



Docetaxel

Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720.** If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name: _____ **Date:** _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Physician's Name: _____
Specialty: _____ **NPI#:** _____
Physician Office Telephone: _____ **Physician Office Fax:** _____

Referring Provider Info: Same as Requesting Provider
Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Rendering Provider Info: Same as Referring Provider Same as Requesting Provider
Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Required Demographic Information:

Patient Weight: _____ *kg*

Patient Height: _____ *cm*

Please indicate the place of service for the requested drug:

- Ambulatory Surgical Home Off Campus Outpatient Hospital
 On Campus Outpatient Hospital Office Pharmacy

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

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**CVS Caremark Specialty Pharmacy • 2211 Sanders Road NBT-6 • Northbrook, IL 60062
Phone: 1-888-877-0518 • Fax: 1-855-330-1720 • www.caremark.com**

Criteria Questions:

1. What is the prescribed medication?
 Taxotere (docetaxel) Docefrez (docetaxel) docetaxel (generic) Other _____

2. What is the patient's diagnosis?

<input type="checkbox"/> Breast cancer	<input type="checkbox"/> Bladder cancer
<input type="checkbox"/> Prostate cancer	<input type="checkbox"/> Urothelial carcinoma of the prostate
<input type="checkbox"/> Non-small cell lung cancer (NSCLC)	<input type="checkbox"/> Upper genitourinary tract tumor
<input type="checkbox"/> Gastric cancer	<input type="checkbox"/> Primary carcinoma of the urethra
<input type="checkbox"/> Esophageal and esophagogastric junction cancer	<input type="checkbox"/> Ewing's sarcoma
<input type="checkbox"/> Epithelial ovarian cancer	<input type="checkbox"/> Osteosarcoma
<input type="checkbox"/> Fallopian tube cancer	<input type="checkbox"/> Small cell lung cancer
<input type="checkbox"/> Primary peritoneal cancer	<input type="checkbox"/> Thyroid carcinoma-anaplastic carcinoma
<input type="checkbox"/> Malignant sex-cord stromal tumor	<input type="checkbox"/> Occult primary tumor (cancer of unknown primary)
<input type="checkbox"/> Malignant germ cell tumor residual disease	<input type="checkbox"/> Grade 1 endometrioid carcinoma
<input type="checkbox"/> Carcinosarcoma (malignant mixed Müllerian tumors)	<input type="checkbox"/> Anal cancer
<input type="checkbox"/> Clear cell carcinoma of the ovary	
<input type="checkbox"/> Mucinous carcinoma of the ovary	
<input type="checkbox"/> Small bowel adenocarcinoma	
<input type="checkbox"/> Low-grade serous carcinoma	
<input type="checkbox"/> Uterine neoplasm (including endometrial carcinoma and uterine sarcoma)	
<input type="checkbox"/> Soft tissue sarcoma (including angiosarcoma, extremity/body wall, head/neck, retroperitoneal/intra-abdominal, pleomorphic rhabdomyosarcoma, dermatofibrosarcoma protuberans (DFSP) with fibrosarcomatous transformation and solitary fibrous tumor)	
<input type="checkbox"/> Head and neck cancer (including very advanced head and neck cancer, cancers of the lip (mucosa), oral cavity, salivary gland, oropharynx, hypopharynx, nasopharynx, glottic larynx, and supraglottic larynx)	
<input type="checkbox"/> Other _____	

3. What is the ICD-10 code? _____

4. Is patient currently receiving treatment with the requested medication? Yes No *If No, skip to #6.*

5. Is there evidence of unacceptable toxicity or disease progression while on the current regimen?
 Yes No *No further questions.*

6. What is the clinical setting in which the requested medication will be used? **Indicate ALL that apply.**

<input type="checkbox"/> Advanced disease
<input type="checkbox"/> Metastatic disease
<input type="checkbox"/> Progressive disease
<input type="checkbox"/> Recurrent disease
<input type="checkbox"/> Recurrent unresectable disease
<input type="checkbox"/> Refractory disease
<input type="checkbox"/> Relapsed disease
<input type="checkbox"/> The patient has had no response to preoperative systemic therapy
<input type="checkbox"/> Unresectable locally recurrent disease
<input type="checkbox"/> None of the above

Complete the following section based on the patient's diagnosis, if applicable.

Section A: Breast Cancer

7. Will the requested medication be given in any of the following regimens? **Indicate ALL that apply.**

<input type="checkbox"/> As a single agent
<input type="checkbox"/> As adjuvant therapy
<input type="checkbox"/> As preoperative therapy
<input type="checkbox"/> In combination with capecitabine
<input type="checkbox"/> In combination with pertuzumab and trastuzumab
<input type="checkbox"/> In combination with trastuzumab
<input type="checkbox"/> None of the above

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8. Will the requested medication be used as a substitute for other taxanes (e.g., paclitaxel or albumin-bound paclitaxel) due to medical necessity? *If Yes, no further questions* Yes No.
9. What is the patient's human epidermal growth factor receptor 2 (HER2) status?
 HER2-positive
 HER2-negative
 Unknown

Section B: Anal Cancer

10. What is the patient's disease histology?
 Squamous cell carcinoma
 Non-squamous cell carcinoma

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X _____

Prescriber or Authorized Signature

Date (mm/dd/yy)

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