

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use POHERDY safely and effectively. See full prescribing information for POHERDY.

POHERDY® (pertuzumab-dpzb) injection, for intravenous use

Initial U.S. Approval: 2025

POHERDY (pertuzumab-dpzb) is biosimilar* to PERJETA® (pertuzumab).

WARNING: LEFT VENTRICULAR DYSFUNCTION and EMBRYO-FETAL TOXICITY

See full prescribing information for complete boxed warning.

- **Left Ventricular Dysfunction:** Pertuzumab products can cause subclinical and clinical cardiac failure manifesting as decreased LVEF and CHF. Evaluate cardiac function prior to and during treatment. Discontinue POHERDY treatment for a confirmed clinically significant decrease in left ventricular function. (2.3, 5.1, 6.1)
- **Embryo-fetal Toxicity:** Exposure to pertuzumab products can cause embryo-fetal death and birth defects. Advise patients of these risks and the need for effective contraception. (5.2, 8.1, 8.3)

INDICATIONS AND USAGE

POHERDY is a HER2/neu receptor antagonist indicated for:

- Use in combination with trastuzumab and docetaxel for treatment of adults with HER2-positive metastatic breast cancer (MBC) who have not received prior anti-HER2 therapy or chemotherapy for metastatic disease. (1.1)
- Use in combination with trastuzumab and chemotherapy as
 - neoadjuvant treatment of adults with HER2-positive, locally advanced, inflammatory, or early stage breast cancer (either greater than 2 cm in diameter or node positive) as part of a complete treatment regimen for early breast cancer. (1.2, 2.2, 14.2)
 - adjuvant treatment of adults with HER2-positive early breast cancer at high risk of recurrence (1.2, 2.2, 14.3)

DOSAGE AND ADMINISTRATION

- **For intravenous infusion only.** Do not administer as an intravenous push or bolus. (2.6)
- HER2 testing: Perform using FDA-approved tests by laboratories with demonstrated proficiency. (2.1)
- The initial POHERDY dose is 840 mg administered as a 60-minute intravenous infusion, followed every 3 weeks thereafter by 420 mg administered as a 30 to 60 minute intravenous infusion. (2.3)
- MBC: Administer POHERDY, trastuzumab or trastuzumab hyaluronidase-oysk, and docetaxel every 3 weeks. (2.3)
- Neoadjuvant: Administer POHERDY, trastuzumab or trastuzumab hyaluronidase-oysk, and chemotherapy preoperatively every 3 weeks for 3 to 6 cycles. (2.3)
- Adjuvant: Administer POHERDY, trastuzumab or trastuzumab hyaluronidase-oysk, and chemotherapy postoperatively every 3 weeks for a total of 1 year (up to 18 cycles). (2.3)

DOSAGE FORMS AND STRENGTHS

- Injection: 420 mg/14 mL (30 mg/mL) solution in a single-dose vial. (3)

CONTRAINDICATIONS

POHERDY is contraindicated in patients with known hypersensitivity to pertuzumab products or to any of its excipients. (4)

WARNINGS AND PRECAUTIONS

- **Infusion-Related Reactions:** Pertuzumab products can cause serious infusion reactions, including fatal events. Monitor for signs and symptoms. If a significant infusion-associated reaction occurs, slow or interrupt the infusion and administer appropriate medical therapies. (5.3)
- **Hypersensitivity Reactions/Anaphylaxis:** Pertuzumab products can cause hypersensitivity reactions, including anaphylaxis. Monitor for signs and symptoms, including angioedema. If a severe hypersensitivity reaction/anaphylaxis occurs, discontinue the infusion immediately and administer appropriate medical therapies. (5.4)

ADVERSE REACTIONS

Metastatic Breast Cancer

- The most common adverse reactions (> 30%) with pertuzumab in combination with trastuzumab and docetaxel were diarrhea, alopecia, neutropenia, nausea, fatigue, rash, and peripheral neuropathy. (6.1)

Neoadjuvant Treatment of Breast Cancer

- The most common adverse reactions (> 30%) with pertuzumab in combination with trastuzumab and docetaxel were alopecia, diarrhea, nausea, and neutropenia. (6.1)
- The most common adverse reactions (>30%) with pertuzumab in combination with trastuzumab and docetaxel when given for 3 cycles following 3 cycles of FEC were fatigue, alopecia, diarrhea, nausea, vomiting, and neutropenia. (6.1)
- The most common adverse reactions (>30%) with pertuzumab in combination with docetaxel, carboplatin, and trastuzumab (TCH) were fatigue, alopecia, diarrhea, nausea, vomiting, neutropenia, thrombocytopenia, and anemia. (6.1)
- The most common adverse reactions (>30%) with pertuzumab in combination with trastuzumab and paclitaxel when given for 4 cycles following 4 cycles of ddAC were nausea, diarrhea, alopecia, fatigue, constipation, peripheral neuropathy, and headache. (6.1)
- The most common adverse reactions (>30%) with pertuzumab in combination with trastuzumab and docetaxel when given for 4 cycles following 4 cycles of FEC were diarrhea, nausea, alopecia, asthenia, constipation, fatigue, mucosal inflammation, vomiting, myalgia, and anemia. (6.1)

Adjuvant Treatment of Breast Cancer

- The most common adverse reactions (>30%) with pertuzumab in combination with trastuzumab and chemotherapy were diarrhea, nausea, alopecia, fatigue, peripheral neuropathy and vomiting. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Organon LLC, a subsidiary of Organon & Co., at 1-844-674-3200 or FDA at 1-800-FDA-1088 or <http://www.fda.gov/medwatch>.

USE IN SPECIFIC POPULATIONS

Females and Males of Reproductive Potential: Verify the pregnancy status of females prior to initiation of POHERDY. (8.3)

See 17 for PATIENT COUNSELING INFORMATION.

*Biosimilar means that the biological product is approved based on data demonstrating that it is highly similar to an FDA-approved biological product, known as a reference product, and that there are no clinically meaningful differences between the biosimilar product and the reference product. Biosimilarity of POHERDY has been demonstrated for the condition(s) of use (e.g., indication(s), dosing regimen(s)), strength(s), dosage form(s), and route(s) of administration) described in its Full Prescribing Information.

Revised: 11/2025

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FULL PRESCRIBING INFORMATION

WARNING: LEFT VENTRICULAR DYSFUNCTION and EMBRYO-FETAL TOXICITY

- **Left Ventricular Dysfunction:** Pertuzumab products can cause subclinical and clinical cardiac failure manifesting as decreased LVEF and CHF. Evaluate cardiac function prior to and during treatment. Discontinue POHERDY treatment for a confirmed clinically significant decrease in left ventricular function [see *Dosage and Administration (2.3)*, *Warnings and Precautions (5.1)* and *Adverse Reactions (6.1)*].
- **Embryo-fetal Toxicity:** Exposure to pertuzumab products can cause embryo-fetal death and birth defects. Advise patients of these risks and the need for effective contraception [see *Warnings and Precautions (5.2)* and *Use in Specific Populations (8.1) (8.3)*].

1 INDICATIONS AND USAGE

1.1 Metastatic Breast Cancer (MBC)

POHERDY is indicated for use in combination with trastuzumab and docetaxel for the treatment of adults with HER2-positive metastatic breast cancer who have not received prior anti-HER2 therapy or chemotherapy for metastatic disease [see *Dosage and Administration (2.2)* and *Clinical Studies (14.1)*].

1.2 Early Breast Cancer (EBC)

POHERDY is indicated for use in combination with trastuzumab and chemotherapy for

- the neoadjuvant treatment of adults with HER2-positive, locally advanced, inflammatory, or early stage breast cancer (either greater than 2 cm in diameter or node positive) as part of a complete treatment regimen for early breast cancer [see *Dosage and Administration (2.2)* and *Clinical Studies (14.2)*].
- the adjuvant treatment of adults with HER2-positive early breast cancer at high risk of recurrence [see *Dosage and Administration (2.2)* and *Clinical Studies (14.3)*].

2 DOSAGE AND ADMINISTRATION

2.1 Evaluation and Testing Before Initiating POHERDY

Assess left ventricular ejection fraction (LVEF) prior to initiation of POHERDY and at regular intervals during treatment [see *Boxed Warning*, *Dosage and Administration (2.4)*, *Warnings and Precautions (5.1)*].

Verify the pregnancy status of females of reproductive potential prior to the initiation of POHERDY [see *Warnings and Precautions (5.2)*, *Use in Specific Populations (8.1, 8.3)*].

2.2 Patient Selection

Select patients based on HER2 protein overexpression or HER2 gene amplification in tumor specimens [see *Indications and Usage (1)* and *Clinical Studies (14)*]. Assessment of HER2

protein overexpression and HER2 gene amplification should be performed using FDA-approved tests specific for breast cancer by laboratories with demonstrated proficiency. Information on the FDA-approved tests for the detection of HER2 protein overexpression and HER2 gene amplification is available at: <http://www.fda.gov/CompanionDiagnostics>.

Improper assay performance, including use of suboptimally fixed tissue, failure to utilize specified reagents, deviation from specific assay instructions, and failure to include appropriate controls for assay validation, can lead to unreliable results.

2.3 Recommended Dosage and Administration

The initial dose of POHERDY is 840 mg administered as a 60-minute intravenous infusion, followed every 3 weeks by a dose of 420 mg administered as an intravenous infusion over 30 to 60 minutes.

When administered with POHERDY, the recommended initial dose of trastuzumab is 8 mg/kg administered as a 90-minute intravenous infusion, followed every 3 weeks by a dose of 6 mg/kg administered as an intravenous infusion over 30 to 90 minutes.

When administered with POHERDY, the recommended initial dose of trastuzumab hyaluronidase-oysk is 600 mg/10,000 units (600 mg trastuzumab and 10,000 units hyaluronidase) administered subcutaneously over approximately 2 to 5 minutes once every three weeks irrespective of the patient's body weight.

Administer POHERDY, trastuzumab or trastuzumab hyaluronidase-oysk, and taxane sequentially. POHERDY and trastuzumab or trastuzumab hyaluronidase-oysk can be given in any order. Administer taxane after POHERDY and trastuzumab or trastuzumab hyaluronidase-oysk. An observation period of 30 to 60 minutes is recommended after each POHERDY infusion and before commencement of any subsequent administration of trastuzumab or trastuzumab hyaluronidase-oysk, or taxane [see *Warnings and Precautions* (5.3)].

In patients receiving an anthracycline-based regimen, administer POHERDY and trastuzumab or trastuzumab hyaluronidase-oysk after completion of the anthracycline.

Metastatic Breast Cancer (MBC)

When administered with POHERDY, the recommended initial dose of docetaxel is 75 mg/m² administered as an intravenous infusion. The dose may be escalated to 100 mg/m² administered every 3 weeks if the initial dose is well tolerated.

