Trastuzumab (IV) Monotherapy - 7 days

INDICATIONS FOR USE:

<table>
<thead>
<tr>
<th>INDICATION</th>
<th>ICD10</th>
<th>Regimen Code</th>
<th>Reimbursement Status</th>
</tr>
</thead>
<tbody>
<tr>
<td>Treatment of patients with HER2 positive metastatic breast cancer (MBC)</td>
<td>C50</td>
<td>00201a</td>
<td>Hospital</td>
</tr>
</tbody>
</table>

TREATMENT:

The starting dose of the drugs detailed below may be adjusted downward by the prescribing clinician, using their independent medical judgement, to consider each patient's individual clinical circumstances.

MBC: Treatment administered every 7 days unless unacceptable toxicity develops.

Facilities to treat anaphylaxis MUST be present when trastuzumab is administered.

Cycle 1 For NEW patients ONLY.

Omit for patients continuing single-agent trastuzumab following a trastuzumab containing chemotherapy regimen

<table>
<thead>
<tr>
<th>Day</th>
<th>Drug</th>
<th>Dose</th>
<th>Route</th>
<th>Diluent &amp; Rate</th>
<th>Cycle</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Trastuzumab</td>
<td>4mg/kg</td>
<td>IV infusion Observe post infusion*</td>
<td>250ml 0.9% sodium chloride** over 90min</td>
<td>1</td>
</tr>
</tbody>
</table>

Cycle 2 and subsequent cycles or for patients who have just completed a trastuzumab containing chemotherapy regimen

<table>
<thead>
<tr>
<th>Day</th>
<th>Drug</th>
<th>Dose</th>
<th>Route</th>
<th>Diluent &amp; Rate</th>
<th>Cycle</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Trastuzumab</td>
<td>2mg/kg</td>
<td>IV infusion Observe post infusion*</td>
<td>250ml 0.9% sodium chloride over 30min</td>
<td>2 and further cycles</td>
</tr>
</tbody>
</table>

*Recommended Observation period: Patients should be observed for at least six hours after the start of the first infusion and for two hours after the start of the subsequent infusions for symptoms like fever and chills or other infusion-related symptoms. Any deviation should be noted in local policies.

** Trastuzumab is incompatible with glucose solution

ELIGIBILITY:

- Indications as above
- HER2 overexpression or HER2 gene amplification as determined by an accurate and validated assay
- ECOG status 0-3

EXCLUSIONS:

- Clinically significant cardiac disease (history of symptomatic ventricular arrhythmias, congestive heart failure or myocardial infarction within previous 12 months).
NCCP Chemotherapy Regimen

- Hypersensitivity to trastuzumab or any of the excipients.
- Patients experiencing dyspnoea at rest due to complications of advanced malignancy and comorbidities may be at increased risk of a fatal infusion reaction.

**PRESCRIPTIVE AUTHORITY:**
The treatment plan must be initiated by a Consultant Medical Oncologist

**TESTS:**

**Baseline tests:**
- FBC
- Cardiac function (LVEF using ECHO or MUGA scan)

**Regular tests:**
- FBC every 6 weeks.
- Cardiac function, liver profile, creatinine every 12 weeks. Where there are signs of cardiac impairment four to eight weekly checks may be more appropriate

**Disease monitoring:**
Disease monitoring should be in line with the patient’s treatment plan and any other test/s as directed by the supervising Consultant.

**DOSE MODIFICATIONS:**
- Any dose modification should be discussed with a Consultant.
- None usually recommended. Discontinue if unacceptable toxicity occurs
- Weight monitored at each visit and dose adjusted accordingly
- If the patient misses a dose of trastuzumab by one week or less, then the usual maintenance dose of 2 mg/kg should be given as soon as possible. Do not wait until the next planned cycle. Subsequent maintenance doses should then be given according to the previous schedule.
- If the patient misses a dose of trastuzumab by more than one week, a re-loading dose of trastuzumab (4 mg/kg) should be given over approximately 90 minutes, at the discretion of the clinician. Subsequent trastuzumab maintenance doses (2 mg/kg) should then be given weekly from that point.
Renal and Hepatic Impairment:

Table 1: Dose modification of trastuzumab in renal and hepatic impairment

<table>
<thead>
<tr>
<th>Renal Impairment</th>
<th>Hepatic Impairment</th>
</tr>
</thead>
<tbody>
<tr>
<td>No dedicated studies of trastuzumab in patients with renal impairment have been conducted.</td>
<td>No dedicated studies of trastuzumab in patients with hepatic impairment have been conducted. Probably no dose reduction necessary.</td>
</tr>
<tr>
<td>Based on a population pharmacokinetic (PK) analysis renal impairment was not shown to affect trastuzumab disposition.</td>
<td></td>
</tr>
</tbody>
</table>

