

## **Pharmacy Prior Authorization Form** Fax completed form to: 877.974.4411 toll free, or 616.942.8206 This form applies to: Medicaid **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. **Dronabinol** Member Last Name: First Name: DOB: \_\_\_\_\_ Gender: \_\_\_\_ Primary Care Physician: Requesting Provider: \_\_\_\_ Prov. Phone: \_\_\_\_\_ Prov. Fax: \_\_\_\_\_ Provider Address: Contact Name: Provider NPI: Date: \_\_\_ Provider Signature: **Product Information** □ New Request □ Continuation Request Start date (or date of next dose): Drug product: ☐ Dronabinol 2.5 mg capsule ☐ Dronabinol 5 mg capsule Date of last dose (if applicable): ☐ Dronabinol 10 mg capsule Dosing frequency: **Precertification Requirements** Before this drug is covered, the patient must meet all of the following requirements: 1. Diagnosis of chemotherapy induced nausea and vomiting and meet the following: Must be receiving chemotherapy; AND • Trial and failure, intolerance or contraindication to an emetic regimen consistent with NCCN guidelines including ondansetron, granisetron, dexamethasone, promethazine, or prochlorperazine. 2. Appetite stimulation in AIDS patients and meet the following: Patient must have AIDS with anorexia associated with weight loss; AND Must have trial and failure, intolerance, or contraindication to megestrol. Note: Authorization for indications, dosing, or a route of administration not approved by the Food and Drug Administration (FDA) or recognized in CMSaccepted compendia (e.g. DrugDex, AHFS, U.S. Pharmacopeia, and also Clinical Pharmacology for oncology indications only) require supporting evidence for coverage. Please provide two published peer-reviewed literature articles supporting the appropriateness of the drug, the dosing of the drug, or the route of administration to be used for the identified indication. **New request Priority Health Precertification Documentation** A. What condition is this drug being requested for? Chemotherapy induced nausea and vomiting ☐ AIDS with anorexia associated with weight loss ☐ Other – the patient's condition is: \_\_\_\_\_ Rationale for use:



В.	Is the patient receiving chemotherapy?
	☐ Yes
	□ No
C.	Please list what antiemetic regimen the member has trialed, date and outcome:
	Ondansetron, date/outcome:
	Granisetron, date/outcome:
	Dexamethasone, date/outcome:
	Promethazine, date/outcome:
	Prochlorperazine, date/outcome:
	Megestrol, date/outcome:
	None of the above, rationale:

## **Additional information**

**Note:** Duration of approval for chemotherapy induced nausea and vomiting will be limited based on the plan of care developed utilizing the chemotherapeutic agents.