Neoadjuvant Treatment of Breast Cancer

Administer POHERDY every 3 weeks for 3 to 6 cycles as part of one of the following treatment regimens [see *Clinical Studies* (14.2, 14.3)]:

- Four preoperative cycles of POHERDY in combination with trastuzumab or trastuzumab hyaluronidase-oysk and docetaxel followed by 3 postoperative cycles of fluorouracil, epirubicin, and cyclophosphamide (FEC)
- Three or four preoperative cycles of FEC alone followed by 3 or 4 preoperative cycles of POHERDY in combination with docetaxel and trastuzumab or trastuzumab hyaluronidase-oysk
- Six preoperative cycles of POHERDY in combination with docetaxel, carboplatin, and

trastuzumab (TCH) or trastuzumab hyaluronidase-oysk (escalation of docetaxel above 75 mg/m² is not recommended)

- Four preoperative cycles of dose-dense doxorubicin and cyclophosphamide (ddAC) alone followed by 4 preoperative cycles of POHERDY in combination with paclitaxel and trastuzumab or trastuzumab hyaluronidase-oysk

Following surgery, administer POHERDY and trastuzumab or trastuzumab hyaluronidase-oysk to complete 1 year of treatment (up to 18 cycles) or until disease recurrence or unmanageable toxicity, whichever occurs first.

Adjuvant Treatment of Breast Cancer

As part of a regimen including standard anthracycline- and/or taxane-based chemotherapy, administer POHERDY in combination with trastuzumab or trastuzumab hyaluronidase-oysk every 3 weeks for a total of 1 year (up to 18 cycles) or until disease recurrence or unmanageable toxicity, whichever occurs first. Administer POHERDY on Day 1 of the first taxane-containing cycle [see *Clinical Studies* (14.3)].

2.4 Important Dosing Considerations

Missed Dose

The recommended dosage modifications for delayed or missed doses are listed in Table 1.

Table 1: Recommendations for Delayed or Missed Doses

Time between two sequential doses	POHERDY	Trastuzumab (intravenous)	Trastuzumab hyaluronidase-oysk
< 6 weeks	Administer POHERDY 420 mg intravenously as soon as possible. Do not wait until the next planned dose.	Administer trastuzumab 6 mg/kg intravenously as soon as possible. Do not wait until the next planned dose.	Administer trastuzumab hyaluronidase-oysk 600 mg/10,000 units subcutaneously as soon as possible.
≥ 6 weeks	Readminister POHERDY loading dose of 840 mg intravenously as a 60 minute infusion, followed by a maintenance dose of 420 mg administered intravenously over a period of 30 to 60 minutes every 3 weeks thereafter.	Readminister trastuzumab loading dose of 8 mg/kg intravenously over approximately 90 minutes, followed by a maintenance dose of 6 mg/kg administered intravenously over a period of 30 or 90 minutes every 3 weeks thereafter.	Do not wait until the next planned dose.

Permanently discontinue POHERDY if trastuzumab or trastuzumab hyaluronidase-oysk treatment is discontinued.

Dose reductions are not recommended for POHERDY.

For chemotherapy dose modifications, see relevant prescribing information.

2.5 Dosage Modification for Adverse Reactions

Left Ventricular Dysfunction

Assess left ventricular ejection fraction (LVEF) prior to initiation of POHERDY and at regular intervals during treatment. The recommended dosage modifications for LVEF decrease are listed in Table 2 [see *Warnings and Precautions* (5.1)].

Table 2: Dose Modifications for Left Ventricular Dysfunction

	Pre-treatment LVEF:	Monitor LVEF every:	Withhold POHERDY and trastuzumab or trastuzumab hyaluronidase-oysk for at least 3 weeks for an LVEF decrease to:		Resume POHERDY and trastuzumab or trastuzumab hyaluronidase-oysk after 3 weeks if LVEF has recovered to:	
Metastatic Breast Cancer	≥ 50%	~12 weeks	Either		Either	
			<40%	40%-45% with a fall of ≥10%-points below pre-treatment value	>45%	40%-45% with a fall of <10%-points below pre-treatment value
Early Breast Cancer	≥ 55%*	~12 weeks (once during neoadjuvant therapy)	<50% with a fall of ≥10%-points below pre-treatment value		Either	
					≥50%	<10% points below pre-treatment value

*For patients receiving anthracycline-based chemotherapy, a LVEF of ≥ 50% is required after completion of anthracyclines, before starting POHERDY and trastuzumab or trastuzumab hyaluronidase-oysk.

Infusion-Related Reactions

The infusion rate of POHERDY may be slowed or interrupted if the patient develops an infusion-related reaction [see *Warnings and Precautions* (5.3)].

Hypersensitivity Reactions/Anaphylaxis

The infusion should be discontinued immediately if the patient experiences a serious hypersensitivity reaction [see *Warnings and Precautions* (5.4)].

2.6 Preparation for Administration

Administer as an intravenous infusion only. Do not administer as an intravenous push or bolus. Do not mix POHERDY with other drugs.

Preparation

Prepare the solution for infusion, using aseptic technique, as follows:

- Parenteral drug products should be inspected visually for particulates and discoloration prior to administration, whenever solution and container permit.
- Withdraw the appropriate volume of POHERDY solution from the vial(s) using a sterile needle and syringe.
- Dilute into a 250 mL 0.9% Sodium Chloride Injection infusion bag.

- POHERDY is compatible with polyvinylchloride (PVC) or polypropylene infusion bags.
- Mix diluted solution by gentle inversion. Do not shake.
- Administer immediately once prepared.
- If the diluted infusion solution is not used immediately, it can be stored refrigerated at 2°C to 8°C (36°F to 46°F) for up to 24 hours.
- Dilute with 0.9% Sodium Chloride Injection only.
- Do not use 5% Dextrose Injection.

3 DOSAGE FORMS AND STRENGTHS

Injection: 420 mg/14 mL (30 mg/mL) clear to slightly opalescent, colorless to pale yellow solution in a single-dose vial.

4 CONTRAINDICATIONS

POHERDY is contraindicated in patients with known hypersensitivity to pertuzumab products or to any of its excipients [see *Warnings and Precautions* (5.4)].

5 WARNINGS AND PRECAUTIONS

5.1 Left Ventricular Dysfunction

Pertuzumab products can cause left ventricular dysfunction, including symptomatic heart failure. Decreases in LVEF have been reported with drugs that block HER2 activity, including pertuzumab products.

Assess LVEF prior to initiation of POHERDY and at regular intervals during treatment to ensure that LVEF is within normal limits. If the LVEF declines and has not improved, or has declined further at the subsequent assessment, consider permanent discontinuation of POHERDY and trastuzumab [see *Dosage Modification for Adverse Reactions* (2.5)].

In the pertuzumab-treated patients with MBC in CLEOPATRA, left ventricular dysfunction occurred in 4% of patients and symptomatic left ventricular systolic dysfunction (LVSD) (congestive heart failure) occurred in 1% of patients. Patients who received prior anthracyclines or prior radiotherapy to the chest area may be at higher risk of decreased LVEF or left ventricular dysfunction.

In patients receiving pertuzumab as a neoadjuvant treatment in combination with trastuzumab and docetaxel in NeoSphere, LVEF decline > 10% and a drop to < 50% occurred in 8% of patients and left ventricular dysfunction occurred in 3% of patients. LVEF recovered to ≥ 50% in all these patients.

In patients receiving neoadjuvant pertuzumab in TRYPHAENA, LVEF decline > 10% and a drop to < 50% occurred in 7% of patients treated with pertuzumab plus trastuzumab and FEC followed by pertuzumab plus trastuzumab and docetaxel, 16% of patients treated with pertuzumab plus trastuzumab and docetaxel following FEC, and 11% of patients treated with pertuzumab in combination with TCH. Left ventricular dysfunction occurred in 6% of patients treated with pertuzumab plus trastuzumab and FEC followed by pertuzumab plus trastuzumab and docetaxel, 4% of patients treated with pertuzumab plus trastuzumab and docetaxel following

FEC, and 3% of patients treated with pertuzumab in combination with TCH. Symptomatic LVSD occurred in 4% of patients treated with pertuzumab plus trastuzumab and docetaxel following FEC, 1% of patients treated with pertuzumab in combination with TCH, and none of the patients treated with pertuzumab plus trastuzumab and FEC followed by pertuzumab plus trastuzumab and docetaxel. LVEF recovered to $\geq 50\%$ in all but one patient.

In patients receiving neoadjuvant pertuzumab in BERENICE, in the neoadjuvant period, LVEF decline $\geq 10\%$ and a drop to $< 50\%$ as measured by ECHO/MUGA assessment occurred in 7% of patients treated with pertuzumab plus trastuzumab and paclitaxel following ddAC, and 2% of patients treated with pertuzumab plus trastuzumab and docetaxel following FEC. Ejection fraction decreased (asymptomatic LVD) occurred in 7% of patients treated with pertuzumab plus trastuzumab and paclitaxel following ddAC and 4% of the patients treated with pertuzumab plus trastuzumab and docetaxel following FEC in the neoadjuvant period. Symptomatic LVSD (NYHA Class III/IV Congestive Heart Failure) occurred in 2% of patients treated with pertuzumab plus trastuzumab and paclitaxel following ddAC and none of the patients treated with pertuzumab plus trastuzumab and docetaxel following FEC in the neoadjuvant period.

In patients receiving adjuvant pertuzumab in APHINITY, the incidence of symptomatic heart failure (NYHA Class III/IV) with a LVEF decline $\geq 10\%$ and a drop to $< 50\%$ was 0.6%. Of the patients who experienced symptomatic heart failure, 47% of pertuzumab-treated patients had recovered (defined as 2 consecutive LVEF measurements above 50%) at the data cutoff. The majority of the events (86%) were reported in anthracycline-treated patients. Asymptomatic or mildly symptomatic (NYHA Class II) declines in LVEF $\geq 10\%$ and a drop to $< 50\%$ were reported in 3% of pertuzumab-treated patients, of whom 80% recovered at the data cutoff.

Pertuzumab products have not been studied in patients with a pretreatment LVEF value of $< 50\%$, a prior history of CHF, decreases in LVEF to $< 50\%$ during prior trastuzumab therapy, or conditions that could impair left ventricular function such as uncontrolled hypertension, recent myocardial infarction, serious cardiac arrhythmia requiring treatment or a cumulative prior anthracycline exposure to $> 360 \text{ mg/m}^2$ of doxorubicin or its equivalent.

5.2 Embryo-Fetal Toxicity

Based on its mechanism of action and findings in animal studies, pertuzumab products can cause fetal harm when administered to a pregnant woman. Pertuzumab products are HER2/neu receptor antagonists. Cases of oligohydramnios and oligohydramnios sequence manifesting as pulmonary hypoplasia, skeletal abnormalities, and neonatal death have been reported with use of another HER2/neu receptor antagonist (trastuzumab) during pregnancy. In an animal reproduction study, administration of pertuzumab to pregnant cynomolgus monkeys during the period of organogenesis resulted in oligohydramnios, delayed fetal kidney development, and embryo-fetal death at exposures 2.5 to 20 times the exposure in humans at the recommended dose, based on C_{max} .

Verify the pregnancy status of females of reproductive potential prior to the initiation of POHERDY. Advise pregnant women and females of reproductive potential that exposure to POHERDY in combination with trastuzumab during pregnancy or within 7 months prior to conception can result in fetal harm, including embryo-fetal death or birth defects. Advise females of reproductive potential to use effective contraception during treatment and for 7 months following the last dose of POHERDY in combination with trastuzumab [see *Use in Specific*

Populations (8.1, 8.3)].

