

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: .....

**Pertuzumab with trastuzumab**

**INITIATION**

Re-assessment required after 12 months

**Prerequisites** (tick boxes where appropriate)

- ☐ The individual has received an initial Special Authority approval for intravenous pertuzumab and trastuzumab for metastatic breast cancer
- and
- ☐ Pertuzumab with trastuzumab to be administered subcutaneously at a maximum dose of 600 mg pertuzumab with 600 mg trastuzumab every three weeks (or equivalent)

or

- ☐ The patient has metastatic breast cancer expressing HER-2 IHC 3+ or ISH+ (including FISH or other current technology)
- and
- ☐ Patient is chemotherapy treatment naïve
- or
- ☐ 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
- ☐ Loading dose of pertuzumab with trastuzumab to be administered subcutaneously at a maximum dose of 1200 mg pertuzumab with 600 mg trastuzumab, respectively
- and
- ☐ Maintenance doses of pertuzumab with trastuzumab to be administered subcutaneously at a maximum dose of 600 mg pertuzumab with 600 mg trastuzumab every three weeks (or equivalent)
- and
- ☐ Pertuzumab with trastuzumab to be discontinued at disease progression

**CONTINUATION**

Re-assessment required after 12 months

**Prerequisites** (tick boxes where appropriate)

- ☐ The individual 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 pertuzumab and trastuzumab

or

- ☐ Individual has previously discontinued treatment with pertuzumab with trastuzumab for reasons other than severe toxicity or disease progression
- and
- ☐ Individual has signs of disease progression
- and
- ☐ Disease has not progressed during previous treatment with pertuzumab with trastuzumab

I confirm that the above details are correct:

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