

Data Sheet

Herclon[®]

Trastuzumab 150 mg and 440 mg powder for concentrate for solution for infusion

Antineoplastic agent

Pharmaceutical Form

Type of dosage form

Powder for concentrate for solution for infusion.

Route of administration

Intravenous infusion.

Sterile/radioactive statement

Sterile product.

Qualitative and Quantitative Composition

Active ingredient

Trastuzumab.

Dosage Preparations: 150 mg single-dose vial and 440 mg multi-dose vial containing powder for concentrate for solution for infusion. Reconstituted Herclon concentrate contains 21 mg/mL of trastuzumab.

Excipients

Herclon 150 mg and 440 mg vials:

L-histidine hydrochloride, L-histidine, a,a-trehalose dihydrate, polysorbate 20.

Solvent vial (for use with the 440 mg vial only):

Water for Injection containing 1.1% benzyl alcohol (Bacteriostatic Water for Injection).

Appearance

Herclon is a white to pale yellow lyophilised powder. The solvent is a clear to slightly opalescent liquid.

Indications

Metastatic breast cancer

Herclon is indicated for the treatment of patients with metastatic breast cancer who have tumours that overexpress HER2:

- as monotherapy for the treatment of those patients who have received one or more chemotherapy regimens for their metastatic disease; or
- in combination with taxanes for the treatment of those patients who have not received chemotherapy for their metastatic disease; or
- in combination with an aromatase inhibitor for the treatment of post-menopausal patients with hormone-receptor positive metastatic breast cancer.

Early breast cancer

Herclon is indicated for the treatment of patients with:

- HER2-positive locally advanced breast cancer in combination with neoadjuvant chemotherapy, followed by adjuvant Herclon; or
- HER2-positive early breast cancer following surgery, sequentially or concurrently with chemotherapy and, if applicable, radiotherapy.

Herclon should only be used in early breast cancer patients with a normal left ventricular ejection fraction.

Advanced gastric cancer

Herclon is indicated in combination with cisplatin and either capecitabine or 5-FU for the treatment of patients with HER2 positive advanced adenocarcinoma of the stomach or gastro-oesophageal junction who have not received prior anti-cancer treatment for their metastatic disease.

Dosage and Administration

HER2 testing is mandatory prior to initiation of Herclon therapy (see Dosage and Administration, Detection of HER2 Overexpression or Gene Amplification).

In order to prevent medication errors it is important to check the vial labels to ensure the medicine being prepared and administered is Herclon (trastuzumab) and not Kadcyła® (trastuzumab emtansine).

Herclon should be administered by a healthcare professional prepared to manage anaphylaxis and adequate life support facilities should be available. Treatment may be administered in an outpatient setting.

In order to improve traceability of biological medicinal products, the trade name of the administered product should be clearly recorded in the patient medical record. Substitution by any other biological medicinal product requires the consent of the prescribing physician.

Herclon should be administered as an intravenous infusion.

DO NOT ADMINISTER HERCLON AS AN INTRAVENOUS PUSH OR BOLUS.

Breast Cancer

Herclon may be used in **metastatic breast cancer** with the once weekly dose schedule, or the, alternative three-weekly schedule. The alternative schedule has been tested in clinical trials but has not been compared directly with the standard weekly regimen.

For treatment of **early breast cancer**, either the once weekly schedule or the three weekly schedule, with loading and subsequent doses, may be used.

Once Weekly Dose Schedule

Loading dose

The recommended initial loading dose of Herclon is 4 mg/kg body weight administered as a 90-minute intravenous infusion. Patients should be observed for fever and chills or other infusion-associated symptoms (see Undesirable Effects). Interruption of the infusion may help control such symptoms. The infusion may be resumed when symptoms abate.

Subsequent doses

The recommended weekly maintenance dose of Herclon is 2 mg/kg body weight. If the initial loading dose was well tolerated, the subsequent doses can be administered as a 30 minute infusion. Patients should be observed for fever and chills or other infusion-associated symptoms (see Undesirable Effects).

Three-Weekly Dose Schedule

Loading dose

The recommended initial loading dose of Herclon is 8 mg/kg body weight administered as a 90-minute intravenous infusion. Patients should be observed for fever and chills or other infusion-associated symptoms (see Undesirable Effects). Interruption of the infusion and/or medication may help to control such symptoms. The infusion may be resumed when symptoms abate.

Subsequent doses

The recommended three-weekly maintenance dose of Herclon is 6 mg/kg body weight administered as a 90-minute intravenous infusion. If the initial loading dose was well tolerated, the subsequent doses can be administered as a 30 minute infusion. Patients should be observed for fevers and chills or other infusion-associated symptoms (see Undesirable Effects).

Switching patients from established weekly schedule to three-weekly doses

The first 6 mg/kg dose should be given a week after the last 2 mg/kg dose. As a precaution, all 6 mg/kg doses should be administered as a 90-minute intravenous infusion. Patients should be observed for fever and chills or other infusion-associated symptoms (see Undesirable Effects). Interruption of the infusion and/or medication may help to control such symptoms. The infusion may

be resumed when symptoms abate. Subsequent maintenance doses of 6 mg/kg are given three-weekly.

Advanced Gastric Cancer

Initial loading dose of Herclon is 8 mg/kg body weight, followed by 6 mg/kg body weight three weeks later and then 6 mg/kg repeated at 3-weekly intervals administered as infusions over approximately 90 minutes. If the initial loading dose was well tolerated, the subsequent doses can be administered as a 30 minute infusion.

In clinical trials, patients with advanced gastric cancer were treated with trastuzumab until progression of disease.

Refer to Clinical Efficacy Studies, Advanced Gastric Cancer for chemotherapy combination dosing.

Duration of treatment

In clinical studies, patients with **metastatic breast cancer** or **advanced gastric cancer** were treated with Herceptin until progression of disease. Patients with **early breast cancer** should be treated for 1 year or until disease recurrence, whichever occurs first. Extending treatment in early breast cancer beyond one year is not recommended).

Missed doses

If the patient misses a dose of Herclon by one week or less, then the usual maintenance dose of Herclon (weekly regimen: 2 mg/kg; three-weekly regimen: 6 mg/kg) should be given as soon as possible. Do not wait until the next planned cycle. Subsequent Herclon maintenance doses (weekly regimen: 2 mg/kg; three-weekly regimen: 6 mg/kg respectively) should then be given according to the previous schedule.

If the patient misses a dose of Herclon by more than one week, a re-loading dose of Herclon should be given over approximately 90 minutes (weekly regimen: 4 mg/kg; three-weekly regimen: 8 mg/kg). Subsequent Herclon maintenance doses (weekly regimen: 2 mg/kg; three-weekly regimen: 6 mg/kg respectively) should then be given according to the previous schedule.

Dose reduction

No reductions in the dose of trastuzumab were made during clinical trials. Patients may continue Herclon therapy during periods of reversible, chemotherapy-induced myelosuppression, but they should be monitored carefully for complications of neutropenia during this time. The specific instructions to reduce or hold the dose of chemotherapy should be followed.

Special Dosage Instructions

Elderly

Data suggest that the disposition of trastuzumab is not altered based on age (see Pharmacokinetics in Special Populations). In clinical trials, elderly patients did not receive reduced doses of trastuzumab.

Children

The safety and efficacy of trastuzumab in paediatric patients have not been established.

Detection of HER2 Overexpression or Gene Amplification

Herclon should only be used in patients whose tumours have HER2 overexpression or HER2 gene amplification.

To ensure accurate and reproducible results, testing must be performed in a specialized laboratory, which can ensure validation of the testing procedures.

HER2 overexpression should be detected using an immunohistochemistry (IHC)-based assessment of fixed tumour blocks. HER2 gene amplification should be detected using *in situ* hybridisation (ISH) of fixed tumour blocks. Examples of ISH include fluorescence *in situ* hybridisation (FISH), chromogenic *in situ* hybridisation (CISH) and silver *in situ* hybridisation (SISH).

For any other method to be used for the assessment of HER2 protein or gene expression, the test method must be precise and accurate enough to demonstrate overexpression of HER2 (it must be able to distinguish between moderate (congruent with 2+) and strong (congruent with 3+) HER2 overexpression).

For full instructions on assay performance and interpretation please refer to the package inserts of validated FISH, CISH and SISH assays. Official recommendations on HER2 testing may also apply.

Breast Cancer

Herclon treatment is only appropriate if there is strong HER2 overexpression, as described by a 3+ score by IHC or a positive ISH result. For patients with an intensity score of 2+ on IHC, confirmation of HER2 positive status by ISH is mandatory.

Advanced Gastric Cancer

Herclon treatment is only appropriate if there is HER2 overexpression, as described by a 3+ IHC score. For cases with a score of less than 3+ by IHC, confirmation of HER2 positive status by ISH is mandatory.

Bright-field ISH technology is recommended for advanced gastric cancer samples to enable evaluation of tumour histology and morphology in parallel. Either FISH or SISH are recommended for detecting HER2 gene amplification in advanced gastric cancer tissue.

Contraindications

Herclon is contraindicated in patients with known hypersensitivity to trastuzumab or to any of its excipients.

Warnings and Precautions

General

Herclon therapy should only be initiated under supervision of a physician experienced in the treatment of cancer patients.

