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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

Zydelig® 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"/> Relapsed chronic lymphocytic leukemia (CLL)	
<input type="checkbox"/> Relapsed follicular B-cell non-Hodgkin lymphoma	
<input type="checkbox"/> Relapsed small lymphocytic lymphoma (SLL)	
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____	
Provider's Specialty:	
Is Zydelig prescribed by or in consultation with an oncologist/hematologist? <input type="checkbox"/> Yes <input type="checkbox"/> No	
For relapsed chronic lymphocytic leukemia (CLL), answer the following:	
Has the patient relapsed on at least one prior therapy (e.g., purine analogues [fludarabine, pentostatin, cladribine], alkylating agents [chlorambucil, cyclophosphamide], or monoclonal antibodies [rituximab])? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Will Zydelig be used in combination with Rituxan (rituximab)? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Is the patient a candidate for Rituxan (rituximab) monotherapy due to presence of other comorbidities (e.g., coronary artery disease, peripheral vascular disease, diabetes mellitus, pulmonary disease [COPD], etc.)? <input type="checkbox"/> Yes <input type="checkbox"/> No	
For relapsed follicular B-cell non-Hodgkin lymphoma, answer the following:	
Has the patient relapsed on at least two prior systemic therapies (e.g., rituximab, alkylating agents [cyclophosphamide, chlorambucil], anthracyclines [doxorubicin, daunorubicin], purine analogs [fludarabine])? <input type="checkbox"/> Yes <input type="checkbox"/> No	
For relapsed small lymphocytic lymphoma (SLL), answer the following:	
Has the patient relapsed on at least two prior systemic therapies (e.g., rituximab, alkylating agents [cyclophosphamide, chlorambucil], anthracyclines [doxorubicin, daunorubicin], purine analogs [fludarabine])? <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 Zydelig therapy? <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.
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: Zydelig_Comm_2017Feb-W