5.3 Infusion-Related Reactions

Pertuzumab products can cause serious infusion reactions, including fatal events [*see Adverse Reactions (6.1)]*.

In CLEOPATRA, on the first day, when only pertuzumab was administered, infusion-related reactions occurred in 13% of patients and <1% were Grade 3 or 4. The most common infusion reactions ($\geq 1\%$) were pyrexia, chills, fatigue, headache, asthenia, hypersensitivity, and vomiting. During the second cycle when all drugs were administered on the same day, the most common infusion reactions in the pertuzumab-treated group ($\geq 1\%$) were fatigue, dysgeusia, hypersensitivity, myalgia, and vomiting.

In APHINITY, when pertuzumab was administered in combination with trastuzumab and chemotherapy on the same day, infusion-related reactions occurred in 21% of patients with <1% of patients experiencing Grade 3-4 events.

Observe patients closely for 60 minutes after the first infusion and for 30 minutes after subsequent infusions of POHERDY. If a significant infusion-related reaction occurs, slow or interrupt the infusion, and administer appropriate medical therapies. Monitor patients carefully until complete resolution of signs and symptoms. Consider permanent discontinuation in patients with severe infusion reactions [*see Dosage and Administration (2.5)]*.

5.4 Hypersensitivity Reactions/Anaphylaxis

Pertuzumab products can cause hypersensitivity reactions, including anaphylaxis.

In CLEOPATRA, the overall frequency of hypersensitivity/anaphylaxis reactions was 11% in pertuzumab-treated patients, with Grade 3 – 4 hypersensitivity reactions and anaphylaxis occurring in 2% of patients.

In NeoSphere, TRYPHAENA, BERENICE, and APHINITY, hypersensitivity/anaphylaxis events were consistent with those observed in CLEOPATRA. In APHINITY, the overall frequency of hypersensitivity/anaphylaxis was 5% in the pertuzumab treated group. The incidence was highest in the pertuzumab plus TCH treated group (8%) with 1% Grade 3 – 4 events.

Observe patients closely for hypersensitivity reactions. Severe hypersensitivity, including anaphylaxis and fatal events, have been observed in patients treated with pertuzumab products [*see Clinical Trials Experience (6.1)]*. Angioedema has been described in post-marketing reports. Medications to treat such reactions, as well as emergency equipment, should be available for immediate use prior to administration of POHERDY. POHERDY is contraindicated in patients with known hypersensitivity to pertuzumab products or to any of its excipients [*see Contraindications (4)]*.

6 ADVERSE REACTIONS

The following adverse reactions are discussed in greater detail in other sections of the label:

- Left Ventricular Dysfunction [*see Warnings and Precautions (5.1)]*
- Embryo-Fetal Toxicity [*see Warnings and Precautions (5.2)]*

- Infusion-Related Reactions [see *Warnings and Precautions* (5.3)]
- Hypersensitivity Reactions/Anaphylaxis [see *Warnings and Precautions* (5.4)]

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in clinical practice.

Metastatic Breast Cancer (MBC)

CLEOPATRA

The safety of pertuzumab in combination with trastuzumab and docetaxel was evaluated in a randomized trial (CLEOPATRA) in patients with HER2-positive metastatic breast cancer [see *Clinical Studies* (14.1)]. Patients received either pertuzumab administered at an initial dose of 840 mg followed by 420 mg every 3 weeks thereafter or placebo in combination with trastuzumab (initial dose of 8 mg/kg, followed by 6 mg/kg every 3 weeks thereafter) and docetaxel (75 mg/m² by intravenous infusion every 3 weeks for 6 cycles). The median duration of study treatment was 18.1 months for patients in the pertuzumab-treated group.

Permanent discontinuation of pertuzumab, trastuzumab, and docetaxel due to adverse reactions occurred in 6% of patients. Adverse reactions that led to permanent discontinuation of pertuzumab, trastuzumab, and docetaxel in >1% of patients were left ventricular dysfunction.

The safety profile of pertuzumab remained unchanged with an additional 2.75 years of follow-up (median total follow-up of 50 months) in CLEOPATRA.

The most common adverse reactions (> 30%) with pertuzumab in combination with trastuzumab and docetaxel were diarrhea, alopecia, neutropenia, nausea, fatigue, rash, and peripheral neuropathy. The most common Grade 3 – 4 adverse reactions (> 2%) were neutropenia, febrile neutropenia, leukopenia, diarrhea, peripheral neuropathy, anemia, asthenia, and fatigue. An increased incidence of febrile neutropenia was observed for Asian patients in both treatment arms compared with patients of other races and from other geographic regions. Among Asian patients, the incidence of febrile neutropenia was higher in the pertuzumab-treated group (26%) compared with the placebo-treated group (12%).

Table 3 summarizes the adverse reactions in CLEOPATRA that occurred ≥ 10% of patients in the pertuzumab-treated group.

Table 3: Adverse Reactions (≥ 10%) in Patients Who Received Pertuzumab in Combination with Trastuzumab and Docetaxel in CLEOPATRA

Adverse Reactions	Pertuzumab + trastuzumab + docetaxel n=407 %		Placebo + trastuzumab + docetaxel n=397 %	
	All Grades %	Grades 3 – 4 %	All Grades %	Grades 3 – 4 %
Gastrointestinal disorders				
Diarrhea	67	8	46	5
Nausea	42	1	42	0.5
Vomiting	24	1	24	2
Stomatitis	19	0.5	15	0.3
Constipation	15	0	25	1
Skin and subcutaneous tissue disorders				
Alopecia	61	0	60	0.3
Rash	34	0.7	24	0.8
Nail disorder	23	1	23	0.3
Pruritus	14	0	10	0
Dry skin	11	0	4	0
Blood and lymphatic system disorders				
Neutropenia	53	49	50	46
Anemia	23	2	19	4
Leukopenia	18	12	20	15
Febrile neutropenia*	14	13	8	7
General disorders and administration site conditions				
Fatigue	37	2	37	3
Mucosal inflammation	28	1	20	1
Asthenia	26	2	30	2
Peripheral edema	23	0.5	30	0.8
Pyrexia	19	1	18	0.5
Nervous system disorders				
Neuropathy peripheral	32	3	34	2
Headache	21	1	17	0.5
Dysgeusia	18	0	16	0
Dizziness	13	0.5	12	0
Metabolism and nutrition disorders				
Decreased appetite	29	2	26	2
Musculoskeletal and connective tissue disorders				
Myalgia	23	1	24	0.8
Arthralgia	15	0.2	16	0.8
Infections and infestations				
Upper respiratory tract infection	17	0.7	13	0
Nasopharyngitis	12	0	13	0.3
Respiratory, thoracic, and mediastinal disorders				
Dyspnea	14	1	16	2
Eye disorders				
Lacrimation increased	14	0	14	0

Psychiatric disorders				
Insomnia	13	0	13	0

* In this table this denotes an adverse reaction that has been reported in association with a fatal outcome

Clinically relevant adverse reactions in < 10% of patients in the pertuzumab-treated group in CLEOPATRA included paronychia (7%).

Adverse Reactions Reported in Patients Receiving Pertuzumab and Trastuzumab After Discontinuation of Docetaxel

In CLEOPATRA, adverse reactions that occurred after discontinuation of docetaxel included diarrhea (19%), upper respiratory tract infection (13%), rash (12%), headache (11%), and fatigue (11%).

Neoadjuvant Treatment of Breast Cancer

NeoSphere

The safety of pertuzumab was evaluated in a randomized trial (NeoSphere) in patients with operable, locally advanced, or inflammatory HER2-positive breast cancer (T2-4d) who were scheduled for neoadjuvant therapy [see *Clinical Studies* (14.2)].

In combination with trastuzumab and docetaxel, pertuzumab was given intravenously at an initial dose of 840 mg, followed by 420 mg every 3 weeks for 4 cycles. After surgery, patients in the pertuzumab plus trastuzumab arm received docetaxel every 3 weeks for 4 cycles prior to FEC.

Permanent discontinuation of neoadjuvant pertuzumab due to an adverse reaction occurred in 0.9% of patients.

The most common adverse reactions (> 30%) were alopecia, neutropenia, diarrhea, and nausea. The most common Grade 3 – 4 adverse reactions (> 2%) were neutropenia, febrile neutropenia, leukopenia, and diarrhea.

Table 4 summarizes the adverse reactions in NeoSphere that occurred $\geq$ 10% of patients who received neoadjuvant pertuzumab with trastuzumab and docetaxel followed by FEC.

Table 4: Adverse Reactions(≥ 10%) in Patients who Received Neoadjuvant Pertuzumab in NeoSphere

Adverse Reactions	Trastuzumab + docetaxel n = 107 %		Pertuzumab + trastuzumab + docetaxel n = 107 %	
	All Grades %	Grades 3 – 4 %	All Grades %	Grades 3 – 4 %
Skin and subcutaneous tissue disorders				
Alopecia	66	0	65	0
Rash	21	2	26	0.9
Blood and lymphatic system disorders				
Neutropenia	64	59	50	45
Leukopenia	21	11	9	5
Gastrointestinal disorders				
Nausea	36	0	39	0
Diarrhea	34	4	46	6
Vomiting	12	0	13	0
Stomatitis	7	0	18	0
General disorders and administration site conditions				
Fatigue	27	0	26	0.9
Mucosal inflammation	21	0	26	2
Asthenia	18	0	21	2
Pyrexia	10	0	17	0
Peripheral edema	10	0	3	0
Musculoskeletal and connective tissue disorders				
Myalgia	22	0	22	0
Arthralgia	8	0	10	0
Nervous system disorders				
Peripheral Sensory Neuropathy	12	0.9	8	0.9
Headache	11	0	11	0
Dysgeusia	10	0	15	0
Psychiatric disorders				
Insomnia	11	0	8	0
Metabolism and nutrition disorders				
Decreased appetite	7	0	14	0

Clinically relevant adverse reactions in < 10% of patients receiving neoadjuvant pertuzumab with trastuzumab and docetaxel followed by FEC included anemia, febrile neutropenia, dizziness, upper respiratory tract infection, and increased lacrimation.

Neoadjuvant Treatment of Breast Cancer

TRYPHAENA

The safety of pertuzumab was evaluated in patients with HER2-positive locally advanced, operable, or inflammatory (T2-4d) breast cancer in TRYPHAENA [see *Clinical Studies (14.2)*].

Adverse reactions resulting in permanent discontinuation of any component of neoadjuvant treatment occurred in 7% of patients receiving pertuzumab in combination with trastuzumab and

docetaxel following FEC, and 8% for patients receiving pertuzumab in combination with TCH.

The most common adverse reactions (>2%) resulting in permanent discontinuation of pertuzumab were left ventricular dysfunction, drug hypersensitivity, and neutropenia.

For pertuzumab administered in combination with trastuzumab and docetaxel for 3 cycles following 3 cycles of FEC, the most common adverse reactions (> 30%) were diarrhea, nausea, alopecia, neutropenia, vomiting, and fatigue. The most common Grade 3 – 4 adverse reactions (> 2%) were neutropenia, leukopenia, febrile neutropenia, diarrhea, left ventricular dysfunction, anemia, dyspnea, nausea, and vomiting.

