SENTARA HEALTH PLANS

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>

Atypical Antipsychotics

Drug Requested: (select one below)

□ asenapine (generic Saphris®)	□ Caplyta [®] (lumateperone)	
□ Fanapt [®] (iloperidone)	□ lurasidone (Latuda®)	
□ paliperidone (generic Invega®)	□ Rexulti [®] (brexpiprazole)	
□ Vraylar® (cariprazine)		
MEMBER & PRESCRIBER INFORMA	ATION: Authorization may be delayed if incomplete.	
Member Name:		
Member Sentara #:	Date of Birth:	
Prescriber Name:		
rescriber Signature: Date:		
Office Contact Name:		
Phone Number: Fax Number:		
DEA OR NPI #:		
DRUG INFORMATION: Authorization m	nay be delayed if incomplete.	
Drug Form/Strength:		
Dosing Schedule:	Length of Therapy:	
Diagnosis:	ICD Code:	
Weight:	Date:	
☐ If diagnosis is any type of depressive disorde	er, please list current antidepressant therapy:	

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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. □ For all antipsychotics used for the treatment of bipolar disorder, major depressive disorder and/or Schizophrenia				
	□ risperidone	□ quetiapine/ER		
	□ ziprasidone	□ olanzapine		
	□ aripiprazole			
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.				
□ For Rexulti requests prescribed for Agitation Associated with Dementia due to Alzheimer's Disease				
Recommended Dosage: Initial: 0.5 mg once daily for 7 days, increase dose on days 8 to 14 to 1 mg once daily, then on day 15 to the target dose of 2 mg once daily. Based on response and tolerability, may increase dose after at least 14 days to the maximum dose of 3 mg once daily.				
Quantity Limit: 1 tablet per day				
	Prescribed by or in consultation with a neurologist of	or psychiatrist		
	Member is 55 years of age or older			
	Member has a confirmed diagnosis consistent with the findings of Alzheimer's disease with magnetic resonance imaging (MRI) or computed tomography of the brain (must submit documentation)			
	Member has/is experiencing signs and symptoms of agitation characterized by increased, often undirected, motor activity, restlessness, aggressiveness, or emotional distress (must submit chart note documentation)			
	Provider continues to monitor member for the occurrence of any medical or neurological conditions (other than Alzheimer's disease) that may be a contributing cause to the member's agitation			
	Provider attests the risk-benefit has been discussed with patients, surrogate decision-makers, families, an or caregivers before starting treatment			
	Member has documented use of non-pharmacological interventions (i.e., music therapy, re-direction of behaviors) prior to starting treatment (must submit chart note documentation)			

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Member has tried and failed at least 30 days of therapy with two (2) of the following:		
□ risperidone	□ quetiapine/ER	
□ ziprasidone	□ olanzapine	
□ aripiprazole		

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. *