Management of adverse events:

Table 2: Dose Modification of trastuzumab for Adverse Events

<table>
<thead>
<tr>
<th>Adverse reactions</th>
<th>Recommended dose modification</th>
</tr>
</thead>
<tbody>
<tr>
<td>LVEF drops 10 ejection fraction points from baseline and to below 50%</td>
<td>Withhold treatment. Repeat LVEF after 3 weeks. No improvement or further decline, consider discontinuation. Discuss with consultant and refer to cardiologist.</td>
</tr>
<tr>
<td>Symptomatic heart failure</td>
<td>Discontinue</td>
</tr>
<tr>
<td>NCI-CTCAE Grade 4 hypersensitivity reactions</td>
<td>Discontinue</td>
</tr>
<tr>
<td>Haematological</td>
<td>Treatment may continue during periods of reversible, chemotherapy-induced myelosuppression. Monitor carefully for any complications of neutropenia.</td>
</tr>
</tbody>
</table>

**SUPPORTIVE CARE:**

**EMETOGENIC POTENTIAL:** Minimal *(Refer to local policy).*

**PREMEDICATIONS:**

Not usually required unless the patient has had a previous hypersensitivity. Paracetamol and antihistamine cover should be considered. Patient should be educated about the possibility of delayed infusion-related symptoms

**OTHER SUPPORTIVE CARE:** No specific recommendations

**ADVERSE EFFECTS / REGIMEN SPECIFIC COMPLICATIONS:**

The adverse effects listed are not exhaustive. Please refer to the relevant Summary of Product Characteristics for full details.

- **Cardiac toxicity:**
  - Trastuzumab has been associated with moderate to severe cardiac failure. Baseline and 3 monthly cardiac function tests are required during treatment especially for those with prior anthracycline exposure.
  - If LVEF drops by greater than or equal to 10 ejection fraction (EF) points from baseline AND to below 50%, treatment should be withheld and a repeat LVEF assessment carried out within approximately 3 weeks. If LVEF has not improved, or declined further, discontinuation of trastuzumab should be strongly considered, unless the benefits for the individual patient are
Trastuzumab and anthracyclines should not be given concurrently in combination due to cardiotoxicity risk.

- The half-life of trastuzumab is approximately 4-5 weeks.

**Trastuzumab infusion-associated symptoms**, usually chills and fever may occur. Stop infusion and consider antihistamine cover. When symptoms have resolved the infusion may be recommenced. For serious reactions, discontinue the trastuzumab infusion and provide supportive therapy such as oxygen, beta-agonists and corticosteroids.

**Pulmonary events:** Severe pulmonary adverse reactions occur in association with the use of trastuzumab and have been associated with a fatal outcome. These events may occur as part of an infusion-related reaction or with a delayed onset. Caution should be exercised for pneumonitis, especially in patients being treated concomitantly with taxanes.

**DRUG INTERACTIONS:**

- A possible interaction with warfarin has been reported. An increased INR and bleeding may occur in patients previously stabilized on warfarin. The interaction was noted in two patients after 8-10 doses of trastuzumab. An INR prior to starting the trastuzumab is recommended, then every 2 weeks for the first 3 months and then monthly if stable. Inform patient to watch for any bleeding. Modification of the warfarin dose may be needed (1).

**Current drug interaction databases should be consulted for more information.**

**ATC CODE:**

Trastuzumab - L01XC03

**REFERENCES:**

1. Nissenblatt MJ. Karp Gl. Bleeding risk with trastuzumab (Herceptin) treatment JAMA 1999;282:2299-301
NCCP Chemotherapy Regimen


<table>
<thead>
<tr>
<th>Version</th>
<th>Date</th>
<th>Amendment</th>
<th>Approved By</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>10/02/2014</td>
<td></td>
<td>Dr Maccon Keane</td>
</tr>
<tr>
<td>2</td>
<td>10/02/2016</td>
<td>Added Disease Monitoring</td>
<td>Dr Maccon Keane</td>
</tr>
<tr>
<td>3</td>
<td>07/02/2018</td>
<td>Updated infusion time recommendations and emetogenic potential. Clarification of dosing in renal and hepatic impairment. Formatting in new NCCP Regimen Template</td>
<td>Prof Maccon Keane</td>
</tr>
<tr>
<td>4</td>
<td>15/01/2020</td>
<td>Reviewed</td>
<td>Prof Maccon Keane</td>
</tr>
</tbody>
</table>

Comments and feedback welcome at oncologydrugs@cancercontrol.ie.

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Tumour Group: Breast
NCCP Regimen Code: 00201
ISM0 Contributor: Prof Maccon Keane
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Version number: 4

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