

Updated: 08/2018 PARP Approved: 10/2018

## Prior Authorization Criteria Oncology Medications

All requests for Oncology Medications require a prior authorization and will be screened for medical necessity and appropriateness using the criteria listed below.

\*\*\*This policy applies to every oncology product that does not have its own specific policy\*\*\*

For all requests for oncology medications all of the following criteria must be met:

- The member has a diagnosis that is indicated in the Food and Drug Administration (FDA) approved package insert, listed in nationally recognized compendia, or peer reviewed medical literature for the determination of medically-accepted indications
- If not indicated as a first line agent, either in the FDA approved package insert, nationally recognized compendia, or peer reviewed medical literature, must provide documentation of previous therapies tried and failed (previous therapies must include those recommended by the FDA approved package insert, nationally recognized compendia, or peer reviewed medical literature)
- Prescribed by, or in consultation with, an oncologist or hematologist
- Unless indicated as monotherapy, must be used in combination with other chemotherapeutic or adjuvant agents according to the FDA approved prescribing information, nationally recognized compendia, or peer reviewed medical literature
- If a test with adequate ability to confirm a disease mutation exists, documentation that the test was performed to confirm the mutation
- The member does not have any contraindications to the requested medication
- For requests to start a new non-formulary agent, the member has had a trial and failure of a formulary agent or a clinically submitted reason for not having a trial of a formulary agent
- The prescribed quantity and dosing regimen is in accordance with the manufacturer's published dosing guidelines, nationally recognized compendia, or peer reviewed medical literature
- **Initial Duration of Approval:** as requested with a maximum of 12 months
- Reauthorization criteria:
  - Documentation that the member had a positive clinical response and is able to tolerate therapy
- **Reauthorization Duration of Approval:** as requested with a maximum of 12 months

Coverage may be provided for any non-FDA labeled indication if it is determined that the use is a medically accepted indication supported by nationally recognized pharmacy compendia or peer-reviewed medical literature for treatment of the diagnosis(es) for which it is prescribed. These requests will be reviewed on a case by case basis to determine medical necessity.

When criteria are not met, the request will be forwarded to a Medical Director for review. The physician reviewer must override criteria when, in their professional judgment, the requested medication is medically necessary.



Updated: 08/2018 PARP Approved: 10/2018

## **ONCOLOGY MEDICATIONS** PRIOR AUTHORIZATION FORM

Please complete and fax all requested information below including any progress notes, laboratory test results, or chart documentation as applicable to Gateway Health<sup>SM</sup> Pharmacy Services. **FAX:** (888) 245-2049

If needed, you may call to speak to a Pharmacy Services Representative.

**PHONE**: (800) 392-1147 Monday through Friday 8:30am to 5:00pm PROVIDER INFORMATION

Requesting Provider:			NPI:				
Provider Specialty:			Office Contact:				
Office Address:			Office Phone:				
			Office Far	x:			
MEMBER INFORMATION							
ember Name: DOB:							
Gateway ID:		Member	weight:	poi	unds or	kg	
REQUESTED DRUG INFORMATION							
Medication:		Strengtl	Strength:				
Frequency:		Duratio	Duration:				
Is the member currently receiving	requested medication? 🗌 Y	Yes No	To Date Medication Initiated:				
Billing Information							
This medication will be billed: at a pharmacy <b>OR</b>							
medically (if medically please provide a JCODE:							
Place of Service: Hospital Provider's office Member's home Other							
Place of Service Information							
Name: Address:			NPI: Phone:				
Address:			Pnone:				
	MEDICAL HISTORY (	Complete fo	n AII no	augsts)			
Diagnosis:		CD-10 Code		quests)			
If a test with adequate ability to confirm a disease mutation exists, was the test performed to confirm the mutation?							
Yes No Not applicable							
Is the requested drug being used in combination with other chemotherapeutic or adjuvant agents?							
☐ Yes, other medications being used: ☐ No							
Does the member have any contraindications to the requested oncology medication?  Yes					s 🔲 No	)	
CURRENT or PREVIOUS THERAPY							
Medication Name	Strength/ Frequency	Dates of Therapy		Status (Discontinued & Why/Current)			
REAUTHORIZATION							
Has the member experienced a positive clinical response and is able to tolerate treatment?  Yes No							
Please describe:							
SUPPORTING INFORMATION or CLINICAL RATIONALE							
Duoganiking Duonid	on Signatura			- Doto-			
Prescribing Provid	er Signature			Date			