



Gemcitabine (for Maryland only)

Prior Authorization Request

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

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 **1-866-814-5506**. 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.

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

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

Additional Demographic Information:

Patient Weight: _____kg

Patient Height:______ft _____inches

Criteria Questions:

1. What is the prescribed medication?

Gemzar Gemzitabine (generic) Other

- 2. What is the patient's diagnosis?
 - Pancreatic adenocarcinoma
 - Breast cancer
 - □ Non-small cell lung cancer (NSCLC)
 - Cholangiocarcinoma (intra- or extra-hepatic)
 - Gallbladder cancer
 - Epithelial ovarian cancer/fallopian tube cancer/primary peritoneal cancer

Bladder cancer (includes primary carcinoma of the urethra, upper genitourinary tract tumors, and urothelial carcinoma of the prostate)

- □ Small cell lung cancer (SCLC)
- □ Soft tissue sarcoma (STS)
- Bone cancer
- □ Head and neck cancer
- Hodgkin lymphoma
- □ Non-Hodgkin's lymphoma (NHL)
- Cervical cancer
- Dermatofibrosarcoma protuberans (DFSP)
- □ Kidney cancer
- □ Malignant pleural mesothelioma
- Occult primary
- Testicular cancer
- □ Thymoma/thymic carcinoma
- Uterine sarcoma
- Other

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- 3. What is the ICD-10 code?
- 4. Would the prescriber like to request an override of the step therapy requirement? \Box Yes \Box No If No, skip to #7
- 5. Has the member received the medication through a pharmacy or medical benefit within the past 180 days? □ Yes □ No ACTION REQUIRED: *Please provide documentation to substantiate the member had a prescription paid for within the past 180 days (i.e. PBM medication history, pharmacy receipt, EOB etc.)*
- 6. Is the medication effective in treating the member's condition? □ Yes □ No *Continue to #7 and complete this form in its entirety.*
- 7. Is gemcitable being prescribed as:
 □ First-line or primary therapy □ Second-line or subsequent therapy □ Other _____

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

Section A: Pancreatic Adenocarcinoma

- 10. What is the intent of treatment with gemcitabine?
 - □ Neoadjuvant therapy
 - □ Therapy for unresectable disease
 - □ Adjuvant therapy
 - □ Therapy for locally advanced, recurrent, or metastatic disease
 - Other _____

Section B: Non-Small Cell Lung Cancer (NSCLC)

11. What is the intent of treatment with gemcitabine?

- □ Neoadjuvant or induction chemotherapy, *no further questions*
- Adjuvant chemotherapy, *no further questions*
- □ Therapy for recurrent, locally advanced, or metastatic disease
- □ Other _____
- 12. Is gemcitabine being prescribed for use as a single agent in a patient with metastatic disease, as subsequent therapy following progression on a cytotoxic regimen? *If Yes, no further questions* \Box Yes \Box No
- 13. Is gemcitabine being prescribed for use as subsequent therapy following epidermal growth factor receptor (EGFR) inhibitor therapy (eg, erlotinib [Tarceva], afatinib [Gilotrif], gefitinib [Iressa])? Yes No *If No, skip to #15*
- 14. Does the patient have sensitizing EGFR mutation-positive tumors? Yes No No further questions
- 15. Is gemcitabine being prescribed for use as subsequent therapy following anaplastic lymphoma kinase (ALK) inhibitor therapy (eg, crizotinib [Xalkori])? □ Yes □ No
- 16. Does the patient have ALK-positive tumors? \Box Yes \Box No

Section C: Cholangiocarcinoma (intra- or extra-hepatic), Gallbladder Cancer

- 17. What is the intent of treatment with gemcitabine?
 - □ Primary treatment for unresectable disease
 - □ Primary treatment for metastatic disease
 - □ Adjuvant therapy for resected disease
 - Other ____

 Section D: Epithelial Ovarian Cancer/Fallopian Tube Cancer/Primary Peritoneal Cancer 18. Is the patient's disease platinum-resistant or platinum-sensitive? Platinum-resistant Platinum-sensitive Unknown 	
Section E: Bladder Cancer, Primary Carcinoma of the Urethra, Upper Genitourinary Tract Tumors, Urothelial Carcinoma of the Prostate 19. Is gencitabine being prescribed for use as neoadjuvant or adjuvant therapy? Yes No	
 Section F: Soft Tissue Sarcoma (STS) 20. What is the soft tissue sarcoma subtype? Angiosarcoma Retroperitoneal/Intra-abdominal soft tissue sarcoma Pleomorphic rhabdomyosarcoma Soft tissue sarcoma of the extremity/superficial trunk Other 	
Section G: Bone Cancer 21. What is the bone cancer type? Osteosarcoma Ewing's sarcoma Other	
Section H: Head and Neck Cancer 22. Does the patient have a diagnosis of nasopharyngeal cancer?	
 Section I: Hodgkin Lymphoma 23. Which type of Hodgkin lymphoma is the patient diagnosed with? Classical Hodgkin lymphoma Nodular lymphocyte-predominant Hodgkin lymphoma Other 	
 Section J: Non-Hodgkin's Lymphoma 24. What is the patient's diagnosis (ie, NHL subtype)? Adult T-cell leukemia/lymphoma (ATLL) AIDS-related B-cell lymphoma Burkitt lymphoma Diffuse large B-cell lymphoma (DLBCL) Extranodal NK/T-cell lymphoma, nasal type Follicular lymphoma Gastric MALT lymphoma Mantle cell lymphoma Mycosis fungoides (MF)/Sezary syndrome (SS) Nongastric MALT lymphoma Peripheral T-cell lymphoma Primary cutaneous B-cell lymphoma Splenic marginal zone lymphoma Other 	

Section K: Cervical Cancer

25. Is gencitabine being prescribed for the neoadjuvant treatment of locally advanced disease? Yes No

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.