Hypersensitivity reactions including anaphylaxis

Severe hypersensitivity reactions have been infrequently reported in patients treated with trastuzumab. Signs and symptoms include anaphylaxis, urticaria, bronchospasm, angioedema, and/or hypotension. In some cases, the reactions have been fatal. The onset of symptoms generally occurred during an infusion, but there have also been reports of symptom onset after the completion of an infusion. Reactions were most commonly reported in association with the initial infusion.

Patients should be observed closely for hypersensitivity reactions. Herclon infusion should be interrupted in all patients with severe hypersensitivity reactions. In the event of a hypersensitivity reaction, appropriate medical therapy should be administered, which may include adrenaline, corticosteroids, antihistamines, bronchodilators and oxygen. Patients should be evaluated and carefully monitored until complete resolution of signs and symptoms.

Infusion-related reactions (IRRs)

Pre-medication may be used to reduce risk of occurrence of IRRs.

Serious IRRs to trastuzumab infusion including dyspnoea, hypotension, wheezing, bronchospasm, tachycardia, reduced oxygen saturation and respiratory distress and supraventricular tachyarrhythmia have been reported (see Undesirable Effects).

Patients should be observed for IRRs. Interruption of an IV infusion may help control such symptoms and the infusion may be resumed when symptoms abate. These symptoms can be treated with an analgesic/antipyretic such as paracetamol and an antihistamine. Serious reactions have been treated successfully with supportive therapy such as oxygen, intravenous fluids, beta-agonists and corticosteroids. In rare cases, these reactions are associated with a clinical course culminating in a fatal outcome. In other patients with acute onset of signs and symptoms, initial improvement was followed by clinical deterioration and delayed reactions with rapid clinical deterioration have also been reported. Fatalities have occurred within hours or up to one week following an infusion.

Patients who are experiencing dyspnoea at rest due to complications of advanced malignancy or comorbidities may be at increased risk of a fatal infusion reaction. Therefore, these patients should be treated with extreme caution and the risk versus the benefit considered for each patient (see Pulmonary reactions).

Pulmonary reactions

Severe pulmonary events leading to death have been reported with the use of trastuzumab in the post-marketing setting. These events may occur as part of an infusion-related reaction (see Infusion-related reactions (IRRs), Allergic reactions and Hypersensitivity) or with delayed onset. In addition, cases of interstitial lung disease including lung infiltrates, acute respiratory distress syndrome, pneumonia, pneumonitis, pleural effusion, respiratory distress, acute pulmonary oedema and respiratory insufficiency have been reported.

Risk factors associated with interstitial lung disease include prior or concomitant therapy with other anti-neoplastic therapies known to be associated with it such as taxanes, gemcitabine, vinorelbine and radiation therapy. Patients with dyspnoea at rest due to complications of advanced malignancy and comorbidities may be at increased risk of pulmonary events. Therefore, these patients should not be treated with Herclon.

Cardiac dysfunction

General considerations

Patients treated with trastuzumab are at increased risk of developing congestive heart failure (CHF) (New York Heart Association [NYHA] class II - IV) or asymptomatic cardiac dysfunction. These events have been observed in patients receiving trastuzumab therapy alone or in combination with paclitaxel following anthracycline (doxorubicin or epirubicin)-containing chemotherapy. This may be moderate to severe and has been associated with death (see Undesirable Effects). In addition, caution should be exercised in treating patients with increased cardiac risk (e.g. hypertension, documented coronary artery disease, CHF, diastolic dysfunction, older age).

As the half-life of trastuzumab is approximately 28 – 38 days, trastuzumab may persist in the circulation for up to 27 weeks after stopping Herclon treatment (see Pharmacokinetics). Patients who receive anthracycline after stopping Herclon may also be at increased risk of cardiac dysfunction. If possible, physicians should avoid anthracycline-based therapy for up to 27 weeks after stopping Herclon. If anthracyclines are used, the patient's cardiac function should be monitored carefully.

Candidates for treatment with Herclon, especially those with prior exposure to anthracycline and cyclophosphamide (AC) exposure), should undergo baseline cardiac assessment including history and physical examination, ECG, echocardiogram, and/or MUGA scan. Monitoring may help to identify patients who develop cardiac dysfunction, including signs and symptoms of CHF. Cardiac assessments, as performed at baseline, should be repeated every 3 months during treatment and every 6 months following discontinuation of treatments until 24 months from the last administration of Herclon.

If LVEF drops 10 percentage points from baseline and to below 50%, Herclon should be withheld and a repeat LVEF assessment performed within approximately 3 weeks. If LVEF has not improved, or declined further, or clinically significant CHF has developed, discontinuation of Herclon should be strongly considered, unless the benefits for the individual patient are deemed to outweigh the risks.

Patients who develop asymptomatic cardiac dysfunction may benefit from more frequent monitoring (e.g. every 6 - 8 weeks). If patients have a continued decrease in left ventricular function, but remain

asymptomatic, the physician should consider discontinuing therapy if no clinical benefit of Herclon therapy has been seen.

The safety of continuation or resumption of trastuzumab in patients who experience cardiac dysfunction has not been prospectively studied. If symptomatic cardiac failure develops during Herclon therapy, it should be treated with the standard medications for this purpose. In the pivotal trials most patients who developed heart failure improved with standard heart failure treatment consisting of angiotensin-converting enzyme (ACE) inhibitor or angiotensin receptor blocker (ARB) and a β -blocker. The majority of patients with cardiac symptoms and evidence of a clinical benefit of trastuzumab treatment continued on weekly therapy with trastuzumab without additional clinical cardiac events.

Early breast cancer

For patients with early breast cancer, cardiac assessments, as performed at baseline, should be repeated every 3 months during treatment and every 6 months following discontinuation of treatment until 24 months from the last administration of Herclon. In patients who receive anthracycline containing chemotherapy further monitoring is recommended, and should occur yearly up to 5 years from the last administration of Herclon, or longer if a continuous decrease of LVEF is observed.

Patients with history of myocardial infarction (MI), angina pectoris requiring medication, history of or present CHF (NYHA II –IV), other cardiomyopathy, cardiac arrhythmia requiring medication, clinically significant cardiac valvular disease, poorly controlled hypertension (hypertension controlled by standard medication eligible), and haemodynamic effective pericardial effusion were excluded from adjuvant breast cancer clinical trials with trastuzumab.

Neoadjuvant-adjuvant treatment

In patients with early breast cancer eligible for neoadjuvant-adjuvant treatment, Herclon should only be used concurrently with anthracyclines in chemotherapy-naïve patients and only with low-dose anthracycline regimens i.e. with maximum cumulative doses of doxorubicin 180 mg/m² or epirubicin 360 mg/m².

If patients have been treated concurrently with low-dose anthracyclines and Herclon in the neoadjuvant setting, no additional cytotoxic chemotherapy should be given after surgery.

Clinical experience in the neoadjuvant-adjuvant setting is limited in patients above 65 years of age.

Adjuvant treatment

Herclon and anthracyclines should not be given concurrently in the adjuvant treatment setting.

In patients with early breast cancer an increase in the incidence of symptomatic and asymptomatic cardiac events was observed when trastuzumab was administered after anthracycline-containing chemotherapy compared to administration with a non-anthracycline regimen of docetaxel and carboplatin. The incidence was more marked when trastuzumab was administered concurrently with taxanes than when administered sequentially to taxanes. Regardless of the regimen used, most symptomatic cardiac events occurred within the first 18 months.

Risk factors for a cardiac event identified in four large adjuvant studies included advanced age (> 50 years), low level of baseline and declining LVEF (< 55%), low LVEF prior to or following the

initiation of paclitaxel treatment, trastuzumab treatment, and prior or concurrent use of anti-hypertensive medications. In patients receiving trastuzumab after completion of adjuvant chemotherapy the risk of cardiac dysfunction was associated with a higher cumulative dose of anthracycline given prior to initiation of trastuzumab and a high body mass index (BMI) ($> 25 \text{ kg/m}^2$).

Metastatic breast cancer

Herclon and anthracycline should not be given concurrently in the metastatic breast cancer setting.

Advanced Gastric Cancer

In advanced gastric cancer, patients with a history of documented congestive heart failure, angina pectoris requiring medication, evidence of transmural myocardial infarction on ECG, poorly controlled hypertension (systolic BP $>180 \text{ mmHg}$ or diastolic BP $>100 \text{ mmHg}$), clinically significant valvular heart disease, high risk uncontrollable arrhythmias, baseline LVEF $<50\%$ (measured by echocardiography or MUGA) were excluded from Study BO18255 (ToGA) according to the study protocol.

Benzyl alcohol

Benzyl alcohol, used as a preservative in Bacteriostatic Water for Injection in the 440 mg multidose vial, has been associated with toxicity in neonates and children up to 3 years old. When administering Herclon to a patient with a known sensitivity to benzyl alcohol, Herclon should be reconstituted with water for injection, and only one dose per Herclon vial should be used. Any unused portion must be discarded. Sterile water for injection, used to reconstitute the 150 mg single dose vials, does not contain benzyl alcohol.

