Trastuzumab in combination with Docetaxel - Metastatic Breast Cancer

(This form must be completed before the first dose is dispensed.)
1. Patient Profile

- Surname: ___________________________________________
- Given Name: _______________________________________
- OHIN: ___________________________  
  Chart Number: ___________________________
- Postal Code: ___________________________
- Height (cm): _______  
  Weight (kg): _______
- BSA (m²): _________  
  Gender:  
  ○ Male  
  ○ Female  
  ○ Other
- Date of Birth: _______ _______ _______
  Day  Month  Year
- Site:
- Attending Physician (MRP- Most Responsible Physician): ___________________________________________

  Requested Prior Approval  □ Yes  
  Patient on Clinical Trial  ○ Yes  ○ No

  Other (specify): ___________________________________________

  Specify Arm:
  ○ Standard of care arm  
  ○ Experimental arm  
  ○ Blinded / Unknown

Request prior approval for enrolment

- Justification for Funding


2. Eligibility Criteria

a. Her2/neu Status

Patient must test positive for Her2/neu as per Cancer Care Ontario criteria. A photocopy of the pathology report of the Her2/neu test must be submitted to Cancer Care Ontario. The report must state clearly the...
report of the Her2/neu test must be submitted to Cancer Care Ontario. The report must state clearly the hospital, date of biopsy and the hospital pathology specimen number of the original material used for the Her2/neu test.

Please specify tests used for detection of Her2/neu:

- IHC 3+  [ ] Yes
- FISH/SISH >= 2  [ ] Yes

b. Combination treatment for patients receiving trastuzumab and docetaxel.

i. Patient has metastatic breast cancer.  [ ] Yes

   AND

ii. [ ] cannot tolerate anthracyclines
   [ ] has failed anthracycline therapy for metastatic disease.
   [ ] has received an anthracycline as adjuvant therapy.

3. Precautions

a. The patient has a normal cardiac ejection fraction  [ ] Yes

b. There is no evidence of extensive lung involvement  [ ] Yes

4. Funded Dose

- Therapy should be initiated with a loading dose of 4 mg/kg, followed by 2 mg/kg IV weekly.
- Patients may be switched to 6 mg/kg IV q3 weeks after they are adequately loaded over a reasonable period of time with weekly dosing.

5. Notes

a. Reimbursement will be discontinued for patients whose disease progresses while being treated with trastuzumab in the metastatic setting.

6. Supporting Documents

To ensure reimbursement of your claim, both the completed enrolment form and a copy of the required documentation (where applicable) must be submitted through CCO e-Claims.