For pertuzumab administered in combination with docetaxel, carboplatin, and trastuzumab (TCH) for 6 cycles, the most common adverse reactions (> 30%) were diarrhea, alopecia, neutropenia, nausea, fatigue, vomiting, anemia, and thrombocytopenia. The most common Grade 3 – 4 adverse reactions (> 2%) were neutropenia, febrile neutropenia, anemia, leukopenia, diarrhea, thrombocytopenia, vomiting, fatigue, ALT increased, hypokalemia, and hypersensitivity.

Table 5 summarizes the adverse reactions in TRYPHAENA that occurred in > 10% of patients who received neoadjuvant pertuzumab with trastuzumab and docetaxel following FEC or who received neoadjuvant pertuzumab in combination with TCH.

Table 5: Adverse Reactions (≥ 10%) in Patients Receiving Neoadjuvant Treatment with Pertuzumab in TRYPHAENA

Adverse Reactions	Pertuzumab + trastuzumab + docetaxel following FEC n = 75 %		Pertuzumab + TCH n = 76 %	
	All Grades %	Grades 3 – 4 %	All Grades %	Grades 3 – 4 %
Gastrointestinal disorders				
Diarrhea	61	5	72	12
Nausea	53	3	45	0
Vomiting	36	3	39	5
Dyspepsia	8	0	22	0
Constipation	23	0	16	0
Stomatitis	17	0	12	0
Skin and subcutaneous tissue disorders				
Alopecia	52	0	55	0
Rash	11	0	21	1
Palmar-Plantar Erythrodysesthesia Syndrome	11	0	8	0
Dry skin	9	0	11	0
Blood and lymphatic system disorders				
Neutropenia	47	43	49	46
Leukopenia	16	12	17	12
Anemia	9	4	38	17
Febrile neutropenia	9	9	17	17
Thrombocytopenia	1	0	30	12

General disorders and administration site conditions				
Fatigue	36	0	42	4
Mucosal inflammation	20	0	17	1
Pyrexia	9	0	16	0
Asthenia	15	1	13	1
Edema peripheral	4	0	9	0
Psychiatric disorders				
Insomnia	13	0	21	0
Nervous system disorders				
Headache	15	0	17	0
Dysgeusia	13	0	21	0
Dizziness	8	1	16	0
Neuropathy peripheral	1	0	11	0
Metabolism and nutrition disorders				
Decreased appetite	11	0	21	0
Respiratory, thoracic, and mediastinal disorders				
Epistaxis	11	0	16	1
Dyspnea	8	3	11	1
Oropharyngeal pain	7	0	12	0
Cough	5	0	12	0
Musculoskeletal and connective tissue disorders				
Myalgia	11	1	11	0
Arthralgia	12	0	7	0
Eye disorders				
Lacrimation increased	5	0	8	0
Investigations				
ALT increased	3	0	11	4
Immune system disorders				
Hypersensitivity	1	0	12	3

Clinically relevant adverse reactions in < 10% of patients who received neoadjuvant pertuzumab with trastuzumab and docetaxel following FEC or who received neoadjuvant pertuzumab in combination with TCH included nail disorder, paronychia, pruritus, upper respiratory tract infection, and nasopharyngitis.

Neoadjuvant Treatment of Breast Cancer

BERENICE

The safety of pertuzumab was evaluated in a two-arm non-randomized study (BERENICE) in patient with HER2-positive locally advanced, inflammatory, or early-stage HER2-positive breast cancer [see *Clinical Studies* (14.2)].

Adverse reactions resulting in permanent discontinuation of any component of neoadjuvant treatment were 14% for patients receiving pertuzumab in combination with trastuzumab and paclitaxel following ddAC and 8% for patients receiving pertuzumab in combination with trastuzumab and docetaxel following FEC. The most common adverse reactions (>1%) resulting in permanent discontinuation of any component of neoadjuvant treatment were peripheral

neuropathy, decreased ejection fraction, diarrhea, neutropenia and infusion-related reaction.

For pertuzumab administered in combination with trastuzumab and paclitaxel for 4 cycles following 4 cycles of ddAC, the most common adverse reactions (> 30%) were nausea, diarrhea, alopecia, fatigue, constipation, peripheral neuropathy and headache. The most common Grade 3 – 4 adverse reactions (> 2%) were neutropenia, febrile neutropenia, decreased neutrophil count, decreased white blood cell count, anemia, diarrhea, peripheral neuropathy, increased ALT, and nausea.

For pertuzumab administered in combination with trastuzumab and docetaxel for 4 cycles following 4 cycles of FEC, the most common adverse reactions (> 30%) were diarrhea, nausea, alopecia, asthenia, constipation, fatigue, mucosal inflammation, vomiting, myalgia, and anemia. The most common Grade 3 – 4 adverse reactions (> 2%) were febrile neutropenia, diarrhea, neutropenia, decreased neutrophil count stomatitis, fatigue, vomiting, mucosal inflammation, neutropenic sepsis and anemia.

Table 6 summarizes the adverse reactions in BERENICE that occurred in $\geq 10\%$ of patients who received neoadjuvant pertuzumab with trastuzumab and paclitaxel following ddAC or who received neoadjuvant pertuzumab with trastuzumab and docetaxel following FEC.

Table 6: Adverse Reactions ($\geq 10\%$) of Patients Receiving Neoadjuvant Pertuzumab in Combination with Trastuzumab and Taxane Chemotherapy Following ddAC or FEC in BERENICE

Adverse Reactions	Pertuzumab + trastuzumab + paclitaxel following ddAC n=199 %		Pertuzumab + trastuzumab + docetaxel following FEC n=198 %	
	All Grades %	Grades 3 – 4 %	All Grades %	Grades 3 – 4 %
Gastrointestinal disorders				
Nausea	71	3	69	2
Diarrhea	67	3	69	10
Constipation	35	0.5	38	0.5
Vomiting	23	1	35	4
Stomatitis	25	0	27	5
Dyspepsia	19	0	16	0
Upper abdominal pain	6	0	13	0
Abdominal pain	5	0	10	0
Gastroesophageal reflux disease	12	0	2	0
Skin and subcutaneous tissue disorders				
Alopecia	62	0	59	0
Rash	14	0	11	0
Dry skin	14	0	10	0
Nail discoloration	15	0	2	0
Palmar-Plantar Erythrodysesthesia Syndrome	6	0	10	0.5
General disorders and administration site conditions				
Fatigue	58	1	38	5
Asthenia	19	2	41	0

Mucosal inflammation	22	1	37	4
Pyrexia	15	0	18	0
Peripheral edema	9	0	12	1
Nervous system disorders				
Peripheral neuropathy	42	3	26	0.5
Headache	30	0.5	14	0.5
Dysgeusia	20	0	19	0.5
Paresthesia	15	0	9	0
Dizziness	12	0	8	0
Blood and lymphatic system disorders				
Anemia	27	3	30	3
Neutropenia	22	12	16	9
Febrile neutropenia	7	7	17	17
Musculoskeletal and connective tissue disorders				
Myalgia	20	0	33	1
Arthralgia	20	0	21	1
Back pain	10	0	9	0
Pain in extremity	10	0	8	0
Bone pain	12	0.5	5	0
Respiratory, thoracic, and mediastinal disorders				
Epistaxis	25	0	19	0
Dyspnea	15	0.5	15	0.5
Cough	20	0.5	9	0
Oropharyngeal pain	10	0	8	0.5
Metabolism and nutrition disorders				
Decreased appetite	20	0	23	0
Psychiatric disorders				
Insomnia	19	0	13	0
Vascular disorders				
Hot flush	19	0	13	0
Injury, poisoning and procedural complications				
Infusion related reaction	16	1	13	1
Eye disorders				
Increased lacrimation	9	0	18	0
Investigations				
Decreased white blood cell count	11	4	3	2
Infections and infestations				
Urinary tract infection	11	1	2	0

Clinically relevant adverse reactions in < 10% of patients who received pertuzumab in combination with trastuzumab and paclitaxel following ddAC or patients receiving pertuzumab in combination with trastuzumab and docetaxel following FEC included pruritus, nail disorder, paronychia, upper respiratory tract infection, and nasopharyngitis.

Adjuvant Treatment of Breast Cancer

APHINITY

The safety of pertuzumab was evaluated in a multicenter, randomized, double-blind, placebo-controlled study (APHINITY) conducted in patients with HER2-positive early breast cancer who had their primary tumor excised prior to randomization [see *Clinical Studies* (14.3)].

Patients were randomized to receive either pertuzumab in combination with trastuzumab and chemotherapy or placebo in combination with trastuzumab and chemotherapy. Investigators selected one of three anthracycline-based or non-anthracycline-based chemotherapy regimens for patients. Pertuzumab and trastuzumab were administered intravenously every 3 weeks starting on Day 1 of the first taxane-containing cycle, for a total of 52 weeks (up to 18 cycles) or until recurrence, withdrawal of consent, or unmanageable toxicity.

Serious adverse reactions (hospitalization) due to diarrhea in the pertuzumab-treated group was 2.4%. The incidence of diarrhea was higher when chemotherapy was administered with pertuzumab (61%) and was higher when administered with non-anthracycline based therapy (85%) than with anthracycline based therapy (67%). The median duration of diarrhea was 8 days. The median duration of Grade ≥ 3 diarrhea was 20 days. The incidence of diarrhea during the period pertuzumab and trastuzumab were administered without chemotherapy was 18% in the pertuzumab-treated group.

Adverse reactions resulting in permanent discontinuation of any study therapy were 13% for patients in the pertuzumab-treated group. Adverse reactions resulting in permanent discontinuation of pertuzumab was 7%. The most common adverse reactions ($> 0.5\%$) resulting in permanent discontinuation of any study treatment were ejection fraction decreased, neuropathy peripheral, diarrhea, and cardiac failure.

When pertuzumab was administered in combination with trastuzumab and chemotherapy, the most common adverse reactions ($> 30\%$) were diarrhea, nausea, alopecia, fatigue, peripheral neuropathy, and vomiting. The most common Grade 3 – 4 adverse reactions ($> 2\%$) were neutropenia, febrile neutropenia, diarrhea, neutrophil count decreased, anemia, white blood cell count decreased, leukopenia, fatigue, nausea, and stomatitis.

Table 7 summarizes the adverse reactions that occurred in $\geq 10\%$ of patients who received adjuvant pertuzumab in combination with trastuzumab and chemotherapy followed by pertuzumab and trastuzumab for a total of 52 weeks (up to 18 cycles) or until recurrence, withdrawal of consent, or unmanageable toxicity.

Table 7: Adverse Reactions ($\geq 10\%$) of Patients Receiving Adjuvant Pertuzumab in Combination with Trastuzumab and Chemotherapy Followed by Pertuzumab and Trastuzumab in APHINITY

Adverse Reactions	Pertuzumab + trastuzumab + chemotherapy n=2364 %		Placebo + trastuzumab + chemotherapy n=2405 %	
	All Grades %	Grades 3 – 4 %	All Grades %	Grades 3 – 4 %
Gastrointestinal disorders				
Diarrhea	71	10	45	4
Nausea	69	2	65	2

Vomiting	32	2	30	2
Constipation	29	0.5	32	0.3
Stomatitis	28	2	24	1
Dyspepsia	14	0	14	0
Abdominal pain	12	0.5	11	0.6
Abdominal pain upper	10	0.3	9	0.2
Skin and subcutaneous tissue disorders				
Alopecia	67	<0.1	67	<0.1
Rash	26	0.4	20	0.2
Pruritus	14	0.1	9	<0.1
Dry skin	13	0.1	11	<0.1
Nail disorder	12	0.2	12	0.1
General disorders and administration site conditions				
Fatigue	49	4	44	3
Mucosal inflammation	23	2	19	0.7
Asthenia	21	1	21	2
Pyrexia	20	0.6	20	0.7
Edema peripheral	17	0	20	0.2
Musculoskeletal and connective tissue disorders				
Arthralgia	29	0.9	33	1
Myalgia	26	0.9	30	1
Pain in extremity	10	0.2	10	0.2
Blood and lymphatic system disorders				
Anemia	28	7	23	5
Neutropenia	25	16	23	16
Febrile neutropenia*	12	12	11	11
Nervous system disorders				
Dysgeusia	26	0.1	22	<0.1
Neuropathy peripheral	33	1	32	1
Headache	22	0.3	23	0.4
Paresthesia	12	0.5	10	0.2
Dizziness	11	0	11	0.2
Metabolism and nutrition disorders				
Decreased appetite	24	0.8	20	0.4
Vascular disorders				
Hot flush	20	0.2	21	0.4
Respiratory, thoracic, and mediastinal disorders				
Epistaxis	18	<0.1	14	0
Cough	16	<0.1	15	<0.1

Dyspnea	12	0.4	12	0.5
Psychiatric disorders				
Insomnia	17	0.3	17	<0.1
Investigations				
Neutrophil count decreased	14	10	14	10
Eye disorders				
Lacrimation increased	13	0	13	<0.1
Infections and infestations				
Nasopharyngitis	13	<0.1	12	0.1
Injury, poisoning and procedural complications				
Radiation skin injury	13	0.3	11	0.3

* In this table this denotes an adverse reaction that has been reported in association with a fatal outcome

Clinically relevant adverse reactions in < 10% of patients who received pertuzumab in combination with trastuzumab and anthracycline-based or non-anthracycline-based chemotherapy regimens included leukopenia, upper respiratory tract infection, and paronychia

Adverse Reactions in Patients Receiving Pertuzumab and Trastuzumab After Discontinuation of Chemotherapy

In APHINITY, adverse reactions that occurred after discontinuation of chemotherapy in > 10% included diarrhea (18%), arthralgia (15%), radiation skin injury (12%), and hot flush (12%).

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Based on its mechanism of action and findings in animal studies, pertuzumab products can cause fetal harm when administered to a pregnant woman. There are no available data on the use of pertuzumab products in pregnant women. However, in post-marketing reports, use of another HER2/neu receptor antagonist (trastuzumab) during pregnancy resulted in cases of oligohydramnios and oligohydramnios sequence manifesting as pulmonary hypoplasia, skeletal abnormalities, and neonatal death. In an animal reproduction study, administration of pertuzumab to pregnant cynomolgus monkeys during the period of organogenesis resulted in oligohydramnios, delayed fetal kidney development, and embryo-fetal deaths at clinically relevant exposures that were 2.5 to 20-fold greater than exposures in humans receiving the recommended dose, based on C_{max} [see Data]. Apprise the patient of the potential risks to a fetus. There are clinical considerations if POHERDY is used in combination with trastuzumab during pregnancy or within 7 months prior to conception [see Clinical Considerations].

The estimated background risk of major birth defects and miscarriage for the indicated population is unknown. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2-4% and 15-20%, respectively.

Clinical Considerations

Fetal/Neonatal Adverse Reactions

Monitor women who received POHERDY in combination with trastuzumab during pregnancy or within 7 months prior to conception for oligohydramnios. If oligohydramnios occurs, perform fetal testing that is appropriate for gestational age and consistent with community standards of care.

Data

Animal Data

Pregnant cynomolgus monkeys were treated on Gestational Day (GD)19 with loading doses of 30 to 150 mg/kg pertuzumab, followed by bi-weekly doses of 10 to 100 mg/kg. These dose levels resulted in clinically relevant exposures of 2.5 to 20-fold greater than exposures in humans receiving the recommended dose, based on C_{max} . Intravenous administration of pertuzumab from GD19 through GD50 (period of organogenesis) was embryotoxic, with dose-dependent increases in embryo-fetal death between GD25 to GD70. The incidences of embryo-fetal loss were 33, 50, and 85% for dams treated with bi-weekly pertuzumab doses of 10, 30, and 100 mg/kg, respectively (2.5 to 20-fold greater than the recommended human dose, based on C_{max}). At Caesarean section on GD100, oligohydramnios, decreased relative lung and kidney weights, and microscopic evidence of renal hypoplasia consistent with delayed renal development were identified in all pertuzumab dose groups. Pertuzumab exposure was reported in offspring from all treated groups, at levels of 29% to 40% of maternal serum levels at GD100.

8.2 Lactation

Risk Summary

There is no information regarding the presence of pertuzumab products in human milk, the effects on the breastfed infant or the effects on milk production. Published data suggest that human IgG is present in human milk but does not enter the neonatal and infant circulation in substantial amounts. Consider the developmental and health benefits of breast feeding along with the mother's clinical need for POHERDY treatment and any potential adverse effects on the breastfed child from POHERDY or from the underlying maternal condition. This consideration should also take into account the elimination half-life of pertuzumab and the trastuzumab wash out period of 7 months.

8.3 Females and Males of Reproductive Potential

Pregnancy Testing

Verify the pregnancy status of females of reproductive potential prior to the initiation of POHERDY.

Contraception

Females

Based on the mechanism of action and animal data, pertuzumab products can cause embryo-fetal harm when administered during pregnancy. Advise females of reproductive potential to use effective contraception during treatment and for 7 months following the last dose of POHERDY in combination with trastuzumab [*see Use in Specific Populations (8.1)*].

8.4 Pediatric Use

The safety and effectiveness of POHERDY have not been established in pediatric patients.

8.5 Geriatric Use

In CLEOPATRA, NeoSphere, TRYPHAENA, BERENICE, and APHINITY, 464 patients who received pertuzumab were ≥ 65 years of age and 47 were ≥ 75 years of age.

The incidence of adverse reactions was increased in patients aged ≥ 65 years of age, compared to patients aged < 65 years of age for decreased appetite, anemia, decreased weight, asthenia, dysgeusia, peripheral neuropathy, and hypomagnesemia.

No overall differences in efficacy of pertuzumab were observed in patients aged ≥ 65 and < 65 years of age. Clinical studies did not include sufficient numbers of patients aged ≥ 75 years to determine if these patients respond differently than younger patients.

Based on a population pharmacokinetic analysis, no significant difference was observed in the pharmacokinetics of pertuzumab between patients < 65 years ($n=306$) and patients ≥ 65 years ($n=175$).

8.6 Renal Impairment

Dose adjustments of POHERDY are not needed in patients with mild (creatinine clearance [CLCr] 60 to 90 mL/min) or moderate (CLCr 30 to 60 mL/min) renal impairment. No dose adjustment can be recommended for patients with severe renal impairment (CLCr less than 30 mL/min) because of the limited pharmacokinetic data available [*see Clinical Pharmacology (12.3)*].

11 DESCRIPTION

Pertuzumab-dpzb is a recombinant humanized monoclonal antibody and HER2/neu receptor antagonist that targets the extracellular dimerization domain (Subdomain II) of the human epidermal growth factor receptor 2 protein (HER2). Pertuzumab-dpzb is produced by recombinant DNA technology in a mammalian cell (Chinese Hamster Ovary) culture. Pertuzumab-dpzb has an approximate molecular weight of 148 kDa.

POHERDY (pertuzumab-dpzb) injection is a sterile, preservative-free, clear to slightly opalescent, colorless to pale yellow solution for intravenous infusion. Each 14 mL single-dose vial contains 420 mg of pertuzumab-dpzb, and histidine (21.14 mg), L-histidine hydrochloride monohydrate (30.1 mg), polysorbate 20 (2.8 mg), sorbitol (420 mg) and Water for Injection, with a pH of 6.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Pertuzumab products target the extracellular dimerization domain (Subdomain II) of the human epidermal growth factor receptor 2 protein (HER2) and, thereby, blocks ligand-dependent heterodimerization of HER2 with other HER family members, including EGFR, HER3, and HER4. As a result, pertuzumab products inhibit ligand-initiated intracellular signaling through two major signal pathways, mitogen-activated protein (MAP) kinase, and phosphoinositide 3-

kinase (PI3K). Inhibition of these signaling pathways can result in cell growth arrest and apoptosis, respectively. In addition, pertuzumab products mediate antibody-dependent cell-mediated cytotoxicity (ADCC).

While pertuzumab products alone inhibited the proliferation of human tumor cells, the combination of pertuzumab products and trastuzumab augmented anti-tumor activity in HER2-overexpressing xenograft models.

12.2 Pharmacodynamics

The exposure-response relationship and time course of pharmacodynamic response for the safety and effectiveness of pertuzumab products have not been fully characterized.

Cardiac Electrophysiology

The effect of pertuzumab with an initial dose of 840 mg followed by a maintenance dose of 420 mg every three weeks on QTc interval was evaluated in a subgroup of 20 patients with HER2-positive breast cancer in CLEOPATRA. At the recommended dose of pertuzumab, a mean increase in the QTc interval >20 ms was not observed. A small increase in the mean QTc interval (i.e., less than 10 ms) cannot be excluded because of the limitations of the trial design.

12.3 Pharmacokinetics

Based on a population PK analysis that included 481 patients, pertuzumab demonstrated linear pharmacokinetics at a dose range of 2 – 25 mg/kg.

With an initial dose of 840 mg followed by a maintenance dose of 420 mg every three weeks thereafter, the steady-state concentration of pertuzumab was reached after the first maintenance dose.

Elimination

The median clearance (CL) of pertuzumab was 0.24 L/day and the median half-life was 18 days.

Specific Populations

No clinically significant differences in the pharmacokinetics of pertuzumab were observed based on age, sex, ethnicity (Japanese vs. non-Japanese), or disease status (neoadjuvant or adjuvant vs. metastatic setting). No dose adjustments based on body weight or baseline albumin level are needed, as the exposure changes are not considered clinically relevant.

Pertuzumab exposure in patients with mild (CLcr 60 to 90 mL/min, n=200) and moderate renal impairment (CLcr 30 to 60 mL/min, n=71) were similar to those in patients with normal renal function (CLcr greater than 90 mL/min, n=200). The pharmacokinetics of pertuzumab in patients with moderate to severe hepatic impairment or severe renal impairment is unknown.

12.6 Immunogenicity

The observed incidence of anti-drug antibodies is highly dependent on the sensitivity and specificity of the assay. Differences in assay methods preclude meaningful comparisons of the incidence of anti-drug antibodies in the studies described below with the incidence of anti-drug antibodies in other studies, including those of pertuzumab or of other pertuzumab products.

Patients in CLEOPATRA were tested at multiple time-points for anti-pertuzumab antibodies. 3%

(13/389) of patients in the pertuzumab-treated group and 7% (25/372) of patients in the placebo-treated group tested positive for anti-pertuzumab antibodies. Of these 38 patients, none experienced anaphylactic/hypersensitivity reactions that were clearly related to anti-pertuzumab antibodies. The presence of pertuzumab in patient serum at the levels expected at the time of anti-drug antibodies sampling can interfere with the ability of this assay to detect anti-pertuzumab antibodies. In addition, the assay may be detecting antibodies to trastuzumab. As a result, data may not accurately reflect the true incidence of anti-pertuzumab antibody development.

In the neoadjuvant period of BERENICE, 0.3% (1/383) of patients treated with pertuzumab tested positive for anti-pertuzumab antibodies. This patient did not experience any anaphylactic/hypersensitivity reactions.

Because of limited immunogenicity data, the clinical impact of anti-pertuzumab antibodies is unknown. There was no identified clinically significant effect of anti-pertuzumab antibodies on the safety of pertuzumab.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Long-term studies in animals have not been performed to evaluate the carcinogenic potential of pertuzumab products.

Studies have not been performed to evaluate the mutagenic potential of pertuzumab products.

No specific fertility studies in animals have been performed to evaluate the effect of pertuzumab products. No adverse effects on male and female reproductive organs were observed in repeat-dose toxicity studies of up to six months duration in cynomolgus monkeys.

14 CLINICAL STUDIES

14.1 Metastatic Breast Cancer

CLEOPATRA (NCT00567190) was a multicenter, double-blind, placebo-controlled trial of 808 patients with HER2-positive metastatic breast cancer. HER2 overexpression was defined as a score of 3+ IHC or FISH amplification ratio of 2.0 or greater as determined by a central laboratory. Patients were randomly allocated 1:1 to receive placebo plus trastuzumab and docetaxel or pertuzumab plus trastuzumab and docetaxel. Randomization was stratified by prior treatment (prior or no prior adjuvant/neoadjuvant anti-HER2 therapy or chemotherapy) and geographic region (Europe, North America, South America, and Asia). Patients with prior adjuvant or neoadjuvant therapy were required to have a disease-free interval of greater than 12 months before trial enrollment.

Pertuzumab was given intravenously at an initial dose of 840 mg, followed by 420 mg every 3 weeks thereafter. Trastuzumab was given intravenously at an initial dose of 8 mg/kg, followed by 6 mg/kg every 3 weeks thereafter. Patients were treated with pertuzumab and trastuzumab until progression of disease, withdrawal of consent, or unacceptable toxicity. Docetaxel was given as an initial dose of 75 mg/m² by intravenous infusion every 3 weeks for at least 6 cycles. The docetaxel dose could be escalated to 100 mg/m² at the investigator's discretion if the initial dose was well tolerated. At the time of the primary analysis, the mean number of cycles of study

treatment administered was 16.2 in the placebo-treated group and 19.9 in the pertuzumab-treated group.

The major efficacy outcome measure of CLEOPATRA was progression-free survival (PFS) as assessed by an independent review facility (IRF). PFS was defined as the time from the date of randomization to the date of disease progression or death (from any cause) if the death occurred within 18 weeks of the last tumor assessment. Additional endpoints included overall survival (OS), PFS (investigator-assessed), objective response rate (ORR) and duration of response.

Patient demographic and baseline characteristics were balanced between the treatment arms. The median age was 54 (range 22 to 89 years), 59% were White, 32% were Asian, and 4% were Black. All were female with the exception of 2 patients (0.2%). Seventeen percent (17%) of patients were enrolled in North America, 14% in South America, 38% in Europe, and 31% in Asia. Tumor prognostic characteristics, including hormone receptor status (positive 48%, negative 50%), presence of visceral disease (78%) and non-visceral disease only (22%) were similar in the study arms. Approximately half of the patients received prior adjuvant or neoadjuvant anti-HER2 therapy or chemotherapy (pertuzumab 46%, placebo 47%). Among patients with hormone receptor positive tumors, 45% received prior adjuvant hormonal therapy and 11% received hormonal therapy for metastatic disease. Eleven percent of patients received prior adjuvant or neoadjuvant trastuzumab.

CLEOPATRA demonstrated a statistically significant improvement in IRF-assessed PFS in the pertuzumab-treated group compared with the placebo-treated group. The results for investigator-assessed PFS were comparable to those observed for IRF-assessed PFS. A statistically significant OS improvement was demonstrated for the pertuzumab-treated group compared with the placebo-treated group) with the final OS analysis. OS results in patient subgroups were consistent with those observed for IRF-assessed PFS with the exception of the subgroup of patients with disease limited to non-visceral metastasis [HR=1.11 (95% CI: 0.66, 1.85)].

Table 8: Efficacy Results from CLEOPATRA

Endpoint	Pertuzumab + trastuzumab + docetaxel n=402	Placebo + trastuzumab + docetaxel n=406
Progression-Free Survival (independent review)		
Number of events (%)	191 (47.5%)	242 (59.6%)
Median (months)	18.5	12.4
Hazard Ratio (95% CI)	0.62 (0.51, 0.75)	
p-value	< 0.0001	
Overall Survival* (final)		
Deaths (%)	168 (41.8%)	221 (54.4%)
Median (months)	56.5	40.8
Hazard Ratio (95% CI)	0.68 (0.56, 0.84)	
p-value	0.0002	
Objective Response Rate (independent review)		
	n = 343	n = 336
Objective response (CR + PR)	275 (80.2%)	233 (69.3%)

Complete response (CR) (%)	19 (5.5%)	14 (4.2%)
Partial Response (PR) (%)	256 (74.6%)	219 (65.2%)
Difference in ORR (95% CI)	10.8% (4.2%, 17.5%)	
p-value	0.0011	
Duration of Response		
Median (months)	20.2	12.5

* Final analysis of overall survival performed when 389 patients had died.

CI=Confidence Interval

Consistent results were observed across several patient subgroups including age (< 65 or ≥ 65 years), race, geographic region, prior adjuvant/neoadjuvant anti-HER2 therapy or chemotherapy (yes or no), and prior adjuvant/neoadjuvant trastuzumab (yes or no). In the subgroup of patients with hormone receptor-negative disease (n=408), the hazard ratio was 0.55 (95% CI: 0.42, 0.72). In the subgroup of patients with hormone receptor-positive disease (n=388), the hazard ratio was 0.72 (95% CI: 0.55, 0.95). In the subgroup of patients with disease limited to non-visceral metastasis (n=178), the hazard ratio was 0.96 (95% CI: 0.61, 1.52).

Figure 1 Kaplan-Meier Curve of Progression-Free Survival for CLEOPATRA (IRF-Assessed)

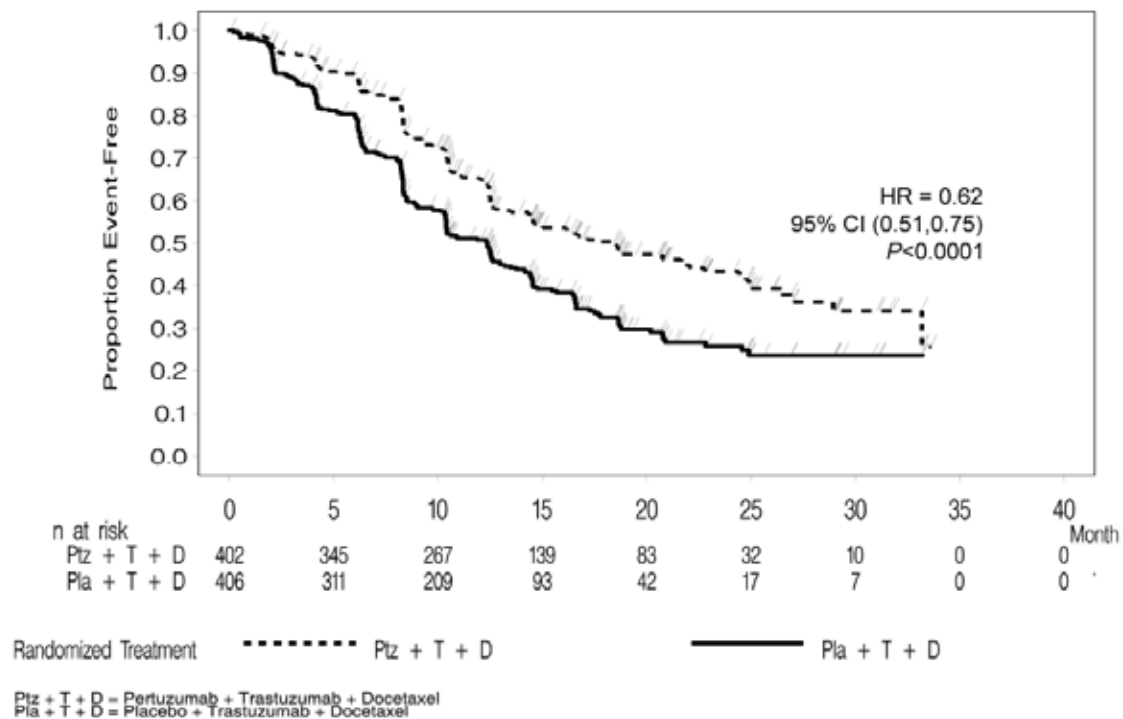
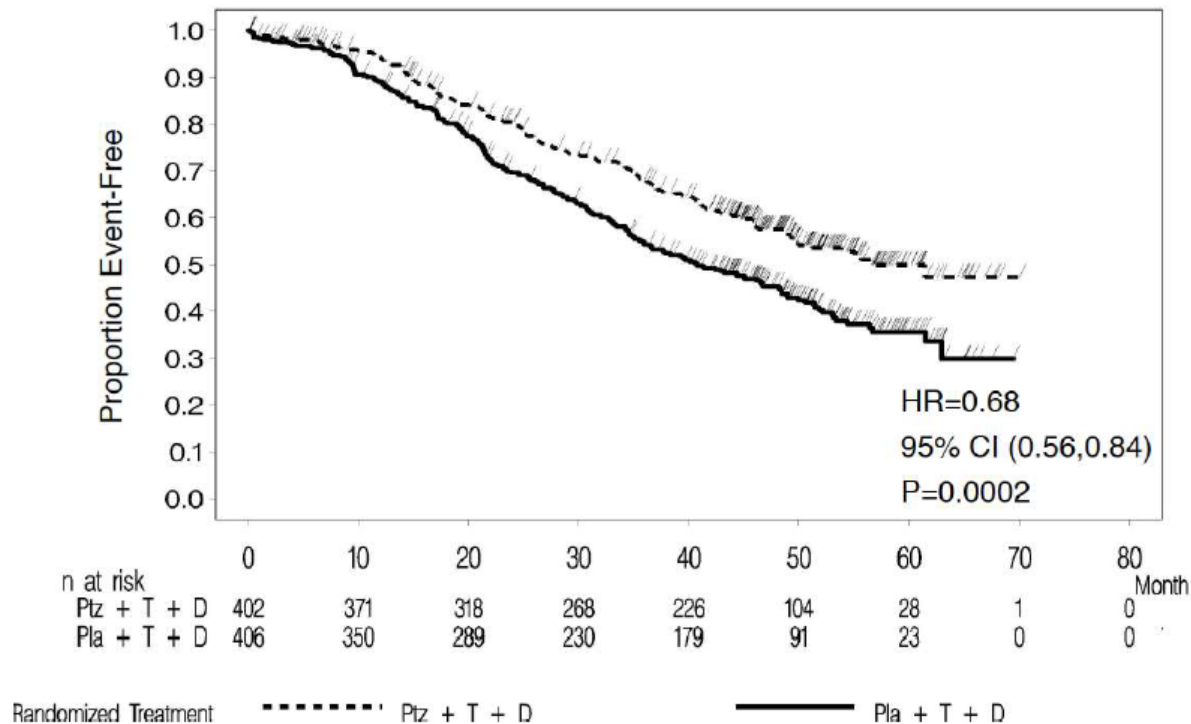


Figure 2 Kaplan-Meier Curve of Overall Survival for CLEOPATRA (Final Analysis)



14.2 Neoadjuvant Treatment of Breast Cancer

NeoSphere

NeoSphere (NCT00545688) was a multicenter, randomized trial conducted in 417 patients with operable, locally advanced, or inflammatory HER2-positive breast cancer (T2-4d) who were scheduled for neoadjuvant therapy. HER2 overexpression was defined as a score of 3+ IHC or FISH amplification ratio of 2.0 or greater as determined by a central laboratory. Patients were randomly allocated to receive 1 of 4 neoadjuvant regimens prior to surgery as follows: trastuzumab plus docetaxel, pertuzumab plus trastuzumab and docetaxel, pertuzumab plus trastuzumab, or pertuzumab plus docetaxel. Randomization was stratified by breast cancer type (operable, locally advanced, or inflammatory) and estrogen receptor (ER) or progesterone receptor (PgR) positivity.

Pertuzumab was given intravenously at an initial dose of 840 mg, followed by 420 mg every 3 weeks for 4 cycles. Trastuzumab was given intravenously at an initial dose of 8 mg/kg, followed by 6 mg/kg every 3 weeks for 4 cycles. Docetaxel was given as an initial dose of 75 mg/m² by intravenous infusion every 3 weeks for 4 cycles. The docetaxel dose could be escalated to 100 mg/m² at the investigator's discretion if the initial dose was well tolerated. Following surgery all patients received 3 cycles of 5-fluorouracil (600 mg/m²), epirubicin (90 mg/m²), and cyclophosphamide (600 mg/m²) (FEC) given intravenously every 3 weeks and trastuzumab administered intravenously every 3 weeks to complete 1 year of therapy. After surgery, patients in the pertuzumab plus trastuzumab arm received docetaxel every 3 weeks for 4 cycles prior to FEC.

The major efficacy outcome measure was pathological complete response (pCR) rate in the breast (ypT0/is) defined as the absence of invasive cancer in the breast and lymph nodes (ypT0/is ypN0) for the efficacy analysis.

Demographics were balanced (median age was 49 – 50 years old, the majority were White (71%) and all were female. Overall, 7% of patients had inflammatory cancer, 32% had locally advanced cancer, and 61% had operable cancer. Approximately half the patients in each treatment group had hormone receptor-positive disease (defined as ER-positive and/or PgR-positive).

Statistically significant improvements in pCR rates were observed in patients receiving pertuzumab plus trastuzumab and docetaxel compared to patients receiving trastuzumab plus docetaxel. The pCR rates and magnitude of improvement with pertuzumab were lower in the subgroup of patients with hormone receptor-positive tumors compared to patients with hormone receptor-negative tumors. The efficacy results are summarized in Table 9.

Table 9: Efficacy Results from NeoSphere

Endpoint	H+T N=107	Ptz+H+T N=107	Ptz+H N=107	Ptz+T N=96
pCR ¹ , n (%) [95% CI] ²	23 (21.5%) [14.1, 30.5]	42 (39.3%) [30.0, 49.2]	12 (11.2%) [5.9, 18.8]	17 (17.7%) [10.7, 26.8]
p-value (with Simes correction for CMH test) ³		0.0063 (vs. H+T)	0.0223 (vs. H+T)	0.0018 (vs. Ptz+H+T)
Hormone receptor-positive subgroup	N=50	N=50	N=51 ⁴	N=46
pCR ¹ , n (%) [95% CI] ²	6 (12.0%) [4.5, 24.3]	11 (22.0%) [11.5, 36.0]	1 (2.0%) [0.1, 10.5]	4 (8.7%) [2.4, 20.8]
Hormone receptor-negative subgroup	N=57	N=57	N=55 ⁴	N=50
pCR ¹ , n (%) [95% CI] ²	17 (29.8%) [18.4, 43.4]	31 (54.4%) [40.7, 67.6]	11 (20.0%) [10.4, 33.0]	13 (26.0%) [14.6, 40.3]

T=docetaxel, Ptz=pertuzumab, H=trastuzumab

CI=Confidence Interval

¹ ypT0/is ypN0 (absence of invasive cancer in the breast and lymph nodes) based on intention-to-treat population

² 95% CI for one sample binomial using Pearson-Clopper method.

³ p-value from Cochran-Mantel-Haenszel (CMH) test, with Simes multiplicity adjustment

⁴ One patient had unknown hormone receptor status. The patient did not achieve a pCR.

TRYPHAENA

TRYPHAENA (NCT00976989) was a three-arm, randomized (1:1:1) study in the neoadjuvant setting conducted in 225 patients with HER2-positive locally advanced, operable, or inflammatory (T2-4d) breast cancer designed primarily to assess cardiac safety. HER2 overexpression was defined as a score of 3+ IHC or FISH amplification ratio of 2.0 or greater as determined by a central laboratory.

Patients were randomly allocated to receive 1 of 3 neoadjuvant regimens prior to surgery as follows: 3 cycles of FEC followed by 3 cycles of docetaxel all in combination with pertuzumab and trastuzumab, 3 cycles of FEC alone followed by 3 cycles of docetaxel and trastuzumab in combination with pertuzumab, or 6 cycles of docetaxel, carboplatin, and trastuzumab (TCH) in combination with pertuzumab. Randomization was stratified by breast cancer type (operable, locally advanced, or inflammatory) and ER and/or PgR positivity.

Pertuzumab was given by intravenous infusion at an initial dose of 840 mg, followed by 420 mg every 3 weeks. Trastuzumab was given by intravenous infusion at an initial dose of 8 mg/kg, followed by 6 mg/kg every 3 weeks. 5-Fluorouracil (500 mg/m²), epirubicin (100 mg/m²), and cyclophosphamide (600 mg/m²) [FEC] were given intravenously every 3 weeks for 3 cycles. In the pertuzumab plus trastuzumab, docetaxel, and FEC arms, docetaxel was given as an initial dose of 75 mg/m² by intravenous infusion every 3 weeks for 3 cycles with the option to escalate to 100 mg/m² at the investigator's discretion if the initial dose was well tolerated. However, in

the pertuzumab plus TCH arm, docetaxel was given intravenously at 75 mg/m² (no escalation was permitted) and carboplatin (AUC 6) was given intravenously every 3 weeks for 6 cycles. Following surgery all patients received trastuzumab to complete 1 year of therapy, which was administered intravenously every 3 weeks.

Demographics were balanced (median age was 49-50 years old, the majority were White [76%]) and all were female. Overall 6% of patients had inflammatory cancer, 25% had locally advanced cancer and 69% had operable cancer, with approximately half the patients in each treatment group having ER-positive and/or PgR-positive disease.

The pCR (ypT0/is ypN0) rates were 56.2% (95% CI: 44.1%, 67.8%), 54.7% (95% CI: 42.7%, 66.2%), and 63.6% (95% CI: 51.9%, 74.3%) for patients treated with pertuzumab plus trastuzumab and FEC followed by pertuzumab plus trastuzumab and docetaxel, pertuzumab plus trastuzumab and docetaxel following FEC, or pertuzumab plus TCH, respectively. The pCR rates were lower in the subgroups of patients with hormone receptor-positive tumors: 41.0% (95% CI: 25.6%, 57.9%), 45.7% (95% CI: 28.8%, 63.4%), and 47.5% (95% CI: 31.5%, 63.9%) than with hormone receptor-negative tumors: 73.5% (95% CI: 55.6%, 87.1%), 62.5% (95% CI: 45.8%, 77.3%), and 81.1% (95% CI: 64.8%, 92.0%), respectively.

BERENICE

A two-arm non-randomized study (BERENICE, NCT02132949) was conducted in 401 patients with HER2-positive locally advanced, inflammatory, or early-stage HER2-positive breast cancer. HER2 overexpression was defined as a score of 3+ IHC or ISH amplification ratio of 2.0 or greater as determined by a central laboratory.

Patients received 1 of 2 neoadjuvant regimens prior to surgery as follows: 4 cycles of dose dense doxorubicin and cyclophosphamide (ddAC) followed by 4 cycles of pertuzumab in combination with trastuzumab and weekly paclitaxel for 12 weeks or 4 cycles of 5-fluorouracil, epirubicin and cyclophosphamide (FEC) followed by 4 cycles of pertuzumab in combination with trastuzumab and docetaxel. The choice of neoadjuvant treatment regimen was made by the Investigator on a site-specific basis. Dosing for the regimens was as follows:

- Pertuzumab was given by intravenous infusion at an initial dose of 840 mg, followed by 420 mg every 3 weeks. Trastuzumab was given by intravenous infusion at an initial dose of 8 mg/kg, followed by 6 mg/kg every 3 weeks.
- In the ddAC cohort, (doxorubicin 60 mg/m² and cyclophosphamide 600 mg/m²) were given intravenously every 2 weeks (ddAC) for 4 cycles with G-CSF (granulocyte colony stimulating factor) support at investigator discretion, followed by paclitaxel 80 mg/m² given intravenously weekly for 12 weeks, with pertuzumab and trastuzumab every 3 weeks from the start of paclitaxel for 4 cycles.
- In the FEC cohort, 5-Fluorouracil (5-FU) (500 mg/m²), epirubicin (100 mg/m²), and cyclophosphamide (600 mg/m²) were given intravenously every 3 weeks for 4 cycles, followed by docetaxel given as an initial dose of 75 mg/m² by intravenous infusion every 3 weeks for 4 cycles with pertuzumab and trastuzumab, and with the option to escalate to 100 mg/m² at the investigator's discretion if the initial dose was well tolerated.

Following surgery, all patients received pertuzumab and trastuzumab administered intravenously every 3 weeks to complete 1 year of therapy.

The median age was 49 years old (range 21-78), 12% of patients were 65 or older, 83% were White, and all but one patient was female (99.8%). Overall 3% of patients had inflammatory cancer, 23% had locally advanced cancer (Stage 3A or greater), 5% were not classified per TNM staging, with approximately two thirds of the patients in each treatment group having ER-positive and/or PgR-positive disease. All patients had an ECOG performance status of 0 or 1.

The pCR (ypT0/is ypN0) rates were 61.8% (95% CI: 54.7, 68.6) and 60.7% (95% CI: 53.6, 67.5) for patients treated with ddAC followed by pertuzumab plus trastuzumab and paclitaxel, or FEC followed by pertuzumab plus trastuzumab and docetaxel, respectively. The pCR rates were lower in the subgroups of patients with hormone receptor-positive tumors: 51.6% (95% CI: 42.6, 60.5%) and 57.3% (95% CI: 48.1, 66.1%) than with hormone receptor-negative tumors: 81.5% (95% CI: 70.0, 90.1%) and 68.0% (95% CI: 56.2, 78.3%), respectively.