Ability to Drive and Use Machines

No studies on the effects on the ability to drive and to use machines have been performed. Patients experiencing infusion-related symptoms should be advised not to drive or use machines until symptoms resolve completely.

Interactions

No formal interaction studies have been performed with trastuzumab in humans. Clinically significant interactions with the concomitant medication used in clinical trials have not been observed. A comparison of serum levels of trastuzumab given in combination with cisplatin, doxorubicin or epirubicin-plus-cyclophosphamide has not suggested the possibility of any interaction.

Administration of paclitaxel in combination with trastuzumab resulted in a slightly less than two-fold decrease in trastuzumab clearance in a non-human primate study and in a 1.5-fold increase in trastuzumab serum levels in clinical studies. Paclitaxel pharmacokinetics determined during the fourth cycle of the alternative 3-weekly trastuzumab regimen ($n = 25$) were not altered appreciably, relative to parameters determined during the initiation of paclitaxel, prior to introduction of trastuzumab. Similarly, docetaxel pharmacokinetics determined during the first dose of trastuzumab in the standard weekly

regimen ($n = 10$) were not altered appreciably relative to those determined 2 weeks earlier for docetaxel alone.

A pharmacokinetic interaction substudy of BO18255 (ToGA) performed in male and female Japanese patients with advanced gastric cancer showed that co-administration of trastuzumab and capecitabine and cisplatin had no significant effects on the pharmacokinetics of the two chemotherapy agents compared with co-administration of the two agents without trastuzumab. The pharmacokinetics of trastuzumab were not evaluated in this study.

Use in Special Populations

Pregnancy

Pregnancy Category: D

Herclon should be avoided during pregnancy and since trastuzumab may persist in the circulation for up to 27 weeks, pregnancy should be avoided for 7 months after the last dose of Herclon, unless the anticipated benefit for the mother outweighs the unknown risk to the foetus.

In the post-marketing setting, cases of foetal renal growth and/or function impairment in association with oligohydramnios, some associated with fatal pulmonary hypoplasia of the foetus, have been reported in pregnant women receiving trastuzumab. Women of childbearing potential should be advised to use effective contraception during treatment with Herclon and for at least 7 months after treatment has concluded. Women who become pregnant should be advised of the possibility of harm to the foetus. If a pregnant woman is treated with Herclon, close monitoring by a multidisciplinary team is desirable.

Nursing mothers

It is not known whether trastuzumab is secreted in human breast milk. As human IgG is secreted into human breast milk, and the potential for harm to the infant is unknown, breast-feeding should be avoided during Herclon therapy and for 7 months after the last dose of Herclon (See Preclinical Safety, Other).

Undesirable Effects

The adverse drug reactions listed in this section fall into the following categories: very common ($\geq 1/10$); common ($\geq 1/100$ to $<1/10$); uncommon ($\geq 1/1,000$ to $<1/100$); rare ($\geq 1/10,000$ to $<1/1,000$); very rare ($<1/10,000$); not known (cannot be estimated from the available data). Within each frequency grouping, adverse reactions are presented in order of decreasing seriousness.

Presented in Table 1 below are adverse reactions that have been reported in association with the use of trastuzumab alone, or in combination with chemotherapy in the below pivotal clinical trials as well as in the post-marketing setting.

Early Breast Cancer

- **BO16348 (HERA)**: Trastuzumab arm (n = 1678). Control arm (n = 1708)
- **B-31/N9831 Joint Analysis**: Trastuzumab arms (n = 2345). Control arm (n = 1673)
- **BCIRG 006**: Trastuzumab arm (n = 2133). Control arm (n = 1041)
- **BO16216 (TanDEM)**: Trastuzumab arm (n = 161). Control arm (n = 161)
- **MO16432 (NOAH)**: Trastuzumab arm (n = 115). Control arm (n = 116)

Metastatic Breast Cancer

- **H0648g / H0649g**: Trastuzumab arms (n = 469 and n = 222 respectively)
- **M77001**: Trastuzumab arm (n = 92). Control arm (n = 94).

Advanced Gastric Cancer

- **BO18255 (ToGA)**: Trastuzumab arm (n = 294). Control arm (n = 290)

All terms included are based on the highest percentage seen in pivotal clinical trials.

Table 1: Adverse reactions

System organ class	Adverse reaction¹	Frequency
Infections and infestations	Nasopharyngitis	Very common
	Infection	Very common
	Neutropenic sepsis	Common
	Cystitis	Common
	Herpes zoster	Common
	Influenza	Common
	Pharyngitis	Common
	Sinusitis	Common
	Skin infection	Common
	Rhinitis	Common
	Upper respiratory tract infection	Common
	Urinary tract infection	Common
	Erysipelas	Common
	Cellulitis	Common
Sepsis	Uncommon	
Neoplasms benign, malignant and unspecified (incl. Cysts and polyps)	Malignant neoplasm progression	Not known
	Neoplasm progression	Not known
Blood and lymphatic system disorders	Febrile neutropenia	Very common
	Anaemia	Very common
	Thrombocytopenia	Common
	White blood cell count decreased / leukopenia	Common
	Neutropenia	Common

System organ class	Adverse reaction¹	Frequency
	Hypoprothrombinaemia	Not known
Immune system disorders	Hypersensitivity	Common
	² Anaphylactic reaction	Not known
	² Anaphylactic shock	Not known
Metabolism and nutrition disorders	Weight Decreased/Weight Loss	Very common
	Weight Increased	Very common
	Decreased appetite	Very common
	Anorexia	Common
	Hyperkalaemia	Not known
Psychiatric disorders	Insomnia	Very common
	Depression	Common
	Anxiety	Common
	Thinking abnormal	Common
Nervous system disorders	Tremor	Very common
	Dizziness	Very common
	Headache	Very common
	Dysgeusia	Very common
	Paraesthesia	Very common
	Hypoaesthesia	Very common
	Peripheral neuropathy	Common
	Hypertonia	Common
	Somnolence	Common
	Ataxia	Common
	Paresis	Rare
	Brain oedema	Not known
Eye disorders	Conjunctivitis	Very common
	Lacrimation increased	Very Common
	Dry eye	Common
	Papilloedema	Not known
	Retinal haemorrhage	Not known
Ear and Labyrinth Disorders Cardiac disorders	Deafness	Uncommon
	³ Blood pressure decreased	Very common
	³ Blood pressure increased	Very common
	³ Heart beat irregular	Very common
	³ Palpitation	Very common
	³ Cardiac flutter	Very common
⁴ Ejection fraction decreased	Very Common	

System organ class	Adverse reaction ¹	Frequency
	² Cardiac failure (congestive)	Common
	^{2,3} Supraventricular tachyarrhythmia	Common
	Cardiomyopathy	Common
	Pericardial effusion	Uncommon
	Cardiogenic shock	Not known
	Pericarditis	Not known
	Bradycardia	Not known
	Gallop rhythm present	Not known
Vascular disorders	Lymphoedema	Very common
	Hot flush	Very common
	^{2,3} Hypotension	Common
	Hypertension	Common
	Vasodilation	Common
Respiratory, thoracic and mediastinal disorders	^{2,3} Wheezing	Very common
	² Dyspnoea	Very common
	Cough	Very common
	Epistaxis	Very common
	Rhinorrhoea	Very common
	Oropharyngeal pain	Very common
	Asthma	Common
	Lung disorder	Common
	² Pleural effusion	Common
	² Pneumonia	Common
	Pneumonitis	Uncommon
	² Pulmonary fibrosis	Not known
	² Respiratory distress	Not known
	² Respiratory failure	Not known
	² Lung infiltration	Not known
	² Acute pulmonary oedema	Not known
	² Acute respiratory distress syndrome	Not known
	² Bronchospasm	Not known
	² Hypoxia	Not known
	² Oxygen saturation decreased	Not known
	Laryngeal oedema	Not known
	² Orthopnoea	Not known
Pulmonary oedema	Not known	
Gastrointestinal disorders	Diarrhoea	Very common

System organ class	Adverse reaction¹	Frequency
	Vomiting	Very common
	Nausea	Very common
	Lip swelling	Very common
	Abdominal pain	Very common
	Stomatitis	Very common
	Pancreatitis	Very common
	Constipation	Very common
	Dyspepsia	Very common
	Haemorrhoids	Common
	Dry mouth	Common
Hepatobiliary disorders	Hepatocellular Injury	Common
	Hepatitis	Common
	Liver Tenderness	Common
	Jaundice	Rare
	Hepatic Failure	Not known
Skin and subcutaneous disorders	Erythema	Very common
	Rash	Very common
	Swelling face	Very common
	Palmar-plantar erythrodysesthesia syndrome	Very common
	Nail disorder	Very common
	Alopecia	Very common
	Dry skin	Common
	Ecchymosis	Common
	Hyperhidrosis	Common
	Maculopapular rash	Common
	Acne	Common
	Onychoclasia	Common
	Pruritus	Common
	Dermatitis	Common
	Urticaria	Uncommon
Angioedema	Not known	
Musculoskeletal and connective tissue disorders	Arthralgia	Very common
	Muscle tightness	Very common
	Myalgia	Very common
	Arthritis	Common
	Back pain	Common
	Bone pain	Common

System organ class	Adverse reaction¹	Frequency
	Muscle spasms	Common
	Neck pain	Common
	Pain in extremity	Common
Renal and urinary conditions	Renal disorder	Common
	Glomerulonephritis membranous	Not known
	Glomerulonephropathy	Not known
	Renal failure	Not known
Pregnancy, puerperium and perinatal disorders	Oligohydramnios	Not known
Reproductive system and breast disorders	Breast inflammation/mastitis	Common
General disorders and administration site conditions	Asthenia	Very common
	Chest pain	Very common
	Chills	Very common
	Fatigue	Very common
	Influenza-like symptoms	Very common
	Infusion related reaction	Very common
	Pain	Very common
	Pyrexia	Very common
	Peripheral oedema	Very common
	Mucosal inflammation	Very common
	Malaise	Common
	Oedema	Common
Injury, poisoning and procedural complications	Nail toxicity	Very common
	Contusion	Common

¹ Adverse drug reactions (ADRs) were identified as events that occurred with at least a 2% difference compared to the control arm in at least one of the major randomised clinical trials

2 Denotes adverse reactions that have been reported in association with a fatal outcome.

3 Denotes adverse reactions that are reported largely in association with Infusion-related reactions. Specific percentages for these are not available.

