.

**Use of samples to initiate therapy does not meet step edit/preauthorization criteria. **

*Previous therapies will be verified through pharmacy paid claims or submitted chart notes. *

Member Name:	
Member Optima #:	
Prescriber Name:	
Prescriber Signature:	Date:
Office Contact Name:	
Phone Number:	Fax Number:
DEA OR NPI #:	

*Approved by the Pharmacy and Therapeutics Committee: 2/20/2020

REVISED/UPDATED: 6/41/2020; 8/26/2022