	Pharmacy Public	Policy Number	MMDP044
		Effective Date	06/01/2022
IOHNS HOPKINS		Review Date	04/20/2022
MEDICINE	<u>Subject</u>	Revision Date	04/20/2022
JOHNS HOPKINS HEALTHCARE	Alimta	Page	1 of 3

This document applies to the following Participating Organizations:

US Family Health Plan

Keywords: Alimta

Table	e of Contents	Page Number
I.	POLICY	1
II.	POLICY CRITERIA	1
III.	AUTHORIZATION PERIOD/LIMITATIONS	2
IV.	EXCLUSIONS	2
V.	RECOMMENDED DOSAGE	2
VI.	CODES	2
VII.	REFERENCES	2
VIII.	APPROVALS	3

I. POLICY

A. Alimta (pemetrexed) will require prior authorization for medical benefit coverage to ensure appropriate use. The process for initiating a prior authorization request can be found in policy PHARM 20.

II. POLICY CRITERIA

- A. Alimta may be approved for patients who meet the following:
 - 1. Bladder Cancer
 - a. Documentation has been submitted showing the patient has locally advanced, metastatic, or relapsed transitional cell urothelium cancer, and Alimta will be used as second-line treatment.
 - 2. Malignant Pleural Mesothelioma (MPM)
 - a. Documentation has been submitted showing one of the following:
 - I. Alimta will be used as a single agent or in combination with cisplatin or carboplatin
 - II. Alimta will be used in combination with bevacizumab and either cisplatin or carboplatin.
 - 3. Non-Small Cell Lung Cancer (Non-Squamous Histology)
 - a. Documentation has been submitted showing the patient has non-squamous non-small cell lung cancer.
 - 4. Ovarian Cancer, Fallopian Tube Cancer, and Primary Peritoneal Cancer
 - a. Documentation has been submitted showing:
 - . The patient has one of the following:
 - persistent or recurrent epithelial ovarian cancer
 - fallopian tube cancer, primary peritoneal cancer
 - carcinosarcoma (malignant mixed Mullerian tumors)
 - clear cell carcinoma
 - grade 1 endometrioid carcinoma
 - low-grade serious carcinoma/ovarian borderline
 - epithelial tumor (low malignant potential) with invasive implants
 - · mucinous carcinoma
 - II. Alimta will be used as a as single agent treatment
 - 5. Primary Central Nervous System (CNS) Lymphoma

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	Johns Hopkins HealthCare LLC Pharmacy Public Medical Management Drug Policies	Policy Number	MMDP044
		Effective Date	06/01/2022
		Review Date	04/20/2022
Subject Alimta		Revision Date	04/20/2022
	Alimta	Page	2 of 3

- Documentation has been submitted showing Alimta will be used as a single agent for treatment of primary CNS lymphoma
- 6. Thymomas and Thymic Carcinomas
 - a. Documentation has been submitted showing Alimta will be used as a single agent for treatment of thymoma or thymic carcinoma
- 7. Malignant Peritoneal Mesothelioma (MPeM)
 - a. Documentation has been submitted supporting a diagnosis of MPeM
- Pericardial Mesothelioma
 - a. Documentation has been submitted supporting pericardial mesothelioma
- 9. Tunica Vaginalis Testis Mesothelioma
 - a. Documentation has been submitted supporting tunica vaginalis testis mesothelioma
- 10. Cervical Cancer
 - a. Documentation has been submitted supporting persistent or recurrent cervical cancer.

III. AUTHORIZATION PERIOD/LIMITATIONS

- A. Initial approval will be limited to 6 months of therapy
- B. Continuation of therapy may be approved in 6-month intervals with documentation showing the patient is continuing to tolerate the regimen and there has not been disease progression while on treatment.

IV. EXCLUSIONS

- A. Alimta will not be covered for the following:
 - 1. Patients with squamous cell NSCLC
 - 2. Any indications or uses that are not FDA-approved, or guideline-supported

V. RECOMMENDED DOSAGE

Please refer to the FDA-approved prescribing information for indication-specific dosing details.

VI. CODES

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Note: The following CPT/HCPCS codes are included below for informational purposes. Inclusion or exclusion of a CPT/HCPCS code(s) below does not signify or imply member coverage or provider reimbursement. The member's specific benefit plan determines coverage.

Medication	HCPCS/CPT Code
Injection, pemetrexed, not otherwise specified, 10 mg	J9305

VII. REFERENCES

1. Alimta [prescribing information]. Indianapolis, IN: Lilly USA, LLC; January 2019.

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	Pharmacy Public Medical Management Drug Policies	Policy Number	MMDP044
		Effective Date	06/01/2022
IOHNS HOPKINS		Review Date	04/20/2022
MEDICINE	<u>Subject</u>	Revision Date	04/20/2022
JOHNS HOPKINS HEALTHCARE	Alimta	Page	3 of 3

2. The NCCN Drugs & Biologics Compendium 2022 National Comprehensive Cancer Network, Inc. Availableat: http://www.nccn.org. Accessed March 1, 2022.

VIII. APPROVALS

Signature on file at JHHC

DATE OF REVISION	SUMMARY OF CHANGE
04/20/2022	Policy Creation

Review Date:04/20/2022

Revision Date: