

# OPTIMA HEALTH PLAN

## 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. Incomplete form will delay authorization process.*

**Drug Requested:**     **Syndros® (dronabinol) Oral Solution**

**DRUG INFORMATION:** *Complete information below. If incomplete, authorization process will be delayed*

Drug Form/Strength/Quantity: \_\_\_\_\_

Dosing Schedule: \_\_\_\_\_ Length of Therapy: \_\_\_\_\_

Diagnosis: \_\_\_\_\_ ICD Code, if applicable: \_\_\_\_\_

**CLINICAL CRITERIA:** *The criteria below **MUST** be met or authorization process will be delayed. Check applicable boxes below to qualify. Chart notes **MUST** be submitted to document trial and failure of prerequisite therapies:*

Patient is 18 years of age or older

**DIAGNOSES:** *Check diagnosis that applies. All criteria following diagnosis **must** be met for approval.*

**Anorexia in patients with AIDS**

Prescriber is an Infectious Disease provider specializing in HIV/AIDS treatment

**AND**

Patient has a diagnosis of wasting syndrome due to AIDS

**AND**

Patient has had a 30 day trial and failure of megestrol acetate

**AND**

Patient has had trial and failure of at least three (3) months of dronabinol generic capsules titrated to maximum effective dose

**Chemotherapy-induced nausea and vomiting**

Prescriber is an Oncologist

**AND**

Patient has a diagnosis of cancer with ongoing chemotherapy treatment

**AND**

Patient has had insufficient response from combination treatment for acute/delayed chemotherapy-induced nausea/vomiting with standard treatment (such as ondansetron, dexamethasone or aprepitant). *Please list therapies tried:* \_\_\_\_\_

**AND**

Patient has had trial and failure of olanzapine for refractory nausea/vomiting

**AND**

Patient has had 30-day trial and failure of dronabinol generic capsules titrated to maximum effective dose

**OR**

Patient has difficulty swallowing capsules due to tumor resection or radiation therapy

*(signature on next page)*

**\*\*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.\***

Patient Name: \_\_\_\_\_

Member Optima #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Prescriber Name: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

**DEA OR NPI #:** \_\_\_\_\_

\*Approved by the Pharmacy and Therapeutics Committee: 1/18/2018  
REVISED/UPDATED: 3/6/2018