Enquiries to Ministry of Health 0800 855 066

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 1 Form SA2424 November 2025

APPLICANT (stamp or sticker acceptable)		PATIENT NHI:	REFERRER Reg No:		
Reg No:		First Names:	First Names:		
Name:		Surname:	Surname:		
Address:		DOB:	Address:		
		Address:			
Fax Number: .			Fax Number:		
Trastuzuma	b emtansine				
Applications o	tion — early breast cancer nly from a relevant specialist or med (tick boxes where appropriate)	dical practitioner on the recommendation of a relevan	nt specialist. Approvals valid for 12 months.		
and	Patient has early breast cancer expressing HER2 IHC3+ or ISH+				
and	Documentation of pathological invasive residual disease in the breast and/or axiliary lymph nodes following completion of surgery and Patient has completed systemic neoadjuvant therapy with trastuzumab and chemotherapy prior to surgery				
and					
and	Disease has not progressed during neoadjuvant therapy				
and	Patient has left ventricular ejection fraction of 45% or greater				
and	Adjuvant treatment with trastuzumab emtansine to be commenced within 12 weeks of surgery				
and	Trastuzumab emtansine to be discontinued at disease progression and				
	Total adjuvant treatment duration	must not exceed 42 weeks (14 cycles)			
Initial application — metastatic breast cancer Applications only from a relevant specialist or any relevant practitioner on the recommendation of a relevant specialist. Approvals valid for 6 months. Prerequisites(tick boxes where appropriate)					
and	Patient has metastatic breast can	cer expressing HER-2 IHC 3+ or ISH+ (including FIS	H or other current technology)		
and	Patient has previously received tra	astuzumab and chemotherapy, separately or in comb	oination		
O		ior therapy for metastatic disease*			
		ase recurrence during, or within six months of comple	eting adjuvant therapy*		
and and	Patient has a good performance s	status (ECOG 0-1)			
O	Patient does not have symp	otomatic brain metastases			
		es and has received prior local CNS therapy			
and					
OI		or funded trastuzumab emtansine or trastuzumab de	rruxtecan treatment		
	Patient has disconting	ued trastuzumab deruxtecan due to intolerance			
		ogress while on trastuzumab deruxtecan			
and	Treatment to be discontinued at d	isease progression			

I confirm the above details are correct and that in signing this form I understand I may be audited.

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APPLICANT (stamp or sticker acceptable)	PATIENT NHI:	REFERRER Reg No:			
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Name:	Surname:	Surname:			
Address:	DOB:	Address:			
	Address:				
Fax Number:		Fax Number:			
Trastuzumab emtansine - continued					
Renewal — metastatic breast cancer					
Current approval Number (if known):					
Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 6 months. Prerequisites(tick boxes where appropriate)					
The cancer has not progressed at any time point during the previous approval period whilst on trastuzumab emtansine					
Treatment to be discontinued at disease progression					
Note: Prior or adjuvant therapy includes anthracyc	line, other chemotherapy, biological drugs, or endocr	ine therapy.			

I confirm the above details are correct and that in signing this form I understand I may be audited.