



Oncology Agents – Phosphatidylinositol 3-Kinase (PI3K) Inhibitors – Oral – Washington Prior Authorization Request Form

Please complete this **entire** form and fax it to: **866-940-7328**. If you have questions, please call **800-310-6826**.
This form may contain multiple pages. Please complete all pages to avoid a delay in our decision.
Allow at least 24 hours for review.

Date of request:	Reference #:	MAS:	
Patient	Date of birth	ProviderOne ID	
Pharmacy name	Pharmacy NPI	Telephone number	Pharmacy name
Prescriber	Prescriber NPI	Telephone number	Prescriber
Medication and strength		Directions for use	Qty/Days supply
<p>1. Is this request for a continuation of existing therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, is there documentation of a positive clinical response? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>2. What is the patient's diagnosis? <input type="checkbox"/> Breast cancer, advanced or metastatic hormone receptor-positive, HER2-negative, PIK3CA- mutated <input type="checkbox"/> Chronic lymphoid leukemia <input type="checkbox"/> Small lymphocytic lymphoma, relapsed or refractory <input type="checkbox"/> Other Specify: _____</p> <p>3. Provide the following for the patient: Indicate disease stage: Indicate disease type (i.e. New onset, refractory, etc.):</p> <p>4. Indicate if prescribed by or in consultation with: <input type="checkbox"/> Oncologist <input type="checkbox"/> Hematologist <input type="checkbox"/> Other Specify: _____</p> <p>For diagnosis of Breast cancer, advanced or metastatic hormone receptor-positive, HER2-negative, PIK3CA- mutated:</p> <p>5. Will the prescribed medication be used in combination with fulvestrant? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>6. Provide documentation of all of the following:</p> <ol style="list-style-type: none">Hormone receptor-positiveHER2-negativePIK3CA-mutated confirmed <p>7. Has cancer progressed while on or after receiving endocrine therapy (e.g. anastrozole, letrozole, exemestane, tamoxifen)? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>For diagnosis of Chronic lymphoid leukemia:</p> <p>8. For duvelisib: Has patient relapsed? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>			



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Is disease refractory? Yes No

Does patient have a history of failure, contraindication, or intolerance to one of the following?

At least two prior chemotherapy regimen containing:

- Bruton tyrosine kinase inhibitor (BTKi)
- Beta cell lymphoma-2 inhibitor (BCL2i)
- Monoclonal antibody (e.g. obinutuzumab, rituximab)
- Other Specify: _____

9. For idelalisib:

Has patient relapsed? Yes No

Has the patient previously taken another PI3K inhibitor without evidence of progression? Yes No

Will the prescribed medication be used in combination with rituximab? Yes No

Does patient have a history of failure, contraindication, or intolerance to one of the following?

At least one prior chemotherapy regimen containing:

- Bruton tyrosine kinase inhibitor (BTKi)
- Beta cell lymphoma-2 inhibitor (BCL2i)
- Other Specify: _____

For diagnosis of Small lymphocytic lymphoma, relapsed or refractory:

10. Is this being used in combination with other chemotherapeutic, radiotherapeutic, or adjuvant agents?

Yes No

If yes, list all therapies:

11. Does patient have a history of failure, contraindication, or intolerance to one of the following?

At least two prior chemotherapy regimen containing:

- Bruton tyrosine kinase inhibitor (BTKi)
- Beta cell lymphoma-2 inhibitor (BCL2i)
- Monoclonal antibody (e.g. obinutuzumab, rituximab)
- Other Specify: _____

12. Indicate for patient:



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Height (cm):	Date taken:	
Weight (kg):	Date taken:	
Body surface area (m ²):	Date taken:	
Chart notes, labs and all diagnostic tests confirming diagnosis are required with this request		
Prescriber signature	Prescriber specialty	Date