4 Observed with combination therapy following anthracyclines and combined with taxanes

The following information is relevant to all indications:

Infusion-related reactions (IRRs) and Hypersensitivity

IRRs such as chills and/or fever, dyspnoea, hypotension, wheezing, bronchospasm, tachycardia, reduced oxygen saturation and respiratory distress were seen in all trastuzumab clinical trials (see Warnings and Precautions).

IRRs may be clinically difficult to distinguish from hypersensitivity reactions.

The rate of IRRs of all grades varied between studies depending on the indication, whether trastuzumab was given concurrently with chemotherapy or as monotherapy and data collection methodology.

In early breast cancer, the rate of IRRs ranged from 18% to 54% in the trastuzumab containing arm compared to 6% to 50% in the comparator arm (which may have contained other chemotherapy). Severe reactions (grade 3 and above) ranged from 0.5% to 6% in the trastuzumab containing arm compared to 0.3% to 5 % in the comparator arm.

In metastatic breast cancer, the rate of IRRs ranged from 49% to 54% in the trastuzumab containing arm compared to 36% to 58% in the comparator arm (which may have contained other chemotherapy), Severe reactions (grade 3 and above) ranged from 5 % to 7% in the trastuzumab containing arm compared to 5% to 6% in the comparator arm.

Anaphylactoid reactions were observed in isolated cases.

Cardiac dysfunction

Congestive heart failure (NYHA II-IV) is a common adverse reaction to Herceptin. It has been associated with fatal outcome. Signs and symptoms of cardiac dysfunction, such as dyspnoea, orthopnoea, increased cough, pulmonary oedema, S3 gallop, or reduced ventricular ejection fraction, have been observed in patients treated with trastuzumab (see Warnings and Precautions).

Early breast cancer

Neoadjuvant-adjuvant setting

In the clinical trial setting, when trastuzumab was administered concurrently with neoadjuvant chemotherapy containing three to four cycles of a neoadjuvant anthracycline (cumulative doxorubicin dose 180 mg/m² or epirubicin dose 360 mg/m²) overall, the incidence of symptomatic cardiac dysfunction was 1.7 % in the trastuzumab arm.

Adjuvant setting

In 3 pivotal clinical trials of adjuvant trastuzumab given in combination with chemotherapy the incidence of grade 3/4 cardiac dysfunction (symptomatic CHF) was similar in patients who were administered chemotherapy alone and in patients who were administered trastuzumab sequentially to a taxane (0.3 - 0.4%). The rate was highest in patients who were administered trastuzumab concurrently with a taxane (2.0%). At 3 years, the cardiac event rate in patients receiving AC→P (doxorubicin plus cyclophosphamide followed by paclitaxel) + H (trastuzumab) was estimated at 3.2%, compared with 0.8% in AC→P treated patients. No increase in the cumulative incidence of cardiac events was seen with further follow-up at 5 years.

At 5.5 years, the rates of symptomatic cardiac or LVEF events were 1.0%, 2.3%, and 1.1% in the AC→D (doxorubicin plus cyclophosphamide, followed by docetaxel), AC→DH (doxorubicin plus cyclophosphamide, followed by docetaxel plus trastuzumab), and DCarbH (docetaxel, carboplatin and trastuzumab) treatment arms, respectively. For symptomatic CHF (Grade 3 - 4), the 5-year rates were 0.6%, 1.9%, and 0.4% in the AC→D, AC→DH, and DCarbH treatment arms, respectively. The overall risk of developing symptomatic cardiac events was low and similar for patients in AC→D and DCarbH arms; relative to both the AC→D and DCarbH arms there was an increased risk of developing a symptomatic cardiac event for patients in the AC→DH arm, being discernable by a continuous increase in the cumulative rate of symptomatic cardiac or LVEF events up to 2.3% compared to approximately 1% in the two comparator arms (AC→D and DCarbH).

When trastuzumab was administered after completion of adjuvant chemotherapy NYHA class III-IV heart failure was observed in 0.6% of patients in the 1-year arm after a median follow-up of 12 months. After a median follow-up of 3.6 years the incidence of severe CHF and left ventricular dysfunction after 1 year trastuzumab therapy remained low at 0.8% and 9.8%, respectively. After a median follow-up of 8 years, the incidence of severe CHF (NYHA III & IV) following 1 year of trastuzumab therapy (combined analysis of the two trastuzumab treatment arms) was 0.89%, and the rate of mild symptomatic and asymptomatic left ventricular dysfunction was 6.35%.

Reversibility of severe CHF (defined as a sequence of at least two consecutive LVEF values ³ 50% after the event) was evident for 70.0% of trastuzumab-treated patients. Reversibility of mild symptomatic and asymptomatic left ventricular dysfunction was demonstrated for 83.1% of trastuzumab-treated patients. Approximately 10% of cardiac endpoints occurred after completion of trastuzumab.

Metastatic breast cancer

Depending on the criteria used to define cardiac dysfunction, the incidence in the pivotal metastatic trials varied between 9% and 12% in the trastuzumab + paclitaxel subgroup, compared with 1% - 4% for the paclitaxel alone subgroup. For trastuzumab monotherapy, the rate was 6% - 9%. The highest rate of cardiac dysfunction was seen in patients receiving trastuzumab + anthracycline/cyclophosphamide (27%), significantly higher than in the anthracycline/cyclophosphamide alone subgroup (7% - 10%). In study M77001 with prospective monitoring of cardiac function, the incidence of symptomatic heart failure was 2.2% in patients receiving trastuzumab and docetaxel, compared with 0% in patients receiving docetaxel alone. Most of the patients (79%) who developed cardiac dysfunction in these trials experienced an improvement after receiving standard treatment for heart failure.

Advanced Gastric Cancer

In Study BO18255 (ToGA), at screening, the median LVEF value was 64% (range 48% - 90%) in the FP arm and 65% (range 50% - 86%) in the trastuzumab + FP arm.

The majority of the LVEF decreases noted in BO18255 Study (ToGA) were asymptomatic, with the exception of 1 patient in the trastuzumab arm whose LVEF decrease coincided with cardiac failure.

Table 2: Summary of LVEF change from baseline (BO18255 Study)

LVEF Decrease [#] : Lowest Post-screening Value	FP (n = 290) (% of patients in each treatment arm)	FP + H (n = 294) (% of patients in each treatment arm)
LVEF decrease ³ 10% to <50%	1.1%	4.6%
Absolute Value <50%	1.1%	5.9%
LVEF decrease ³ 10% to ³ 50%	11.8%	16.5%

FP: Fluoropyrimidine/cisplatin; FP+H: Fluoropyrimidine/cisplatin + trastuzumab; [#]Only includes patients whose method of assessment at that visit is the same as at their initial assessments (FP: n = 187 and FP + H: n = 237)

Table 3: Cardiac Adverse Events (BO18255 Study)

	FP (n = 290) (% of patients in each treatment arm)	FP + H (n = 294) (% of patients in each treatment arm)
Total Cardiac Events	6%	6%
³ Grade 3 NCI-CTC(v3.0) criteria	3% ^a	1% ^b

FP: Fluoropyrimidine/cisplatin; FP+H: Fluoropyrimidine/cisplatin + trastuzumab; ^a 9 patients experienced 9 Events; ^b 4 patients experienced 5 Events

Overall, there were no significant differences in cardiac dysfunction between the treatment arm and the comparator arm.

Haematological toxicity

Breast Cancer

Monotherapy – Study H0649g

Haematological toxicity is infrequent following the administration of trastuzumab monotherapy in the metastatic setting, WHO Grade 3 leucopenia, thrombocytopenia and anaemia occurring in < 1% of patients. No WHO Grade 4 toxicities were observed.

Combination Therapy – Studies H0648g and M77001

WHO Grade 3 or 4 haematological toxicity was observed in 63% of patients treated with trastuzumab and an anthracycline/cyclophosphamide compared to an incidence of 62% in patients treated with the anthracycline/cyclophosphamide combination without trastuzumab.

There was an increase in WHO Grade 3 or 4 haematological toxicity in patients treated with the combination of trastuzumab and paclitaxel compared with patients receiving paclitaxel alone (34% vs. 21%). Haematological toxicity was also increased in patients receiving trastuzumab and docetaxel, compared with docetaxel alone (32% grade 3/4 neutropenia versus 22%, using HCl-CTC criteria). The incidence of febrile neutropenia/neutropenic sepsis was also increased in patients treated with trastuzumab + docetaxel (23% versus 17% for patients treated with docetaxel alone).

Early breast cancer setting – BO16348 (HERA) Study

Using NCI-CTC criteria, in the HERA trial, 0.4% of trastuzumab-treated patients experienced a shift of 3 or 4 grades from baseline, compared with 0.6% in the observation arm.

Advanced Gastric Cancer

The most frequently reported adverse events categorised under the Blood and Lymphatic System Disorders SOC (Grade ≥3) are shown below (Table 4) by trial treatment.

Table 4: Blood and Lymphatic System Disorders (SOC) Adverse Events >1%

	FP (n=290) (% of patients in each treatment arm)	FP + H (n=294) (% of patients in each treatment arm)
Neutropenia	30%	27%
Anaemia	10%	12%
Febrile neutropenia	3%	5%
Thrombocytopenia	3%	5%

FP: Fluoropyrimidine/cisplatin; FP+H: Fluoropyrimidine/cisplatin + trastuzumab

The total percentage of patients who experienced an adverse event of ³ Grade 3 NCI CTCAE v3.0 categorised under this SOC were 38% in the FP arm and 40% in the FP + H arm.

Overall, there were no significant differences in haematotoxicity between the treatment arm and the comparator arm.

Hepatic and renal toxicity

Breast Cancer

WHO Grade 3 or 4 hepatic toxicity was observed in 12% of patients following administration of trastuzumab as single agent, in the metastatic setting. This toxicity was associated with progression of disease in the liver in 60% of these patients.

WHO Grade 3 or 4 hepatic toxicity was less frequently observed among patients receiving trastuzumab and paclitaxel than among patients receiving paclitaxel (7% compared with 15%). No WHO Grade 3 or 4 renal toxicity was observed.

Advanced Gastric Cancer

In Study BO18255 (ToGA) no significant differences in hepatic and renal toxicity were observed between the two treatment arms.

NCI-CTCAE (v3.0) grade ≥3 renal toxicity was not significantly higher in patients receiving trastuzumab than those in the fluoropyrimidine/cisplatin arm (3% and 2% respectively).

NCI-CTCAE (v3.0) grade ≥3 adverse events in the Hepatobiliary Disorders SOC: Hyperbilirubinaemia was the only reported adverse event and was not significantly higher in patients receiving trastuzumab than those in the fluoropyrimidine/cisplatin arm (1% and <1% respectively).

Diarrhoea

Breast Cancer

Monotherapy – Study H0649G

Of patients treated with trastuzumab monotherapy in the metastatic setting, 27% experienced diarrhoea. In the HERA trial (median follow-up 1 year), 7% of trastuzumab-treated patients had diarrhoea.

Combination Therapy – Studies H0648g and M77001

An increase in the incidence of diarrhoea, primarily mild to moderate in severity, has been observed in patients receiving trastuzumab in combination with chemotherapy compared with patients receiving chemotherapy-alone or trastuzumab-alone.

Early breast cancer setting – HERA Study

In the HERA trial, 8% of trastuzumab treated patients experienced diarrhoea during the first year of treatment.

Advanced Gastric Cancer

In Study BO18255 (ToGA), 109 patients (37%) in the trastuzumab treatment arm versus 80 patients (28%) in the comparator arm experienced any grade diarrhoea. Four percent (4%) of patients in the fluoropyrimidine/cisplatin arm experienced Grade 3 diarrhoea vs. 9% in the trastuzumab arm.

Infection

An increased incidence of infections, primarily mild upper respiratory infections of minor clinical significance or catheter infections, has been observed primarily in patients treated with trastuzumab + paclitaxel compared with patients receiving paclitaxel alone or trastuzumab-alone.

Overdosage

There is no experience with overdosage in human clinical trials. Single doses higher than 10 mg/kg have not been tested.

Treatment of overdose should consist of general supportive measures.

For information on the management of overdose, contact the Poisons Information Centre (in Australia call 13 11 26; in New Zealand call 0800 767 766).

Pharmacological Properties and Effects

Pharmacodynamic Properties

Mechanism of action

Trastuzumab is a recombinant humanised monoclonal antibody that selectively targets the extracellular domain of the human epidermal growth factor receptor 2 protein (HER2). The antibody is an IgG₁

isotype that contains human framework regions with the complementarity-determining regions of a murine anti-p185 HER2 antibody that binds to HER2.

The HER2 proto-oncogene or c-erbB2 encodes for a single transmembrane spanning, receptor-like protein of 185 kDa, which is structurally related to the epidermal growth factor receptor. Overexpression of HER2 is observed in 25% - 30% of primary breast cancers and 6.8% - 42.6% of advanced gastric cancers. A consequence of HER2 gene amplification is an increase in HER2 protein expression on the surface of these tumour cells, which results in a constitutively activated HER2 protein.

Studies indicate that patients whose tumours have amplification or overexpression of HER2 have a shortened disease-free survival compared to patients whose tumours do not have amplification or overexpression of HER2.

Trastuzumab has been shown, both in *in vitro* assays and in animals, to inhibit the proliferation of human tumour cells that overexpress HER2. *In vitro*, trastuzumab-mediated antibody-dependent cell-mediated cytotoxicity (ADCC) has been shown to be preferentially exerted on HER2 overexpressing cancer cells compared with cancer cells that do not overexpress HER2.

Clinical efficacy studies

Early Breast Cancer

Trastuzumab in combination with neoadjuvant-adjuvant chemotherapy

The use of trastuzumab for the neoadjuvant-adjuvant treatment of locally advanced breast cancer has been studied in Study MO16432 (NOAH), a multicentre randomized trial, designed to investigate the concurrent administration of trastuzumab with neoadjuvant chemotherapy, including both an anthracycline and a taxane, followed by adjuvant trastuzumab, up to a total treatment duration of 1 year. The study recruited patients with newly diagnosed locally advanced (Stage III) or inflammatory breast cancer. Patients with HER2+ tumours were randomized to receive either neoadjuvant chemotherapy concurrently with neoadjuvant-adjuvant trastuzumab (n = 116), or neoadjuvant chemotherapy alone (n = 118).

Trastuzumab was administered concurrently with 10 cycles of neoadjuvant chemotherapy as follows;

- Doxorubicin (60 mg/m²) and paclitaxel (150 mg/m²) in combination with trastuzumab (8 mg/kg loading dose, followed by 6 mg/kg maintenance, administered 3-weekly) for 3 cycles, followed by
- Paclitaxel (175 mg/m²) and trastuzumab (6mg/kg, administered 3-weekly) for 4 cycles, followed by
- CMF on day 1 and 8 every 4 weeks for 3 cycles, in combination with 4 cycles of trastuzumab (6mg/kg administered 3-weekly), followed by
- up to 7 additional cycles of trastuzumab (6mg/kg, administered 3-weekly) alone to complete 1 year after starting trastuzumab

The primary endpoint for the study, event-free survival (EFS), was defined as the time from randomization to disease recurrence or progression (local, regional, distant or contralateral), or death of any cause. The efficacy results from NOAH (full analysis population, defined as all patients who were randomized in the study following the intent-to-treat principle, with the exception of 3 patients

whose data could not be evaluated) are summarized in the table below. The median duration of follow-up in the trastuzumab arm was 3.8 years.

Table 5: Overview of Efficacy Analyses MO16432 (NOAH)

Parameter	Chemo + trastuzumab n = 115	Chemo only n = 116	p-value	HR (95% CI)
Event-free survival (EFS)				
No. patients with event	46	59	p = 0.0275	0.65 (0.44, 0.96)
Total pathological complete response[^] (95% CI)	40% (31.0, 49.6)	20.7% (13.7, 29.2)	p = 0.0014	

[^] defined as absence of any invasive cancer both in the breast and axillary nodes; HR: hazard ratio

The addition of trastuzumab to neoadjuvant chemotherapy, followed by adjuvant trastuzumab for a total duration of 52 weeks, resulted in a 35% reduction in the risk of disease recurrence/progression. The hazard ratio translates into an absolute benefit, in terms of 3-year event-free survival rate estimates of 13 percentage points (65 % vs. 52 %) in favour of the trastuzumab arm.

To date, results are not available comparing the efficacy of trastuzumab administered with chemotherapy in the adjuvant setting with that obtained in the neoadjuvant/adjuvant setting.

Trastuzumab in combination with adjuvant chemotherapy

The use of trastuzumab in the setting of early breast cancer (after surgery and in association with chemotherapy and, if applicable radiotherapy) has been studied in four multicentre, randomised, phase III trials of patients with HER2 positive breast cancer who have completed surgery. In these clinical trials, localised breast cancer was limited to operable, primary adenocarcinoma of the breast with positive axillary nodes or node negative disease with additional indicators of a higher degree of risk.

The design of these studies is summarized in Table 6 and efficacy results are presented in Tables 7 - 10.

