

SENTARA HEALTH PLANS

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

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

Drug Requested: Auvelity™ (dextromethorphan HBr and bupropion HCl ER tablets 45 mg/105 mg)

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.

Member Name: _____

Member Sentara #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

NPI #: _____

DRUG INFORMATION: Authorization may be delayed if incomplete.

Drug Name/Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Weight (if applicable): _____ Date weight obtained: _____

Recommended Dosage: One tablet twice a day separated by at least 8 hours.

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.

- Member is 18 years of age or older
- Member has a diagnosis of major depressive disorder (MDD)
- Member must **NOT** have hypersensitivity to bupropion, dextromethorphan, or any component of the requested medication
- Provider attests that member has been screened for personal or family history of bipolar disorder, mania, and hypomania
- Provider attests that member is **NOT** undergoing abrupt discontinuation of alcohol, benzodiazepines, barbiturates, or antiepileptic drugs

(Continued on next page)

- Member will **NOT** take a monoamine oxidase inhibitor (MAOI) within 14 days of Auvelity™
- Member does **NOT** have any of the following:
 - A seizure disorder
 - A diagnosis of bulimia or anorexia nervosa
 - A diagnosis of severe hepatic or severe renal impairment
- Member has had at least a 30-day trial and failure of a serotonin-norepinephrine reuptake inhibitor (SNRI) (e.g., venlafaxine, desvenlafaxine, duloxetine) (**verified by chart notes or pharmacy paid claims**)
- Member has had at least a 30-day trial and failure of a selective serotonin reuptake inhibitor (SSRI) (e.g., citalopram, sertraline, fluoxetine) (**verified by chart notes or pharmacy paid claims**)
- Member has had at least a 30-day trial and failure of one other antidepressant agent (e.g., bupropion, mirtazapine, TCA) (**verified by chart notes or pharmacy paid claims**)

Check each drug that has been tried. If not checked, authorization process will be delayed.		
<input type="checkbox"/> bupropion	<input type="checkbox"/> citalopram	<input type="checkbox"/> desvenlafaxine
<input type="checkbox"/> duloxetine	<input type="checkbox"/> escitalopram	<input type="checkbox"/> fluoxetine
<input type="checkbox"/> mirtazapine	<input type="checkbox"/> paroxetine	<input type="checkbox"/> sertraline
<input type="checkbox"/> venlafaxine ER	<input type="checkbox"/> Other: _____	

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.****