

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.

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/tab/soln (maximum quantity per fill=60)	<input type="checkbox"/> meclizine	
<input type="checkbox"/> metoclopramide (tab/soln)	<input type="checkbox"/> Phenadoz[®] supp (AG) (members over 2 years of age)	<input type="checkbox"/> Prochlorperazine (tab/syrup)	
<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"/> Anzemet[®]
<input type="checkbox"/> aprepitant capsule/pack	<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	<input type="checkbox"/> dimenhydrinate
<input type="checkbox"/> Emend[®] Bi Pak	<input type="checkbox"/> Emend[®] susp	<input type="checkbox"/> Emend[®] Tri-fold pack	<input type="checkbox"/> granisetron
<input type="checkbox"/> granisetron	<input type="checkbox"/> Kytril[®]	<input type="checkbox"/> Marinol[®]	<input type="checkbox"/> metoclopramide ODT
<input type="checkbox"/> Metozolv[®] ODT	<input type="checkbox"/> palonosetron (generic Aloxi[®])	<input type="checkbox"/> Phenergan[®]	<input type="checkbox"/> prochlorperazine supp
<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"/> Syndros[™]	<input type="checkbox"/> Tigan[®]	<input type="checkbox"/> Transderm-Scop[®]	<input type="checkbox"/> trimethobenzamide
<input type="checkbox"/> Varubi[®]	<input type="checkbox"/> Vistaril[®]	<input type="checkbox"/> Zofran[®] ODT/soln/tab	<input type="checkbox"/> Zuplenz[®] film

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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: _____

DEA OR NPI #: _____

DRUG INFORMATION: Authorization may be delayed if incomplete.

Drug Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Weight: _____ Date: _____

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. 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. 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, dexamethasone, 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

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

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

*****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.****