Table 6: Clinical Trials in Early Breast Cancer

	HERA trial n = 3386	NSAPB B-31 and NCCTG N9831 trials (joint analysis) n = 3763	BCIRG 006 n = 3222
Eligible patients	Node positive or node negative [n = 1098] and tumour size >1 cm; <i>Protocol initially unrestricted but amended and node negative patients with tumours ≤1 cm [n =93, 8.5%] and node negative patients with tumours >1 and ≤2 cm [n = 509,46.4%] were included</i>	Node positive or node negative [n = 190] and tumour size · >2 cm regardless of hormonal status; or · >1 cm and ER–ve [n = 63 node-negative and tumour size ≤2 cm])	Node positive or node negative and at least 1 of the following: · tumour size > 2 cm and ER and PR -ve, or · histologic and/or nuclear grade 2-3, or · age < 35 years.
Trastuzumab	Loading dose 8 mg/kg,	Loading dose 4 mg/kg,	Loading dose 4 mg/kg,

	HERA trial <i>n</i> = 3386	NSAPB B-31 and NCCTG N9831 trials (joint analysis) <i>n</i> = 3763	BCIRG 006 <i>n</i> = 3222
dosage regimen	followed by 6 mg/kg (q3w)	followed by 2 mg/kg (q1w)	followed by 2 mg/kg (q1w). After chemo, 6 mg/kg (q3w)
Duration of trastuzumab treatment	1yr or 2yrs	52 weeks	52 weeks
Chemotherapy regimen(s)	Various	AC (q3w) followed by IV paclitaxel as a continuous IV infusion (AC→P). Paclitaxel: 80 mg/m ² q1w for 12 weeks or 175 mg/m ² q3w for 4 cycles (day 1 of each cycle)	AC followed by docetaxel (AC→D) or docetaxel and carboplatin (DCarb) Docetaxel (IV infusion over 60 min): (AC→D): 100 mg/m ² q3w for 4 cycles or (DCarb): 75 mg/m ² q3w for 6 cycles Carboplatin (at target AUC): 6 mg/mL/min (IV infusion over 30 - 60 min) q3w for a total of 6 cycles.
Timing of trastuzumab in relation to chemotherapy	After completion of (neo)adjuvant ^a	Concurrent (AC→PH) or sequential (AC→P→H)	Concurrent (AC→DH and DCarbH)
Median follow-up	1 year (initial evaluation) 8 years (follow-up evaluation)	2 years	3 years

AC = doxorubicin + cyclophosphamide; q3w = every 3 weeks; q1w = weekly chemo = chemotherapy; ^a 89% of subjects received adjuvant chemotherapy; 5% received neoadjuvant chemotherapy and 6% received a combination of neoadjuvant and adjuvant chemotherapy.

The HERA trial was designed to compare 1 and 2 years of 3-weekly trastuzumab treatment vs. observation in patients with HER2 positive breast cancer following surgery, established chemotherapy and radiotherapy (if applicable). In addition, a comparison of 2 years trastuzumab treatment vs. 1 year trastuzumab treatment was performed. Patients assigned to receive trastuzumab were given an initial loading dose of 8 mg/kg, followed by 6 mg/kg every 3 weeks for either 1 or 2 years. The efficacy results from the HERA trial are summarized in the following table:

Table 7: Efficacy Results from the HERA Trial at 12 months¹ and 8 years² of median follow up

Parameter	Observation	Trastuzumab 1yr treatment	p-value	HR (95% CI)
<u>Disease free survival</u>				
No. of patients with event (1 year ¹)	12.9%	7.5%	<0.0001	0.54 (0.44,0.67)
No of patients with event (8 year ²)	33.6%	27.7%	<0.0001	0.76 (0.67, 0.86)

Parameter	Observation	Trastuzumab 1yr treatment	p-value	HR (95% CI)
Overall Survival				
No. of patients with event (1 year ¹)	2.4%	1.8%	0.24	0.75 (0.47, 1.21)
No. of patients with event (8 year ²)	20.6%	16.3%	0.0005	0.76 (0.65, 0.88)

HR: Hazard ratio; ¹ co-primary endpoint of DFS of 1 year vs. observation met the pre-defined statistical boundary; ² final analysis (includes crossover of 52% of patients from the observation arm to Herceptin).

The HERA study included a subgroup of patients ($n = 602$) with small tumours (<2 cm) and node-negative disease. In this subgroup, the relative risk reduction was similar to the overall trial population (HR = 0.50; 95% CI 0.21 - 1.15). However, the benefit in terms of absolute difference in rate of recurrence after 1 year of follow-up was smaller (2.7% recurrence rate with trastuzumab vs. 5.5% with observation).

In the final analysis (8 year median follow up) extending trastuzumab treatment for a duration of 2 years did not show additional benefit over treatment for 1 year [DFS HR in the intent to treat (ITT) population of 2 years vs. 1 year = 0.99 (95% CI: 0.87, 1.13); p-value = 0.90 and OS HR = 0.98 (0.83, 1.15); p-value = 0.78]. The rate of asymptomatic cardiac dysfunction was increased in the 2-year treatment arm (8.1% vs. 4.6% in the 1-year treatment arm). More patients experienced at least one grade 3 or 4 adverse event in the 2-year treatment arm (20.4%) compared with the 1-year treatment arm (16.3%).

The efficacy results from the joint analysis of the NCCTG 9831 and NSABP B-31 trials at the time of the definitive analysis of DFS are summarized in the following table:

Table 8: Summary of Efficacy results from NSAPB B-31 and NCCTG N9831 trials (joint analysis) at the time of the definitive DFS analysis

Parameter	AC→P	AC→PH	p-value	HR (95% CI)
Disease recurrence				
Rate (trastuzumab vs. observation)	15.5%	8.0%	< 0.0001	0.48 (0.39, 0.59)
Survival				
Deaths (trastuzumab vs. observation)	5.5%	3.7%	0.014	0.67 (0.48, 0.92)

A: doxorubicin; C: cyclophosphamide; P: paclitaxel; H: trastuzumab; HR: Hazard ratio

* At median duration of follow up was 1.8 years for the patients in the AC→P arm and 2.0 years for patients in the AC→PH arm.

** p value for OS did not cross the pre-specified statistical boundary for comparison of AC→PH vs. AC→P

For the primary endpoint, DFS, the addition of Herceptin to paclitaxel chemotherapy resulted in a 52 % decrease in the risk of disease recurrence. The hazard ratio translates into an absolute benefit, in terms of 3-year disease-free survival rate estimates of 11.8 percentage points (87.2 % versus 75.4 %) in favour of the AC→PH (Herceptin) arm.

At the time of a safety update, after a median of 3.5-3.8 years follow up, an analysis of DFS reconfirms the magnitude of the benefit shown in the definitive analysis of DFS. Despite the cross-over to Herceptin in the control arm, the addition of Herceptin to paclitaxel chemotherapy resulted in a 52 %

decrease in the risk of disease recurrence. The addition of Herceptin to paclitaxel chemotherapy also resulted in a 37 % decrease in the risk of death.

The pre-planned final analysis of OS from the joint analysis of studies NSABP B-31 and NCCTG N9831 was performed when 707 deaths had occurred (median follow-up 8.3 years in the AC→P H group). Treatment with AC→PH resulted in a statistically significant improvement in OS compared with AC→P (stratified HR=0.64; 95% CI [0.55, 0.74]; log-rank p-value < 0.0001). At 8 years, the survival rate was estimated to be 86.9% in the AC→PH arm and 79.4% in the AC→P arm, an absolute benefit of 7.4% (95% CI 4.9%, 10.0%).

The final OS results from the joint analysis of studies NSABP B-31 and NCCTG N9831 are summarized in the table below:

Table 9 Final Overall Survival Analysis from the joint analysis of trials NSABP B-31 and NCCTG N9831

Parameter	AC→P (n=2032)	AC→PH (n=2031)	p-value	HR (95% CI)
Death (OS event)				
No. patients with event (%)	418 (20.6%)	289 (14.2%)	< 0.0001	0.64 (0.55, 0.74)

A: doxorubicin; C: cyclophosphamide; P: paclitaxel; H: Herceptin

DFS analysis was also performed at the final analysis of OS from the joint analysis of studies NSABP B-31 and NCCTG N9831. The updated DFS analysis results (stratified HR = 0.61; 95% CI [0.54, 0.69]) showed a similar DFS benefit compared to the definitive primary DFS analysis, despite 24.8% patients in the AC→P arm who crossed over to receive Herceptin. At 8 years, the disease-free survival rate was estimated to be 77.2% (95% CI: 75.4, 79.1) in the AC→PH arm, an absolute benefit of 11.8% compared with the AC→P arm.

