

Date of last dose (if applicable):

Dosing frequency:

Medicare Part B vs. Part D determination form

Fax completed form to: 877.974.4411 toll free, or 616.942.8206 **Medicare Part B Medicare Part D** This form applies to: **Urgent** (life threatening) Non-Urgent (standard review) This request is: Urgent means the standard review time may seriously jeopardize the life or health of the patient or the patient's ability to regain maximum function. Granisetron oral Member First Name: _____ Gender: _____ Last Name: Primary Care Physician: Prov. Phone: ______ Prov. Fax: _____ Requesting Provider: Provider Address: Provider NPI: Contact Name: Provider Signature: **Drug** information Start date (or date of next dose):

Part B vs. Part D Coverage Determination Criteria

granisetron oral tablet

This drug requires prior authorization because it may be covered differently under Medicare Part B (medical benefit) or Part D (prescription drug benefit) depending on the patient's circumstances. To determine which benefit the drug is covered under, Priority Health needs to know the use and setting of this drug.

For this drug to be covered under Medicare Part B, the patient must meet the following criteria:

- 1. Must be used for the prevention of chemotherapy-induced nausea and vomiting;
- 2. Must be administered within 2 hours of chemotherapy and continued no more than 48 hours after chemotherapy; and
- 3. Must be used as a full therapeutic replacement for intravenous (IV) anti-emetic drugs that would have otherwise been administered at the time of chemotherapy.

For this drug to be covered under Medicare Part D, the patient must meet the following criteria:

- 1. Must not meet criteria for Medicare Part B coverage (see above)
- 2. Must be used for a medically accepted indication*

Drug product:



Medically accepted indication*

This drug is only covered under Medicare Part D when it is used for a medically accepted indication. A medically accepted indication is a use of the drug that is *either*:

- approved by the Food and Drug Administration. (That is, the Food and Drug Administration has approved the drug for the diagnosis or condition for which it is being prescribed.)
 or —
- supported by certain reference books. (These reference books are the American Hospital Formulary Service Drug Information, DRUGDEX Information System, and Lexi-Drugs)

Priority Health Precertification Documentation			
Α.		t condition is this drug being requested for? Prevention of chemotherapy-induced nausea and vomiting 1. Will granisetron oral be administered within 2 hours of chemotherapy? Yes	
	1.		
	2.	Will granisetron oral be continued for more than 48 hours after chemotherapy? Yes No	
	3.	Will granisetron oral be used as a full therapeutic replacement for IV anti-emetic drugs that would have otherwise been administered at the time of chemotherapy? Yes No	
	 Prevention of postoperative nausea and vomiting Prevention of radiation-induced nausea and vomiting 		
		- the patient's condition is:tionale for Other use:	

Additional information

Note: Criteria are found in the Medicare Benefit Policy Manual, Chapter 15 (Covered Medical and Other Health Services), section 50.5.4 (Oral Anti-Nausea (Anti-Emetic) Drugs)



Priority Health Medicare Part D Exception Request (exceptions to the above criteria)
Do you believe one or more of the prior authorization requirements should be waived? Tes No If yes, you must provide a statement explaining the medical reason why the exception should be approved.
Would granisetron oral likely be the most effective option for this patient? No Yes, because:
If the patient is currently using granisetron oral, would changing the patient's current regimen likely result in adverse effects for the patient? No Yes, because: