

# SENTARA COMMUNITY PLAN (MEDICAID)

## 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.**

**The Sentara Health Plans Oncology Program is administered by OncoHealth**

- ❖ **For any oncology indications,** the most efficient way to submit a prior authorization request is through the **OncoHealth OneUM Provider Portal** at <https://oneum.oncohealth.us>. Fax to **1-800-264-6128**.  
OncoHealth can also be contacted at Phone: 1-888-916-2616

## ANTIEMETIC/ANTIVERTIGO DRUGS

**Drug Requested:** (Check below the drug that applies)

PREFERRED MEDICATIONS			
(***)PREFERRED Dronabinol and Diclegis Require Prior Authorization)			
<input type="checkbox"/> Diclegis® (doxylamine succinate/vitamin B6) ***	<input type="checkbox"/> ondansetron ODT (4mg, 8mg)/tab/soln	<input type="checkbox"/> meclizine (OTC, Rx)	
<input type="checkbox"/> metoclopramide (tab/soln/vial)	<input type="checkbox"/> Phenadoz® supp (AG) (members over 2 years of age)	<input type="checkbox"/> Prochlorperazine tab	
<input type="checkbox"/> promethazine (AG) (members over 2 years of age)	<input type="checkbox"/> dronabinol capsule***		
All Non-Preferred Medications Require Prior Authorization			
<input type="checkbox"/> Akynzeo®	<input type="checkbox"/> Aloxi®	<input type="checkbox"/> Antivert®	<input type="checkbox"/> Aponvie™
<input type="checkbox"/> Aprepitant capsule/pack	<input type="checkbox"/> Barhemsys®	<input type="checkbox"/> Bonjesta®	<input type="checkbox"/> Cesamet®
<input type="checkbox"/> Cinvanti™	<input type="checkbox"/> Compazine® supp/tab	<input type="checkbox"/> Compro®	<input type="checkbox"/> dimenhydrinate tab, vial
<input type="checkbox"/> Emend® Bi Pak	<input type="checkbox"/> Emend® susp	<input type="checkbox"/> Emend® Tri-fold pack	<input type="checkbox"/> Focinvez™
<input type="checkbox"/> Fosaprepitant vial	<input type="checkbox"/> granisetron <input type="checkbox"/> Granisol soln	<input type="checkbox"/> Kytril®	<input type="checkbox"/> Marinol®
<input type="checkbox"/> metoclopramide ODT	<input type="checkbox"/> Nereus	<input type="checkbox"/> ondansetron 16mg ODT	<input type="checkbox"/> palonosetron (generic Aloxi®)
<input type="checkbox"/> prochlorperazine supp, vial	<input type="checkbox"/> promethazine 50mg supp, vial, ampule	<input type="checkbox"/> Reglan®	<input type="checkbox"/> Sancuso® patch
<input type="checkbox"/> scopolamine (generic Transderm-Scop®)	<input type="checkbox"/> Sustol®	<input type="checkbox"/> Syndros™	<input type="checkbox"/> trimethobenzamide
<input type="checkbox"/> Transderm-Scop®	<input type="checkbox"/> Varubi®	<input type="checkbox"/> Vistaril®	<input type="checkbox"/> Zofran® ODT/soln/tab

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**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: \_\_\_\_\_

**DIAGNOSIS AND 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.

1. Does the member have a diagnosis of severe, chemotherapy-induced nausea and vomiting?  Yes  No
2. If diagnosis is AIDS-related wasting, has member tried and failed megestrol acetate oral suspension **OR** has a contraindication, intolerance, drug-drug interaction?  Yes  No
3. Does the member have nausea or vomiting related to radiation therapy, moderate to highly emetogenic chemotherapy, or post-operative nausea and vomiting?  Yes  No
4. Member has tried and failed therapeutic doses of, or has adverse effects or contraindications to **TWO (2)** different conventional antiemetics (e.g., **promethazine, prochlorperazine, meclizine, metoclopramide, etc.**)?  Yes  No
5. Member has hyperemesis (pregnancy-related nausea/vomiting)?  Yes  No
6. **Bonjesta®/Diclegis®:** length of approval, Estimated Delivery Date (EDD)  
Member must be pregnant and at least 18 years of age  Yes  No  
Estimated Delivery Date: \_\_\_\_\_  
If requesting **Bonjesta®**, member must have tried and failed **Diclegis®**  Yes  No

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7. Does the member have diabetic gastroparesis? If yes, list why oral metoclopramide cannot be used.  Yes  No

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8. For **ondansetron 16 mg ODT**:

Has the member tried and failed or been intolerant to ondansetron 8 mg ODT?  Yes  No

9. Provide clinical evidence that the **Preferred** drug(s) will not provide adequate benefit and list pharmaceutical drugs attempted and outcome.

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10. For **Nereus (tradipitant)**:

Is this medication being used to treat or prevent motion sickness?  Yes  No

Has the member tried and failed meclizine and promethazine?  Yes  No

***\*\*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.\****