

Kadcyla® (ado-trastuzumab) **Precertification Request**

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

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

1-888-267-3277

For Medicare Advantage Part B:

Plassa indicata:	☐ Start of treatment: S		/ /	Tor precertification i	cvicw.	PI	ease Use M	ledicare Request For	n
riease ilidicate.	☐ Continuation of thera			/ /					
Precertification Re	equested By:	•			ne:		Fax:		
A. PATIENT INFOR	MATION								
First Name:			L	₋ast Name:					
Address:			(City:			State:	ZIP:	
Home Phone:		Wor	k Phone:			Cell Phone:			
DOB:	Allergies:					E-mail:			
Current Weight:	Ibs or	kgs	Height: _	inches	or _	cms			
B. INSURANCE INF	ORMATION								
Aetna Member ID #	# :		_ Does patient have o	_	_	∕es □ No			
-			If yes, provide ID#:		Car	rier Name:			_
Insured:			Insured:						
Medicare: Yes	☐ No If yes, provide ID	#:	I	Medicaid: 🗌 Yes	☐ No	If yes, provi	de ID #:		_
C. PRESCRIBER IN	FORMATION								
First Name:			Last Name:			(Check One	i	□ D.O. □ N.P. □ P	Α.
Address:			ı	City:			State:	ZIP:	
Phone:	Fax:		St Lic #:	NPI #:		DEA #:		UPIN:	
Provider E-mail:			Office Contact Nam	e:			Phone	э :	
Specialty (Check of	ne): Oncologist	Other: _							
D. DISPENSING PR	OVIDER/ADMINISTRATIO	N INFORM	MATION	<u> </u>					
Place of Administr				-		der/Pharmacy:	Patient Se	elected choice	
☐ Self-administered ☐ Physician's Office				☐ Physician's Office ☐ Retail Pharmacy					
Outpatient Infus						-	Other		_
Center Name: Phone:					Name:				
	ime:								_
Address:									_
	ode(s) (CPT):			TIN:			PIN: _		_
E. PRODUCT INFOR									
Request is for: Dose:	Kadcyla <u>(</u> ado-trastuzum	iab emtai	•	uency:					
	DRMATION – Please indica	te primary		<u> </u>	nlicable	a			
Primary ICD Code:			ndary ICD Code:	arry other where ap	piloabit	Other ICD C	ode.		_
	RMATION – Required clinica			in its entirety for all	precer				
	inical documentation requ		en mast se sempleted	in no <u>ominoty</u> for all	procor	anoadon roquos			
	pidermal growth factor rece		R2) status? ☐ HER2 p	ositive	negativ	e 🗌 Unknown			
☐ Breast cancer									
☐ For early brea	ast cancer ☐ Yes ☐ No Will the req	uested dru	ug be used as adiuvant	treatment?					
-	How many months has the				on?				
For non-early						–			
$\qquad \qquad \longrightarrow \mid$	Please indicate the clinical	setting in v		-				disease mic therapy Other	
,	What is the place in therapy	in which	_				•		
	Will the requested drug be i			_	_				
☐ Non-small cell lu	=			7.4.	_				
	he clinical setting in which t						se ∐ Meta	static disease	
What is the place in therapy in which the requested drug be used? ☐ First-line treatment ☐ Subsequent treatment ☐ Subsequent treatment ☐ Yes ☐ No Will the requested drug be used as a single agent?									



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For Medicare Advantage Part B: Please Use Medicare Request Form

Patient First Name	Patient Last Name	Patient Phone	Patient DOB						
G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.									
Salivary gland tumor Please indicate the clinical setting in which the requested drug will be used: ☐ Recurrent disease ☐ Other ☐ Yes ☐ No Will the requested drug be used as a single agent? For Continuation Requests (clinical documentation required): ☐ Yes ☐ No Has the patient experienced disease progression or unacceptable toxicity while on the current regimen? ☐ Yes ☐ No Is the requested drug being used as adjuvant treatment of early breast cancer? ☐ How many months of the requested medication has the patient received?									
H. ACKNOWLEDGEMENT									
Request Completed By (Signature Requ	uired):		Date:/						
Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent 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.