SENTARA 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) on this request</u>. All other information may be filled in by office staff; fax to <u>1-800-750-9692</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If information provided is not complete, correct, or legible, authorization may be delayed.</u>

<u>Drug Requested</u> : (Select drug below)	
□ Nayzilam [®] (midazolam nasal spray)	□ Valtoco® (diazepam nasal spray)
MEMBER & PRESCRIBER INFORMA	ATION: Authorization may be delayed if incomplete.
Member Name:	
Member Sentara #:	Date of Birth:
Prescriber Name:	
Prescriber Signature:	Date:
Office Contact Name:	
Phone Number:	
DEA OR NPI #:	
DRUG INFORMATION: Authorization ma	
Drug Form/Strength:	
Dosing Schedule:	Length of Therapy:
Diagnosis:	ICD Code, if applicable:

Recommended Dosing:

- Nayzilam® nasal spray: Administer one spray (5 mg dose) into one nostril. Optional second dose after 10 minutes. Maximum dose: 2 doses per episode, 1 episode every 3 days, 5 episodes per month.
- Valtoco® (diazepam) nasal spray: Initial Dose: 5 mg and 10 mg doses are administered as a single spray intranasally into one nostril. Administration of 15 mg and 20 mg doses requires two nasal spray devices, one spray (7.5 mg or 10 mg) into each nostril. A second dose, when required, maybe administered at least 4 hours after the initial dose. Maximum dose: 2 doses per episode, 1 episode every 5 days, 5 episodes per month.
- Quantity Limit for Nayzilam and Valtoco: 10 spray units per 30 days

CLINICAL CRITERIA: 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.

Authorization Criteria

(Continued on next page)

	Member must meet ONE of the following age requirements:
	☐ If requesting Nayzilam®, member must be 12 years of age or older
	☐ If requesting Valtoco®, member must be 6 years of age or older
	AND
	Prescribing physician is a neurologist or has consulted with a neurologist
	AND
	Member has a diagnosis of epilepsy
	AND
	Member will be using Nayzilam® or Valtoco® for the acute treatment of intermittent episodes of frequent seizure activity (i.e., seizure clusters, acute repetitive seizures) which are distinct from the member's usual seizure pattern with epilepsy (chart notes must be submitted for documentation of seizure activity)
	<u>AND</u>
	Member is currently receiving maintenance antiepileptic medication(s) (e.g., lamotrigine, levetiracetam topiramate, oxcarbazepine)
	<u>AND</u>
	Prescriber agrees to assess the member before prescribing concomitant opioid therapy to limit opioid dosages and durations to the minimum required
	<u>AND</u>
	Dose does not exceed the FDA-approved maximum dose
	<u>AND</u>
	Nayzilam® and Valtoco® will NOT be used concomitantly
'vcl	usion Criteria: Patients with known hypersensitivity to midazolam and acute

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.

narrow-angle glaucoma