## **Besponsa<sup>®</sup>** (inotuzumab ozogamicin) Medication Precertification Request

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(All fields must be completed and legible for precertification review.)

**Aetna Precertification Notification** Phone: 1-866-752-7021 FAX: 1-888-267-3277

For Medicare Advantage Part B: Please Use Medicare Request Form

Fax:

Please indicate:	Start of treatment: Start date / /	_		
	Continuation of therapy: Date of last treatment	/	/	
Precertification Re	equested By:		Phone:	

A. PATIENT INFORMATION									
First Name:			L	Last N	lame:				
Address:			(	City:			State:	ZIP:	
Home Phone:		Work	Phone:			Cell Phone:			
DOB:	Allergies:					Email:			
Current Weight:	lbs ork	gs	Height:	_	inches or	cms			
B. INSURANCE INFORMATIO	N								
Aetna Member ID #:			Does patient have o	other o	coverage?	∕es 🗌 No			
Group #:			If yes, provide ID#:		•				
Insured:			Insured:						
Medicare: 🗌 Yes 🗌 No If	yes, provide ID #:		N	Medic	aid: 🗌 Yes 🔲 N	No If yes, prov	vide ID #:		
C. PRESCRIBER INFORMATI	ON								
First Name:			Last Name:			(Check One	e): 🗌 M.D. 🗌	D.O. 🗌 N.P. 🗌 P.A.	
Address:				C	City:		State:	ZIP:	
Phone:	Fax:		St Lic #:	Ν	NPI #:	DEA #:	U	PIN:	
Provider Email:	•		Office Contact Name	ne:			Phone:		
Specialty (Check one):	Oncologist	Other							
D. DISPENSING PROVIDER/A	DMINISTRATIGON IN	FORM	ATION						
Place of Administration:					Dispensing Provid	der/Pharmacv:	Patient Selec	ted choice	
Self-administered Physician's Office								etail Pharmacy	
Outpatient Infusion Center	-				Specialty Phar		Other:	-	
Center Name:						-			
Home Infusion Center	Phone:				Name:				
Agency Name:					Address:				
Administration code(s) (CF	PT):			_	Phone:				
Address:					TIN:		PIN:		
E. PRODUCT INFORMATION									
Request is for Besponsa (in	otuzumab ozogami	cin):	Dose:		Frequency:				
F. DIAGNOSIS INFORMATION	I – Please indicate pri	mary I	CD Code and specify a	any of	ther where applicable	e.			
Primary ICD Code:	5	Secon	dary ICD Code:			_ Other ICD C	ode:		
G. CLINICAL INFORMATION -	<ul> <li>Required clinical info</li> </ul>	ormatic	on must be completed	in its <u>e</u>	<u>entirety</u> for all precer	tification reques	sts.		
For All Requests (clinical doc	umentation required	<u>):</u>							
Yes No Does the patient	nt have a documented	l diagn	osis of acute lymphob	olastic	leukemia (ALL)?				
For Initiation Requests (Clinic		-							
☐ Yes ☐ No Is there docum	•			e lymp	hoblastic leukemia (	B-ALL)?			
☐ Yes ☐ No Does the patien	in: 🔲 Relapsed 🔲 F								
☐ Yes ☐ No ☐ Unknown Is	•		•	g or an	alysis to identify the	CD22 protein o	n the surface of	the B-cell?	
Please indicate the Philadelphia					, ,	·			
Philadelphia chromosome-p									
Philadelphia chromosome-n	egative (Ph-) disease								
	a dim an i								
Please indicate the requested r	egimen.								
In combination with cyclophe	osphamide. dexameth	asone	. vincristine. methotrex	xate. a	and cytarabine with c	or without blinati	umomab		
In combination with a tyrosir								tive disease	
Other						-	-		
Yes No Will the patient	receive more than 6 t	reatme	ent cycles of the reque	ested o	drug?				



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Patient First Name	Patient Last Name	Patient Phone	Patient DOB
G. CLINICAL INFORMATION (co	ontinued) – Required clinical information must b	e completed in its <u>entirety</u> for all precer	tification requests.
For Continuation Requests (clin	ical documentation required):		
☐ Yes ☐ No Is there evidence	of disease progression or unacceptable toxicity	while on the current regimen?	
Please indicate the number of cyc	les the patient has already received:		
H. ACKNOWLEDGEMENT			
Request Completed By (Signa	ature Required):		Date: / /
	a request for authorization of coverage of a viding materially false information or conceal		

insurance act, which is a crime and subjects such person to criminal and civil penalties. The plan may request additional information or clarification, if needed, to evaluate requests.