

Phesgo® (pertuzumab, trastuzumab, and hyaluronidase-zzxf) Medication Precertification Request

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

Aetna Precertification Notification Phone: <u>1-866-752-7021</u> (TTY: <u>711</u>)

FAX: <u>1-888-267-3277</u>

For Medicare Advantage Part B: Please Use Medicare Request Form

Please indicate:		:: Start date// nerapy: Date of last treatme		_		
Precertification R	equested By:		Pho	one:	Fax:	
A. PATIENT INFOR	RMATION					
First Name:			Last Name:			
Address:			City:		State:	ZIP:
Home Phone:		Work Phone:	1	Cell Pho	ne:	
DOB:	Allergies:			Email:		
Current Weight:	lbs or	kgs He	eight: inche	s or	cms	
B. INSURANCE INI	FORMATION					
Aetna Member ID	#:	Does patient	have other coverage?	☐ Yes ☐ I	No	
Group #:		If yes, provide	e ID#:	Carrier Name	e:	
Insured:		Insured:				
Medicare: Tes	☐ No If yes, provide	ID #:	Medicaid: Te	s 🗌 No If ye	s, provide ID #:	
C. PRESCRIBER IN	NFORMATION					
First Name:		Last Name:		(Chec	k One): M.D.	D.O. N.P. P.A
Address:			City:		State:	ZIP:
Phone:	Fax:	St Lic #:	NPI #:	DEA	#:	UPIN:
Provider Email:		Office Contac	t Name:		Phone	4
Specialty (Check of	one):	☐ Other:				
D. DISPENSING PR	ROVIDER/ADMINISTRA	TION INFORMATION				
Center Na	sion Center Phorame: Center Phor	n's Office ne:	☐ Physici ☐ Specia ☐ Name: Address: _	ian's Office Ity Pharmacy	macy: Patient Sea	macy
☐ Administration of	code(s) (CPT):		TIN:		PIN:	
E. PRODUCT INFO	RMATION					
Request is for: Ph	nesgo (pertuzumab, tras	tuzumab, and hyaluronidase	-zzxf) Dose:	Fre	quency:	
F. DIAGNOSIS INF	ORMATION – Please inc	licate primary ICD Code and s	pecify any other where a	pplicable.		
Primary ICD Code:		Secondary ICD Code	:	Other I	CD Code:	
G. CLINICAL INFO	RMATION – Required cli	nical information must be com	pleted in its <u>entirety</u> for a	Il precertification re	equests.	
Breast cancer What is the hu Please select t ☐ Adjuvant th ☐ Yes ☐ Neoadjuvar ☐ Yes ☐ How ma ☐ Yes ☐ How ma ☐ Yes ☐ Treatment of	man epidermal growth the clinical setting in whereapy No Is the disease any months has the pating No Will the request (pre-operative) treatment (pre-operative) treatment No Is the disease any months has the pating No Will the request of recurrent disease, means and manufactures.	factor receptor 2 (HER2) state ich the requested medication node-positive or at high risk ent received therapy with the sted drug be used in combina	for recurrence? e requested medication? ation with chemotherapy ory, or early stage (either e requested medication? ation with chemotherapy ation with chemotherapy ase had no response to be used? Metastatic	? /? er greater than 2 of ? /? preoperative systems disease	cm in diameter or r temic therapy	
				.2 .144 115 1000011		



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Patient First Name	Patient Last Name	Patient Phone	Patient DOB					
G CLINICAL INFORMATION (continued)	 Required clinical information must be completed. 	eted in its entirety for all precertification	requests					
 G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests. Other, Please explain: 								
For Continuation Requests (clinical documentation required):								
Yes No Is there evidence of unacceptable toxicity or disease progression on the current regimen?								
Please select the clinical setting in which the requested medication is being used:								
☐ Neoadjuvant (pre-operative) treatment of breast cancer								
How many months of the requested medication has the patient received?								
☐ Yes ☐ No Has the patient received the requested drug for 12 months (52 weeks or greater)?								
☐ Adjuvant therapy								
How many months of the requested medication has the patient received?								
☐ Yes ☐ No Has the patient received the requested drug for 12 months (52 weeks or greater)?								
☐ Treatment of recurrent disease, metastatic disease or the disease had no response to preoperative systemic therapy								
Other, Please explain:								
H. ACKNOWLEDGEMENT								
Request Completed By (Signature Requ	uired):		Date:/					
any insurance company by providing mate	t for authorization of coverage of a medical erially false information or conceals materia cts such person to criminal and civil penaltie	i information for the purpose of misl						

The plan may request additional information or clarification, if needed, to evaluate requests.