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Please note: All information below is required to process this request.

Mon-Fri: 5am to 10pm Pacific / Sat: 6am to 3pm Pacific

Imbruvica® Prior Authorization Request Form

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:	Specialty:	
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:

Medication Information (required)		
Medication Name:	Strength:	Dosage Form:
<input type="checkbox"/> Check if requesting brand	Directions for Use:	
<input type="checkbox"/> Check if request is for continuation of therapy		

Clinical Information (required)	
Select the diagnosis below:	
<input type="checkbox"/> Chronic lymphocytic leukemia (CLL)	<input type="checkbox"/> Marginal zone lymphoma (MZL)
<input type="checkbox"/> Chronic graft versus host disease (cGVHD)	<input type="checkbox"/> Small lymphocytic lymphoma (SLL)
<input type="checkbox"/> Mantle cell lymphoma (MCL)	<input type="checkbox"/> Waldenström's macroglobulinemia/lymphoplasmacytic lymphoma
<input type="checkbox"/> Other diagnosis: _____	ICD-10 Code(s): _____
Prescriber's Specialty:	
Select if Imbruvica is prescribed by or in consultation with one of the following specialists:	
<input type="checkbox"/> Hematologist	<input type="checkbox"/> Oncologist
<input type="checkbox"/> Physician experienced in the management of transplant patients	
For chronic graft versus host disease, answer the following:	
Has the patient had trial and failure of at least one or more lines of systemic therapy (e.g., corticosteroids, mycophenolate)? <input type="checkbox"/> Yes <input type="checkbox"/> No	
For mantle cell lymphoma (MCL), answer the following:	
Has the patient received at least one prior therapy for MCL (e.g., Rituxan [rituximab])? <input type="checkbox"/> Yes <input type="checkbox"/> No	
For marginal zone lymphoma (MZL), answer the following:	
Has the patient received at least one prior anti-CD20-based therapy for MZL [e.g., Rituxan (rituximab), Zevalin (ibritumomab), Gazyva (obinutuzumab), etc.]? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Reauthorization:	
If this is a reauthorization request, answer the following question:	
Does the patient show evidence of progressive disease while on Imbruvica therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Quantity Limit Requests:	
What is the quantity requested per DAY? _____	
What is the reason for exceeding the plan limitations?	
<input type="checkbox"/> Titration or loading dose purposes	
<input type="checkbox"/> Patient is on a dose-alternating schedule (e.g., one tablet in the morning and two tablets at night, one to two tablets at bedtime)	
<input type="checkbox"/> Requested strength/dose is not commercially available	
<input type="checkbox"/> Other: _____	

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.
For urgent or expedited requests please call 1-800-711-4555.
This form may be used for non-urgent requests and faxed to 1-800-527-0531.

This document and others if attached contain information that is privileged, confidential and/or may contain protected health information (PHI). The Provider named above is required to safeguard PHI by applicable law. The information in this document is for the sole use of OptumRx. Proper consent to disclose PHI between these parties has been obtained. If you received this document by mistake, please know that sharing, copying, distributing or using information in this document is against the law. **If you are not the intended recipient, please notify the sender immediately.**

Office use only: Imbruvica_Comm_2018Jun-W