HOSPITAL MEDICINES LIST RESTRICTIONS CHECKLIST

July 2025

Use this checklist to determine if a patient meets the restrictions for funding in the **hospital setting**. For more details, refer to Section H of the Pharmaceutical Schedule. For community funding, see the Special Authority Criteria.

	PATIENT:
e:	Name:
d:	NHI:
stuzumab	(Herzuma)
assessment	arly breast cancer required after 12 months ick boxes where appropriate)
and	The patient has early breast cancer expressing HER-2 IHC 3+ or ISH + (including FISH or other current technology Maximum cumulative dose of 106 mg/kg (12 months' treatment)
assessment	N – early breast cancer* required after 12 months ick boxes where appropriate)
and	The patient has metastatic breast cancer expressing HER-2 IHC 3+ or ISH+ (including FISH or other current technology The patient received prior adjuvant trastuzumab treatment for early breast cancer The patient has not previously received lapatinib treatment for HER-2 positive metastatic breast cancer The patient discontinued lapatinib within 3 months due to intolerable side effects and the cancer did not progress whilst on lapatinib He cancer has not progressed at any time point during the previous 12 months whilst on trastuzumab Trastuzumab will not be given in combination with pertuzumab Trastuzumab to be administered in combination with pertuzumab Patient has not received prior treatment for their metastatic disease and has had a treatment-free interval of at least 12 months between prior (neo)adjuvant chemotherapy treatment and diagnosis of metastatic breast cancer
and	The patient has good performance status (ECOG grade 0-1) Trastuzumab to be discontinued at disease progression
or and	Patient has previously discontinued treatment with trastuzumab in the metastatic setting for reasons other than severe toxicity or disease progression Patient has signs of disease progression Disease has not progressed during previous treatment with trastuzumab

I confirm that the above details are correct:

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Signed.	I Jata:	
Jigrieu		

I confirm that the above details are correct:

Signed: Date:

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July 2025

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e:	Name:		
l:	NHI:		
tuzumab	(Herzuma) - continued		
TIATION - massessment	etastatic breast cancer required after 12 months ick boxes where appropriate)		
and 1	The patient has metastatic breast cancer expressing HER-2 IHC 3+ or ISH+ (including FISH or other current technology)		
The patient has not previously received lapatinib treatment for HER-2 positive metastatic breast cancer The patient discontinued lapatinib within 3 months due to intolerable side effects and the cancer did not progress whilst on lapatinib			
and	Trastuzumab will not be given in combination with pertuzumab		
or	Trastuzumab to be administered in combination with pertuzumab Patient has not received prior treatment for their metastatic disease and has had a treatment-free interval of at least 12 months between prior (neo)adjuvant chemotherapy treatment and diagnosis of metastatic breast cancer		
and	The patient has good performance status (ECOG grade 0-1)		
ATINUATION assessment	The patient has good performance status (ECOG grade 0-1) Frastuzumab to be discontinued at disease progression I – metastatic breast cancer required after 12 months ick boxes where appropriate)		
NTINUATION assessment	Trastuzumab to be discontinued at disease progression N – metastatic breast cancer required after 12 months		
NTINUATION assessment requisites (t	Trastuzumab to be discontinued at disease progression I – metastatic breast cancer required after 12 months ick boxes where appropriate) O The patient has metastatic breast cancer expressing HER-2 IHC 3+ or ISH+ (including FISH or other current technology) O The cancer has not progressed at any time point during the previous 12 months whilst on trastuzumab		
NTINUATION assessment requisites (the and and and and and and and and and assessment ass	Trastuzumab to be discontinued at disease progression N - metastatic breast cancer required after 12 months ick boxes where appropriate) The patient has metastatic breast cancer expressing HER-2 IHC 3+ or ISH+ (including FISH or other current technology) The cancer has not progressed at any time point during the previous 12 months whilst on trastuzumab Trastuzumab to be discontinued at disease progression Patient has previously discontinued treatment with trastuzumab for reasons other than severe toxicity or disease progression Patient has signs of disease progression		

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PRESCI	RIBER	PATIENT:				
Name:		Name:				
Ward: .		NHI:				
Trastuzumab (Herzuma) - continued						
CONTINUATION – gastric, gastro-oesophageal junction and oesophageal cancer Re-assessment required after 12 months						
	uisites (tick boxes where appropriate)					
	O The cancer has not progressed at any time point during the previous 12 months whilst on trastuzumab					
a	Trastuzumab to be discontinued at disease progression					