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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

Gazyva® 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"/> Follicular lymphoma (FL)	
<input type="checkbox"/> Small lymphocytic lymphoma (SLL)	
<input type="checkbox"/> Other diagnosis: _____	ICD-10 Code(s): _____
Clinical Information:	
Is this request for continuation of prior Gazyva therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Has the patient used Gazyva within the last 120 days? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Is Gazyva prescribed by or in consultation with a hematologist/oncologist? <input type="checkbox"/> Yes <input type="checkbox"/> No	
For chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL), answer the following:	
Will Gazyva be used in combination with chlorambucil? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Has the patient previously been treated for CLL? <input type="checkbox"/> Yes <input type="checkbox"/> No	
For follicular lymphoma, answer the following:	
Will Gazyva be used as part of second-line extended dosing? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Will Gazyva be used in combination with bendamustine? <input type="checkbox"/> Yes <input type="checkbox"/> No	
If "yes" to the above question, will treatment be followed by Gazyva monotherapy? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Has the patient relapsed or is refractory to a rituximab-containing regimen? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Does the patient have stage II bulky, III, or IV follicular lymphoma? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Has the patient been treated with prior therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Will Gazyva be used in combination with chemotherapy until the patient has at least achieved partial remission? <input type="checkbox"/> Yes <input type="checkbox"/> No	
If "yes" to the above question, will treatment be followed by Gazyva monotherapy? <input type="checkbox"/> Yes <input type="checkbox"/> No	

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.
 If the patient is not able to meet the above standard prior authorization requirements, please call 1-800-711-4555.
 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: Gazyva_CMS_2019Jan-W