

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](#).

**PRESCRIBER**

Name: .....

Ward: .....

**PATIENT:**

Name: .....

NHI: .....

**Trastuzumab (Herzuma)**

**INITIATION – early breast cancer**

Re-assessment required after 12 months

**Prerequisites** (tick boxes where appropriate)

- ☐ The patient has early breast cancer expressing HER-2 IHC 3+ or ISH + (including FISH or other current technology  
and  
☐ Maximum cumulative dose of 106 mg/kg (12 months' treatment)

**CONTINUATION – early breast cancer\***

Re-assessment required after 12 months

**Prerequisites** (tick boxes where appropriate)

- ☐ The patient has metastatic breast cancer expressing HER-2 IHC 3+ or ISH+ (including FISH or other current technology  
and  
☐ The patient received prior adjuvant trastuzumab treatment for early breast cancer  
and  
☐ The patient has not previously received lapatinib treatment for HER-2 positive metastatic breast cancer  
or  
☐ The patient discontinued lapatinib within 3 months due to intolerable side effects and the cancer did not progress whilst on lapatinib  
or  
☐ The cancer has not progressed at any time point during the previous 12 months whilst on trastuzumab  
and  
☐ Trastuzumab will not be given in combination with pertuzumab  
or  
☐ Trastuzumab to be administered in combination with pertuzumab  
and  
☐ 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)  
and  
☐ Trastuzumab to be discontinued at disease progression  
or  
☐ Patient has previously discontinued treatment with trastuzumab in the metastatic setting for reasons other than severe toxicity or disease progression  
and  
☐ Patient has signs of disease progression  
and  
☐ Disease has not progressed during previous treatment with trastuzumab

Note: \* For patients with relapsed HER-2 positive disease who have previously received adjuvant trastuzumab for early breast cancer

I confirm that the above details are correct:

Signed: ..... Date: .....

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](#).

**PRESCRIBER**

Name: .....

Ward: .....

**PATIENT:**

Name: .....

NHI: .....

**Trastuzumab (Herzuma) - continued**

**INITIATION – metastatic breast cancer**

Re-assessment required after 12 months

**Prerequisites** (tick boxes where appropriate)

- ☐ The patient has metastatic breast cancer expressing HER-2 IHC 3+ or ISH+ (including FISH or other current technology)
- and
- ☐ The patient has not previously received lapatinib treatment for HER-2 positive metastatic breast cancer
- or
- ☐ 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
- and
- ☐ 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)
- and
- ☐ Trastuzumab to be discontinued at disease progression

**CONTINUATION – metastatic breast cancer**

Re-assessment required after 12 months

**Prerequisites** (tick boxes where appropriate)

- ☐ The patient has metastatic breast cancer expressing HER-2 IHC 3+ or ISH+ (including FISH or other current technology)
- and
- ☐ The cancer has not progressed at any time point during the previous 12 months whilst on trastuzumab
- and
- ☐ Trastuzumab to be discontinued at disease progression
- or
- ☐ Patient has previously discontinued treatment with trastuzumab for reasons other than severe toxicity or disease progression
- and
- ☐ Patient has signs of disease progression
- and
- ☐ Disease has not progressed during previous treatment with trastuzumab

**INITIATION – gastric, gastro-oesophageal junction and oesophageal cancer**

Re-assessment required after 12 months

**Prerequisites** (tick boxes where appropriate)

- ☐ The patient has locally advanced or metastatic gastric, gastro-oesophageal junction or oesophageal cancer expressing HER-2 IHC 2+ FISH+ or IHC3+ (or other current technology)
- and
- ☐ Patient has an ECOG score of 0-2

I confirm that the above details are correct:

Signed: ..... Date: .....

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](#).

**PRESCRIBER**

Name: .....

Ward: .....

**PATIENT:**

Name: .....

NHI: .....

**Trastuzumab (Herzuma) - continued**

**CONTINUATION – gastric, gastro-oesophageal junction and oesophageal cancer**

Re-assessment required after 12 months

**Prerequisites** (tick boxes where appropriate)

- ☐ The cancer has not progressed at any time point during the previous 12 months whilst on trastuzumab  
**and**  
☐ Trastuzumab to be discontinued at disease progression

I confirm that the above details are correct:

Signed: ..... Date: .....