The efficacy results from the BCIRG 006 are summarized in the following tables:

Table 10: Overview of Efficacy Analyses BCIRG 006 AC→D versus AC→DT

Parameter	AC→D n = 1073	AC→DH n = 1074	p-value	HR (95% CI)
Disease-free survival (DFS)				
No. patients with event	195	134	<0.0001	0.61 (0.49, 0.77)
Death (OS event)				
No. patients with event	80	49	0.0024	0.58 (0.40,0.83)

AC→D = doxorubicin plus cyclophosphamide, followed by docetaxel; AC→DH = doxorubicin plus cyclophosphamide, followed by docetaxel plus trastuzumab; CI = confidence interval

Table 11: Overview of Efficacy Analyses BCIRG 006 AC→D versus DCarbH

Parameter	AC→D n = 1073	DCarbH n = 1075	p-value	HR (95% CI)
Disease-free survival (DFS)				
No. patients with event	195	145	0.0003	0.67 (0.54, 0.83)

Parameter	AC→D n = 1073	DCarbH n = 1075	p-value	HR (95% CI)
<u>Death (OS event)</u>				
No. patients with event	80	56	0.00182	0.66 (0.47,0.93)

AC→D = doxorubicin plus cyclophosphamide, followed by docetaxel; DCarbH = docetaxel, carboplatin and trastuzumab; CI = confidence interval

Metastatic Breast Cancer

Trastuzumab monotherapy or in combination with chemotherapy

Trastuzumab monotherapy has been used in clinical trials for patients with metastatic breast cancer who have tumours that overexpress HER2 and who have failed one or more chemotherapy regimens for their metastatic disease.

Trastuzumab has also been used in clinical trials in combination with paclitaxel or an anthracycline (doxorubicin or epirubicin) plus cyclophosphamide (AC) as first line therapy for patients with metastatic breast cancer who have tumours that overexpress HER2.

Patients who had previously received anthracycline-based adjuvant chemotherapy were treated with paclitaxel (175 mg/m² infused over 3 hours) with or without trastuzumab. Patients could be treated with trastuzumab until progression of disease.

Trastuzumab monotherapy, when used as second- or third-line treatment of women with metastatic breast cancer which overexpresses HER2, results in an overall tumour response rate of 15% and a median survival of 13 months.

The use of trastuzumab in combination with paclitaxel as first-line treatment of women with metastatic breast cancer that overexpresses HER2 significantly prolongs the median time to disease progression, compared with patients treated with paclitaxel alone. The increase in median time to disease progression for patients treated with paclitaxel is 3.9 months (6.9 months vs. 3.0 months). Tumour response and one year survival rate are also increased for trastuzumab in combination with paclitaxel versus paclitaxel alone.

Trastuzumab has also been studied in a randomised, controlled trial, in combination with docetaxel, as first-line treatment of women with metastatic breast cancer. The combination of trastuzumab and docetaxel significantly increased response rate (61% vs. 34%) and prolonged the median time to disease progression, (by 5.6 months) compared with patients treated with docetaxel alone. Median survival was also significantly increased in patients receiving the combination, compared with those receiving docetaxel alone (31.2 months vs. 22.7 months).

Trastuzumab in combination with anastrozole

The TAnDEM trial was a multi-centre, randomised, open-label, phase III study comparing trastuzumab + anastrozole with anastrozole alone for the first-line treatment of metastatic breast cancer in HER2 overexpressing, hormone-receptor (i.e. oestrogen-receptor (ER) and/or progesterone-receptor (PR)) positive post-menopausal patients. Two hundred and seven patients were randomised to receive oral anastrozole (1 mg/day) with or without trastuzumab (4 mg/kg loading dose, followed by 2 mg/kg weekly). Patients who had previously received trastuzumab were excluded from this trial.

Median progression free survival was doubled in the trastuzumab + anastrozole arm compared to the anastrozole alone arm (4.8 months vs. 2.4 months; $p = 0.0016$). For the other parameters, the improvements seen for trastuzumab + anastrozole were; overall response (16.5% vs. 6.7%); clinical benefit rate (42.7% vs. 27.9%); time to progression (4.8 months vs. 2.4 months). For time to response and duration of response no difference could be recorded between the arms. There was no significant difference in overall survival, however, more than half of the patients in the anastrozole alone arm crossed over to a trastuzumab-containing regimen after progression of disease.

Advanced Gastric Cancer

Study BO18255 (ToGA) was a randomised, open-label, multicentre phase III study investigating trastuzumab in combination with a fluoropyrimidine and cisplatin (FP) versus chemotherapy alone as first-line therapy in patients with HER2 positive, inoperable, locally advanced or recurrent and/or metastatic adenocarcinoma of the stomach or gastro-oesophageal junction.

Patients were eligible if they had 3+ levels of HER2 overexpression based on a 0 - 3+ scale by IHC assessment of tumour tissue and/or those whose tumours had HER2 gene amplification as determined by a FISH test (see Dosage and Administration, Detection of HER2 Overexpression or Gene Amplification).

After satisfying the screening eligibility criteria, including assessment of HER2 status, patients were randomly assigned (1:1) to receive either trastuzumab (8 mg/kg loading dose, followed by 6 mg/kg every 3 weeks) + fluoropyrimidine/cisplatin (FP+H) or FP alone. The chemotherapy regimen was chosen between 5-FU/cisplatin and capecitabine/cisplatin at the investigator's discretion and could be determined on an individual patient basis.

The efficacy results from ToGA are summarised in Table 11. The primary endpoint was overall survival, defined as the time from the date of randomization to the date of death from any cause. At the time of analysis a total of 349 randomised patients had died: 182 patients (62.8%) in the control arm and 167 patients (56.8%) in the treatment arm. The majority of the deaths were due to events related to the underlying cancer.

Overall survival was significantly improved in the FP + H arm compared to the FP arm ($p = 0.0046$, log-rank test). The median survival time was 11.1 months with FP and 13.8 months with FP + H. The risk of death was decreased by 26% (HR = 0.74; 95% CI 0.60 - 0.91) for patients in the FP + H arm compared to the FP arm.

Post-hoc subgroup analyses indicate that targeting tumours with higher levels of HER2 protein (IHC 2+/FISH+ and IHC 3+/regardless of FISH status) results in a greater treatment effect. The median overall survival for the high HER2 expressing group was 11.8 months versus 16 months, HR = 0.65 (95% CI 0.51 - 0.83) and the median progression free survival was 5.5 months vs. 7.6 months, HR = 0.64 (95% CI 0.51 - 0.79).

Table 12: Summary of Efficacy from Study BO18255

Trastuzumab dosage regimen	Every 3 weeks			
Chemotherapy regimens (FP)	<ul style="list-style-type: none"> • Capecitabine: 1000 mg/m² orally twice daily for 14 days every 3 weeks for 6 cycles (Days 1 to 15 of each cycle). • 5-FU: 800 mg/m²/day as a continuous IV infusion over 5 days, given every 3 weeks for 6 cycles (Days 1 to 5 of each cycle). The 5-FU infusion could be started at the same time as the cisplatin infusion on Day 1. • Cisplatin: 80 mg/m² every 3 weeks for 6 cycles (on Day 1 of each cycle) as a 2h IV infusion with hydration and premedication (steroids and anti-emetics). 			
Efficacy Parameters	FP n = 290	FP+H n = 294	HR (95% CI)	p-value
Overall Survival, Median months	11.1	13.8	0.74 (0.60-0.91)	0.0046
Progression-Free Survival, Median months	5.5	6.7	0.71 (0.59-0.85)	0.0002
Time to Disease Progression, Median months	5.6	7.1	0.70 (0.58-0.85)	0.0003
Overall Response Rate, %	34.5	47.3	1.70 ^a (1.22, 2.38)	0.0017
Duration of Response, Median months	4.8	6.9	0.54 (0.40-0.73)	<0.0001

FP: Fluoropyrimidine/cisplatin; FP+H: Fluoropyrimidine/cisplatin + trastuzumab; ^a Odds ratio

Progression-free-survival: time between day of randomization and first documentation of progressive disease (PD) or date of death, whichever occurred first. *Time to disease progression*: time between randomization and first occurrence of PD. *Overall response*: occurrence of either a confirmed complete (CR) or a partial (PR) best overall response as determined by RECIST criteria from confirmed radiographic evaluations of target and non-target lesions. *Duration of response*: time from when response (CR or PR) was first documented to the first documented disease progression. This was only calculated for patients who had a best overall response of CR or PR.

Immunogenicity

Human anti-trastuzumab antibodies were detected in 1 of 903 patients, who had no allergic manifestations.

Pharmacokinetic Properties

The pharmacokinetics of trastuzumab has been studied in patients with breast cancer (metastatic and early breast cancer) and advanced gastric cancer. Formal drug-drug interaction studies have not been performed for trastuzumab.

Breast cancer

Short duration intravenous infusions of 10, 50, 100, 250 and 500 mg trastuzumab once weekly in patients demonstrated dose-dependent pharmacokinetics. Mean half-life increased and clearance decreased with increased dose level.

Steady State Pharmacokinetics in Breast Cancer

A population pharmacokinetic method, using data from Phase I, Phase II and pivotal Phase III studies, was used to estimate the steady state pharmacokinetics in metastatic breast cancer patients. For a typical patient (body weight 68 kg) the clearance of trastuzumab was 0.241 L/day and volume of distribution of the central (V_c) and peripheral (V_p) compartments were 3.02 L, and 2.68 L respectively, with a corresponding elimination half-life ranging from approximately 28-38 days. These indicate steady state pharmacokinetics should therefore be reached by approximately 27 weeks, with mean predicted AUC at steady state (over a three week period) of 1822 mg•day/L with weekly dosing and 1912 mg•day/L with 3-weekly (once every three weeks) dosing. The estimated mean peak and trough concentrations were 113 mg/L and 69.9 mg/L (weekly) and 184 mg/L and 52.9 mg/L (3-weekly) respectively. Comparable steady state mean C_{max} of 225 µg/mL and mean C_{min} of 68.9 µg/mL a day 21 of cycle 18, the last cycle of 1 year treatment, have been reported in localised breast cancer patients administered trastuzumab 3-weekly.

