

Regimen Monograph

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A - Regimen Name

# PACL(W)+PERT+TRAS Regimen

PACLitaxel (weekly)-Pertuzumab-Trastuzumab

**Disease Site** Breast

**Intent** Curative

**Regimen Category** **Evidence-informed :**

Regimen is considered appropriate as part of the standard care of patients; meaningfully improves outcomes (survival, quality of life), tolerability or costs compared to alternatives (recommended by the Disease Site Team and national consensus body e.g. pan-Canadian Oncology Drug Review, pCODR). Recommendation is based on an appropriately conducted phase III clinical trial relevant to the Canadian context OR (where phase III trials are not feasible) an appropriately sized phase II trial. Regimens where one or more drugs are not approved by Health Canada for any indication will be identified under Rationale and Use.

This **Regimen Abstract** is an **abbreviated** version of a Regimen Monograph and contains only top level information on usage, dosing, schedule, cycle length and special notes (if available). Information in regimen abstracts is accurate to the extent of the ST-QBP regimen master listings, and has not undergone the full review process of a regimen monograph. Full regimen monographs will be published for each ST-QBP regimen as they are developed.

**Rationale and Uses** Neoadjuvant treatment of early stage HER2-positive breast cancer

**Supplementary Public Funding** [trastuzumab](#)  
New Drug Funding Program (Trastuzumab (Biosimilar) - Adjuvant Treatment for Breast Cancer) ([NDFP Website](#))

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**B - Drug Regimen**

**Note:** Different trastuzumab products are **NOT INTERCHANGEABLE**.

**Cycle 1 - Trastuzumab and Pertuzumab Loading Dose:**

[PERTuzumab](#)<sup>1</sup>                      840 mg                      IV over 60 minutes                      Day 1

(This drug is not currently publicly funded for this regimen and intent)

*Then,*

[trastuzumab](#)<sup>1</sup>                      8 mg /kg                      IV over 90 minutes                      Day 1

*Then,*

[PACLitaxel](#)<sup>1</sup>                      80 mg /m<sup>2</sup>                      IV over 1 hour                      Days 1, 8, 15

**Cycle 2 and Onwards - Trastuzumab and Pertuzumab Maintenance Dose (Q3W):**

[PERTuzumab](#)<sup>1, 2</sup>                      420 mg                      IV over 30\* to 60 minutes                      Day 1

(This drug is not currently publicly funded for this regimen and intent)

(\* if previous 60-minute infusion well-tolerated)

*Then,*

[trastuzumab](#)<sup>1, 2</sup>                      6 mg /kg                      IV over 30 minutes\*\*                      Day 1

(\*\* if previous 90-minute infusion well-tolerated)

*Then,*

[PACLitaxel](#)<sup>1</sup>                      80 mg /m<sup>2</sup>                      IV over 1 hour                      Days 1, 8, 15

(1) Based on the product monograph, pertuzumab and trastuzumab may be administered in any order; however, the taxane should be given after pertuzumab and trastuzumab.

(2) If delayed by ≥ 3 weeks (i.e. ≥ 6 weeks from last dose), re-load with loading dose.

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## C - Cycle Frequency

### REPEAT EVERY 21 DAYS

For 4 cycles, unless disease progression or unacceptable toxicity

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## D - Premedication and Supportive Measures

**Antiemetic Regimen:** Low

- Also refer to [CCO Antiemetic Recommendations](#).

**Screen for hepatitis B virus in all cancer patients starting systemic treatment.** Refer to the [hepatitis B virus screening and management](#) guideline.

**Pre-medications (prophylaxis for infusion reaction):**

PACLitaxel:

To be given 30-60 minutes prior to PACLitaxel infusion:

- Dexamethasone 10 mg IV, starting in cycle 1
- Diphenhydramine 25-50 mg IV/PO
- Ranitidine 50 mg IV OR Famotidine 20 mg IV

\* Consider discontinuing pre-medications for PACLitaxel if there was no IR in the first 2 doses.

Pertuzumab and Trastuzumab

- No specific premedications recommended.
- For patients who experienced prior infusion reactions, consider premedication with corticosteroids, antihistamines, and/or antipyretics before subsequent infusions.

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## J - Administrative Information

Approximate Patient Visit	3 to 5 hours (Day 1); 2 hours (paclitaxel-only days)
Pharmacy Workload (average time per visit)	23.58 minutes
Nursing Workload (average time per visit)	53.167 minutes

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## K - References

Nitz UA, Gluz O, Christgen M, et al. De-escalation strategies in HER2-positive early breast cancer (EBC): final analysis of the WSG-ADAPT HER2+/HR- phase II trial: efficacy, safety, and predictive markers for 12 weeks of neoadjuvant dual blockade with trastuzumab and pertuzumab ± weekly paclitaxel. *Ann Oncol.* 2017 Nov 1;28(11):2768-72.

Paclitaxel drug monograph, Ontario Health (Cancer Care Ontario).

Pertuzumab drug monograph, Ontario Health (Cancer Care Ontario).

Reimbursement recommendation: pertuzumab. Canada's Drug Agency, October 2025.

Trastuzumab drug monograph, Ontario Health (Cancer Care Ontario).

**April 2026** new ST-QBP regimen

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## M - Disclaimer

### **Regimen Abstracts**

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### **Regimen Monographs**

Refer to the [New Drug Funding Program](#) or [Ontario Public Drug Programs](#) websites for the most up-to-date public funding information.

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