14.3 Adjuvant Treatment of Breast Cancer

APHINITY (NCT01358877) was a multicenter, randomized, double-blind, placebo-controlled study conducted in 4804 patients with HER2-positive early breast cancer who had their primary tumor excised prior to randomization. Patients were then randomized to receive pertuzumab or placebo, in combination with adjuvant trastuzumab and chemotherapy. Randomization was stratified by the following factors: region, nodal status, protocol version, central hormone receptor status, and adjuvant chemotherapy regimen.

Investigators selected one of the following anthracycline-based or non-anthracycline-based chemotherapy regimens for individual patients:

- 3 or 4 cycles of FEC (5-FU 500-600 mg/m², epirubicin 90-120 mg/m², cyclophosphamide 500-600 mg/m²) or FAC (5-FU 500-600 mg/m², doxorubicin 50 mg/m², cyclophosphamide 500-600 mg/m²), followed by 3 or 4 cycles of docetaxel (75 mg/m² which could be escalated to 100 mg/m² every 3 weeks) or 12 cycles of weekly paclitaxel (80 mg/m²).
- 4 cycles of AC (doxorubicin 60 mg/m² and cyclophosphamide 500-600 mg/m²) or EC (epirubicin 90-120 mg/m² and cyclophosphamide 500-600 mg/m²) either every 3 weeks or every 2 weeks with GCSF support, followed by docetaxel (100 mg/m² for 3 cycles or 75 mg/m² for first cycle and 100 mg/m² for subsequent three cycles, or 75 mg/m² for four cycles) or 12 cycles of weekly paclitaxel (80 mg/m²).
- 6 cycles of docetaxel (75 mg/m²) in combination with carboplatin (AUC 6)

Pertuzumab and trastuzumab were administered intravenously every 3 weeks starting on Day 1 of the first taxane-containing cycle, for a total of 52 weeks (up to 18 cycles) or until recurrence, withdrawal of consent, or unmanageable toxicity.

After completion of chemotherapy, patients received radiotherapy and/or hormone therapy as per investigator's discretion.

The major efficacy outcome of the study was invasive disease-free survival (IDFS), defined as the time from randomization to first occurrence of ipsilateral local or regional invasive breast cancer recurrence, distant recurrence, contralateral invasive breast cancer, or death from any cause. Additional efficacy endpoints were IDFS including second primary non-breast cancer, disease-free survival (DFS), and overall survival (OS).

Demographics were balanced between the two treatment arms. The median age was 51 years (range 18-86), 13% of patients were 65 or older, and over 99% of patients were female. Sixty-three percent of patients had node-positive disease, 64% had hormone receptor-positive disease, and 71% were White. All patients had an ECOG performance status of 0 or 1. Seventy-eight percent received an anthracycline containing regimen.

Pertuzumab-treated patients and placebo-treated patients both received a median number of 18 cycles of anti-HER2 therapy. After a median follow-up of 45.4 months, a statistically significant improvement in IDFS was demonstrated in patients randomized to receive pertuzumab compared with patients randomized to receive placebo. The efficacy results from APHINITY are summarized in Tables 10 and 11 and in Figure 3.

Table 10 Efficacy Results from APHINITY

	Pertuzumab + trastuzumab + chemotherapy N=2400	Placebo + trastuzumab + chemotherapy N=2404
Invasive Disease Free Survival (IDFS) Number (%) of patients with event HR [95% CI] ¹ p-value (Log-Rank test, stratified ¹) 3 year event-free rate ² , % [95% CI]	171 (7.1%) 0.82 [0.67, 1.00] 0.047 94.1 [93.1, 95.0]	210 (8.7%) 0.82 [0.67, 1.00] 0.047 93.2 [92.2, 94.3]
IDFS including second primary non-breast cancer Number (%) of patients with event HR [95% CI] ¹ 3 year event-free rate ² , % [95% CI]	189 (7.9%) 0.83 [0.68, 1.00] 93.5 [92.5, 94.5]	230 (9.6%) 0.83 [0.68, 1.00] 92.5 [91.4, 93.6]
Disease Free Survival (DFS) Number (%) of patients with event HR [95% CI] ¹ 3 year event-free rate ² , % [95% CI]	192 (8.0%) 0.82 [0.68, 0.99] 93.4 [92.4, 94.4]	236 (9.8%) 0.82 [0.68, 0.99] 92.3 [91.2, 93.4]
Overall Survival (OS)³ Number (%) of patients with event HR [95% CI] ¹ 3 year event-free rate ² , % [95% CI]	80 (3.3%) 0.89 [0.66, 1.21] 97.7 [97.0, 98.3]	89 (3.7%) 0.89 [0.66, 1.21] 97.7 [97.1, 98.3]

HR=Hazard Ratio, CI=Confidence Interval

¹ All analyses stratified by nodal status, protocol version, central hormone receptor status, and adjuvant chemotherapy regimen. Stratification factors are defined according to the randomization data for IDFS.

² 3-year event-free rate derived from Kaplan-Meier estimates

³ Data from first interim analysis

Figure 3 Kaplan-Meier Curve of Invasive Disease Free Survival from APHINITY (ITT Population)

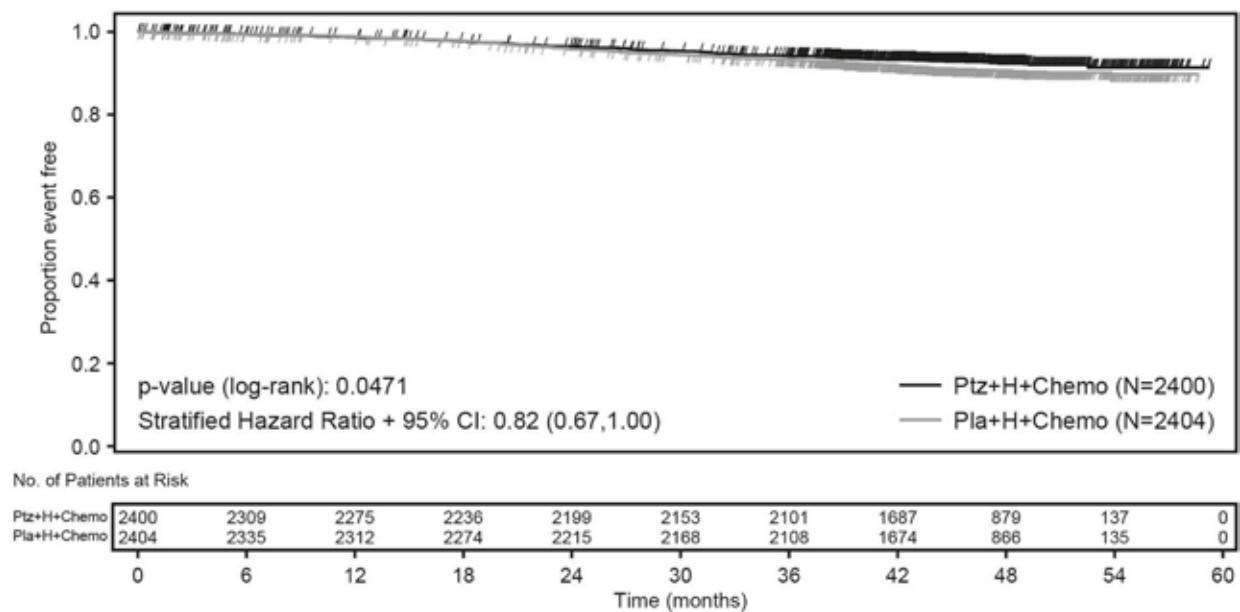


Table 11: Efficacy Results by Baseline Disease Characteristics and Adjuvant Chemotherapy from APHINITY¹

Population	Number of events/Total N (%)		IDFS at 3 year (%, 95% CI)		Unstratified HR (95% CI)
	Pertuzumab + trastuzumab + chemotherapy	Placebo + trastuzumab + chemotherapy	Pertuzumab + trastuzumab + chemotherapy	Placebo + trastuzumab + chemotherapy	
Hormone Receptor Status					
Negative	71/864 (8.2%)	91/858 (10.6%)	92.8 (90.8, 94.3)	91.2 (89.0, 92.9)	0.76 (0.56, 1.04)
Positive	100/1536 (6.5%)	119/1546 (7.7%)	94.8 (93.5, 95.8)	94.4 (93.1, 95.4)	0.86 (0.66, 1.13)
Nodal Status					
Negative	32/897 (3.6%)	29/902 (3.2%)	97.5 (96.3, 98.4)	98.4 (97.3, 99.0)	1.13 (0.68, 1.86)
Positive	139/1503 (9.2%)	181/1502 (12.1%)	92.0 (90.5, 93.3)	90.2 (88.5, 91.6)	0.77 (0.62, 0.96)
Adjuvant Chemotherapy Regimen					
Anthracycline	139/1865 (7.4%)	171/1877 (9.1%)	93.8 (92.6, 94.8)	93.0 (91.8, 94.1)	0.82 (0.66, 1.03)
Non- Anthracycline	32/535 (6.0%)	39/527 (7.4%)	94.9 (92.6, 96.6)	94.0 (91.5, 95.8)	0.82 (0.51, 1.31)

¹ Exploratory analyses without adjusting multiple comparisons, therefore, results are considered descriptive.

16 HOW SUPPLIED/STORAGE AND HANDLING

How Supplied

POHERDY (pertuzumab-dpzb) injection is supplied as a 420 mg/14 mL (30 mg/mL) single-dose vial containing a sterile, preservative-free, clear to slightly opalescent, colorless to pale yellow solution. NDC pending.

Storage and Handling

Store vials refrigerated at 2°C to 8°C (36°F to 46°F) in the original carton until time of use to protect from light. Do not freeze. Do not shake.

17 PATIENT COUNSELING INFORMATION

Left Ventricular Dysfunction

- Advise patients to contact a health care professional immediately for any of the following: new onset or worsening shortness of breath, cough, swelling of the ankles/legs, swelling of the face, palpitations, weight gain of more than 5 pounds in 24 hours, dizziness or loss of consciousness [*see Warnings and Precautions (5.1)*].

Embryo-Fetal Toxicity

- Advise pregnant women and females of reproductive potential that exposure to POHERDY in combination with trastuzumab during pregnancy or within 7 months prior to conception can result in fetal harm [*see Warnings and Precautions (5.2)*]. Advise female patients to contact their healthcare provider with a known or suspected pregnancy [*see Use in Specific Populations (8.1)*].
- Advise females of reproductive potential to use effective contraception during treatment and for 7 months following the last dose of POHERDY in combination with trastuzumab [*see Use in Specific Populations (8.3)*].

Hypersensitivity and Anaphylaxis

- Advise patients to contact their healthcare provider immediately and to report any symptoms of hypersensitivity or anaphylaxis including angioedema, breathing problems, or chest pain [*see Warnings and Precautions (5.4)*].

POHERDY® (pertuzumab-dpzb)

Manufactured by:

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