

Pemetrexed Products (alimta)

Prior Authorization Request

Your patient's benefit plan requires prior authorization for certain medications. In order to make appropriate medical necessity determinations, your patient's diagnosis and other clinical information is required. Please complete the information requested on the form below and fax this form along with supporting clinical documentation to Priority Partners, toll-free at 1-866-212-4756 to initiate the review process. If you have questions regarding the prior authorization please contact Priority Partners at 888-819-1043 Option 4.

Patient's Name:	Date:
Patient's ID:	Patient's Date of Birth:
Physician's Name:	
Specialty:	NPI#:
Physician Office Telephone:	Physician Office Fax:
<u>Referring</u> Provider Info: ☐ Same as Requesting Prov Name:	
Fax:	Phone:
Rendering Provider Info: ☐ Same as Referring Provider Name:	
Fax:	Phone:
	ts in accordance with FDA-approved labeling, evidence-based practice guidelines.
Patient Weight:kg	
Patient Height:cm	
Please indicate the place of service for the requested drug Ambulatory Surgical (POS Code 24) Off Campus Outpatient Hospital (POS Code 19) Office (POS Code 11)	g: ☐ Home (POS Code 12) ☐ On Campus Outpatient Hospital (POS Code 22)
Drug Information:	
Strength/Measure	<u> </u>
, 0,	_Route of administration
Dosing frequency	_
What is the ICD-10 code?	
Clinical Criteria Questions:	
1. What is the diagnosis?	
☐ Bladder cancer (transitional cell urothelium cancer),	Continue to 2
☐ Cervical cancer, <i>Continue to 2</i>	
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Send completed form to: Priority Partners Fax: 1-866-212-4756

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. Pemetrexed Products (alimta) SGM 1900-A - 01/2024.



☐ Fallopian tube cancer, <i>Continue to 2</i>
☐ Non-small cell lung cancer (non-squamous histology), including leptomeningeal metastases, <i>Continue to 2</i> ☐ Pleural or peritoneal mesothelioma, including pericardial mesothelioma and tunica vaginalis testis mesothelioma, <i>Continue to 2</i>
☐ Primary central nervous system (CNS) lymphoma, Continue to 2
☐ Primary peritoneal cancer, <i>Continue to 2</i>
☐ Thymoma or thymic carcinoma, <i>Continue to 2</i>
☐ Other, please specify, Continue to 2
 2. Is this a request for continuation of therapy with the requested medication? ☐ Yes, Continue to 3 ☐ No, Continue to 4
3. Is there evidence of unacceptable toxicity or disease progression while on the current regimen? ☐ Yes, <i>No Further Questions</i> ☐ No, <i>No Further Questions</i>
4. What is the diagnosis?
☐ Bladder cancer (transitional cell urothelium cancer), <i>Continue to 8</i>
☐ Cervical cancer, <i>Continue to 13</i> ☐ Epithelial ovarian cancer (including carcinosarcoma [malignant mixed Mullerian tumor], clear cell carcinoma of the ovary, grade 1 endometrioid carcinoma, low-grade serous carcinoma/ovarian borderline epithelial tumor [low malignant potential], or mucinous carcinoma of the ovary), <i>Continue to 10</i>
☐ Fallopian tube cancer, <i>Continue to 10</i>
☐ Non-small cell lung cancer (non-squamous histology), including leptomeningeal metastases, <i>Continue to 5</i> ☐ Pleural or peritoneal mesothelioma, including pericardial mesothelioma and tunica vaginalis testis mesothelioma, <i>Continue to 6</i>
☐ Primary central nervous system (CNS) lymphoma, Continue to 12
☐ Primary peritoneal cancer, <i>Continue to 10</i>
☐ Thymoma or thymic carcinoma, <i>Continue to 7</i>
5. What is the histology for the disease?
□ Non-squamous histology, No further questions
☐ Squamous histology, No further questions
 6. Will the requested medication be given in any of the following regimens? ☐ As a single agent, <i>No further questions</i> ☐ In combination with cisplatin or carboplatin, <i>No further questions</i>
\square In combination with bevacizumab (Avastin) and either cisplatin or carboplatin, <i>No further questions</i>
☐ In combination with durvalumab (Imfinzi) and either cisplatin or carboplatin, <i>No further questions</i>
☐ Other, please specify, <i>No further questions</i>

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Prescriber or Authorized Signature	Date (mm/dd/yy)
X	
I attest that this information is accurate and true, and that documentation information is available for review if requested by Priority Partners.	supporting this
, 10 January	
Other, please specify, No further questions	ons
☐ Persistent disease, <i>No further questions</i> ☐ Recurrent disease, <i>No further questions</i>	
☐ Metastatic disease, No further questions	
13. What is the clinical setting in which the requested medication will be us	sed?
 12. Will the requested medication be given as single agent? ☐ Yes, No Further Questions ☐ No, No Further Questions 	
 11. Will the requested medication be given as single agent? ☐ Yes, No Further Questions ☐ No, No Further Questions 	
☐ Other, please specify, Continue to 11	
☐ Recurrent disease, Continue to 11	
10. What is the clinical setting in which the requested medication will be us ☐ Persistent disease, <i>Continue to 11</i>	sed?
 9. Will the requested medication be given as second-line treatment? ☐ Yes, No Further Questions ☐ No, No Further Questions 	
☐ Other, please specify, Continue to 9	
☐ Relapsed disease, <i>Continue to 9</i>	
☐ Metastatic disease, <i>Continue to 9</i>	
☐ Locally advanced disease, <i>Continue to 9</i>	
8. What is the clinical setting in which the requested medication will be use	ed?
☐ Yes, No Further Questions ☐ No, No Further Questions	
7. Will the requested medication be given as a single agent?	

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