It is expected that serum trastuzumab levels will fall to less than 5% of the trough levels at steady state approximately 27 weeks (190 days or 5 elimination half-lives) after a dose discontinuation.

Gastric cancer

Short duration IV infusions of 8 mg/kg followed by 6 mg/kg trastuzumab every 3 weeks in patients with advanced gastric cancer demonstrated concentration-dependent clearance comprised of predominantly linear and non-linear components. At very low serum concentrations (below 10 µg/mL) non-linear clearance comprises nearly all of the total clearance (7-fold higher than linear clearance). At higher concentrations (25 µg/mL) the non-linear clearance component decreases to approximately one-half of total clearance, whilst at higher concentrations (> 75 µg/L) clearance becomes predominantly linear.

Steady State Pharmacokinetics in Advanced Gastric Cancer

A two compartment population pharmacokinetic method, using data from the Phase III study BO18255 (ToGA) was used to estimate the steady state pharmacokinetics in patients with advanced gastric cancer administered trastuzumab 3-weekly at a loading dose of 8 mg/kg followed by a 3-weekly maintenance dose of 6 mg/kg. At high serum concentrations, total clearance is dominated by linear clearance and the half-life is approximately 26 days. Steady state C_{min} is reached in approximately 168 days, due to the non-linear clearance component. The mean predicted steady-state AUC value (over a period of 3 weeks at steady state) is approximately 1213 mg•day/L, and the mean steady-state C_{max} and C_{min} values are approximately 132 mg/L and 27.6 mg/L respectively. It is expected that serum trastuzumab levels will fall to less than 5% of the trough levels at steady state, approximately 19 weeks after a dose discontinuation.

There are no data on the level of circulating extracellular domain of the HER2 receptor (shed antigen) in the serum of gastric cancer patients.

Special Populations

Detailed pharmacokinetic studies in the elderly and those with renal or hepatic impairment have not been carried out. The data from Study H0649g suggest that the disposition of trastuzumab is not altered by patient characteristics such as age or serum creatinine. The population pharmacokinetic

analysis also shows that the estimated creatinine clearance (Cockcroft and Gault) does not correlate with the pharmacokinetics of trastuzumab.

Elderly

Age has been shown to have no effect on the disposition of trastuzumab (see Dosage and Administration).

Preclinical Safety

Teratogenicity

Reproduction studies have been conducted in cynomolgus monkeys at doses up to 25 times that of the weekly human maintenance dose of 2 mg/kg trastuzumab and have revealed no evidence of impaired fertility or harm to the foetus. However, when assessing the risk of reproductive toxicity to humans, it is also important to consider the significance of the rodent form of the HER2 receptor in normal embryonic development and the embryonic death in mutant mice lacking this receptor. Placental transfer of trastuzumab during the early (days 20 - 50 of gestation) and late (days 120 - 150 of gestation) foetal development period was observed.

Other

Lactation

A study conducted in lactating cynomolgus monkeys at doses 25 times that of the weekly human maintenance dose of 2 mg/kg trastuzumab demonstrated that trastuzumab is secreted in the milk. The presence of trastuzumab in the serum of infant monkeys was not associated with any adverse effects on their growth or development from birth to 1 month of age.

Pharmaceutical Particulars

Storage

Vials

Store vials at 2°C – 8°C.

This medicine should not be used after the expiry date shown on the pack.

Shelf-life of the reconstituted solution

Herclon 440 mg vial

Reconstituted solutions made with Bacteriostatic Water for Injection for the 440 mg vial of Herclon, as supplied, are stable for **28 days when stored refrigerated at 2 °C – 8 °C**. The reconstituted solution contains preservative and is therefore suitable for multiple use. Any remaining reconstituted solution

should be discarded after 28 days. If sterile water is used to reconstitute the 440 mg vial, the solution is stable for only **24 hours**, and must be discarded thereafter.

Do not freeze the reconstituted solution.

Herclon 150 mg vial

The 150 mg vials are reconstituted with sterile water for injection and are for single use only.

The reconstituted product is physically and chemically stable for **48 hours at 2 °C – 8 °C** (refrigerate, do not freeze) after dissolving with water for injections.

From a microbiological point of view, the reconstituted solution should be further diluted immediately. If not further diluted immediately, in-use storage times and conditions prior to dilution are the responsibility of the user. The shelf-life of the reconstituted solution is **24 hours at 2 °C – 8 °C**, unless reconstitution has taken place in controlled and validated aseptic conditions in which case the shelf-life is **48 hours at 2 °C – 8 °C**.

Shelf-life of the solution for infusion containing the reconstituted product

The infusion solution (0.9% sodium chloride infusion solution) containing the reconstituted product is physically and chemically stable for **24 hours (do not store above 30 °C)**.

From a microbiological point of view, the Herclon infusion solution should be used immediately. If not used immediately, in-use storage times and conditions prior to use are the responsibility of the user and would normally not be longer than **24 hours at 2 °C to 8 °C**, unless reconstitution and dilution have taken place in controlled and validated aseptic conditions.

Special Instructions for Use, Handling and Disposal

Appropriate aseptic technique should be used.

The 440 mg vial of Herclon is reconstituted with 20 mL of Bacteriostatic Water for Injection, containing 1.1% benzyl alcohol, as supplied. This yields a solution for multiple use, containing 21 mg/mL trastuzumab, at a pH of approximately 6.0. Use of other reconstitution solvents should be avoided.

The 150 mg vial of Herclon is reconstituted with 7.2 mL of sterile water for injection.

Herclon should be carefully handled during reconstitution. Causing excessive foaming during reconstitution or shaking the reconstituted Herclon may result in problems with the amount of Herclon that can be withdrawn from the vial.

Instructions for reconstitution – 440 mg vial:

1. Using a sterile syringe, slowly inject 20 mL of sterile Bacteriostatic Water for Injection into the vial containing the lyophilised Herclon, directing the stream into the lyophilised cake.
2. Swirl vial gently to aid reconstitution. **DO NOT SHAKE!**

Instructions for reconstitution – 150 mg vial:

- 1) Using a sterile syringe, slowly inject 7.2 mL of sterile water for injection into the vial containing the lyophilised Herclon, directing the stream into the lyophilised cake.
- 2) Swirl vial gently to aid reconstitution. DO NOT SHAKE!

Slight foaming of the product upon reconstitution is not unusual. Allow the vial to stand undisturbed for approximately 5 minutes. The reconstituted Herclon results in a colourless to pale yellow transparent solution and should be essentially free of visible particles.

Instructions for dilution

Weekly schedule

Determine the volume of the reconstituted solution required based on a loading dose of 4 mg trastuzumab/kg body weight, or a maintenance dose of 2 mg trastuzumab/kg body weight:

$$\text{Volume (mL)} = \frac{\text{Body weight (kg)} \times \text{dose (4 mg/kg for loading or 2 mg/kg for maintenance)}}{21 \text{ (mg/mL, concentration of reconstituted solution)}}$$

Three-weekly schedule

Determine the volume of the reconstituted solution required based on a loading dose of 8 mg trastuzumab/kg body weight, or a maintenance dose of 6 mg trastuzumab/kg body weight:

$$\text{Volume (mL)} = \frac{\text{Body weight (kg)} \times \text{dose (8 mg/kg for loading or 6 mg/kg for maintenance)}}{21 \text{ (mg/mL, concentration of reconstituted solution)}}$$

Preparation and Stability of the Admixture

The appropriate amount of reconstituted solution should be withdrawn from the vial and added to an infusion bag containing 250 mL of 0.9% sodium chloride. Dextrose (5%) solution should not be used (see Incompatibilities). The bag should be gently inverted to mix the solution in order to avoid foaming. Parenteral medicines should be inspected visually for particulates and discoloration prior to administration. Once the infusion is prepared it should be administered immediately (see Storage).

Incompatibilities

No incompatibilities between Herclon and polyvinylchloride, polyethylene or polypropylene bags have been observed.

Dextrose (5%) solution should not be used since it causes aggregation of the protein.

Herclon should not be mixed or diluted with other medicines.

Disposal

The release of medicines into the environment should be minimised. Medicines should not be disposed of via wastewater and disposal through household waste should be avoided. Unused or expired medicine should be returned to a pharmacy for disposal.

Medicine Classification

Prescription medicine.

Packs

Herclon 150 mg single-dose vials: each pack contains one vial of Herclon (150 mg trastuzumab).

Herclon 440 mg multi-dose vials: each pack contains one vial of Herclon (440 mg trastuzumab) and one 20 mL vial of Bacteriostatic Water for Injection.

Name and Address

Roche Products (New Zealand) Limited
PO Box 109113 Newmarket
Auckland 1149
NEW ZEALAND

Medical enquiries: 0800 656 464

Date of Preparation

25 June 2014