

Herzuma[®]
(trastuzumab-pkrb)
For Injection • 420 mg/vial • 150 mg/vial

UNDERSTANDING
TREATMENT

WITH HERZUMA

A GUIDE FOR GETTING STARTED

Please see Important Safety Information throughout and full [Prescribing Information](#), including BOXED WARNINGS, for HERZUMA to discuss with your doctor.

teva

GET TO KNOW

HERZUMA

BEING DIAGNOSED WITH CANCER CAN BE OVERWHELMING. LEARNING MORE ABOUT WHAT CAN BE DONE TO TREAT IT MAY HELP YOU BE PREPARED.

Whether you are getting ready to start treatment for HER2+ breast cancer with HERZUMA or you are considering it, this guide can help you:

GET THE FACTS
about HERZUMA and how it may help

LEARN
what to expect from treatment with HERZUMA

UNDERSTAND
why your doctor may prescribe HERZUMA

FIND OUT
about helpful resources

HER2+ = Human Epidermal growth factor Receptor 2-positive.

APPROVED USES

Patients are selected for therapy with HERZUMA based on an FDA-approved test.

Adjuvant Breast Cancer

HERZUMA is a prescription medicine used for the treatment of early-stage breast cancer that is Human Epidermal growth factor Receptor 2-positive (HER2+) and has spread into the lymph nodes, or is HER2-positive and has not spread into the lymph nodes. If it has not spread into the lymph nodes, the cancer needs to be estrogen receptor/progesterone receptor (ER/PR)-negative or have one high-risk feature. HERZUMA can be used in several different ways: • As part of a treatment course including the chemotherapy drugs doxorubicin, cyclophosphamide, and either paclitaxel or docetaxel. This treatment course is known as “AC → TH” • With the chemotherapy drugs docetaxel and carboplatin. This treatment course is known as “TCH” • Alone after treatment with multiple other therapies, including an anthracycline (doxorubicin)-based therapy (a type of chemotherapy)

See Approved Uses statement continued on next page.

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.

WHAT IS HERZUMA?

HERZUMA is a prescription drug used in adults to treat HER2+ breast cancer. HERZUMA is not chemotherapy, though it is sometimes used with chemotherapy. HERZUMA is a type of treatment called an antibody therapy. This means HERZUMA is able to treat HER2+ breast cancer cells, which can multiply rapidly.



HERZUMA MAY WORK TO TREAT HER2+ BREAST CANCER IN THE FOLLOWING WAYS:

- By binding to the HER2 receptor on cancer cells
- By telling the immune system it's okay to destroy cancer cells
- By destroying cancer cells on its own

HERZUMA may also harm some healthy cells in the body. Talk to your doctor about any concerns you may have.

BIOLOGICS AND BIOSIMILARS

Biologics are complex drugs produced from living cells.

Biosimilars are FDA-approved biological products that are highly similar to and have no clinically meaningful differences from existing FDA-approved biologic drugs.

HERZUMA is a biosimilar to Herceptin® (trastuzumab) for HER2+ breast cancer.

APPROVED USES (continued)

Patients are selected for therapy with HERZUMA based on an FDA-approved test.

Metastatic Breast Cancer

HERZUMA has 2 approved uses in metastatic breast cancer: • HERZUMA in combination with the chemotherapy drug paclitaxel is approved for the first-line treatment of Human Epidermal growth factor Receptor 2-positive (HER2+) metastatic breast cancer • HERZUMA alone is approved for the treatment of HER2-positive breast cancer in patients who have received one or more chemotherapy courses for metastatic disease

IMPORTANT SAFETY INFORMATION

HERZUMA can cause serious side effects, including:

- **HEART PROBLEMS:** Trastuzumab products can cause heart problems—such as congestive heart failure or reduced heart function—with or without symptoms. The risk for and seriousness of these heart problems were highest in people who received both a trastuzumab product and a certain type of chemotherapy (anthracycline). In a study of adjuvant (early) breast cancer, one patient died of significantly weakened heart muscle. Your healthcare provider will check for signs of heart problems before, during, and after treatment with HERZUMA

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.



ABOUT HER2+ BREAST CANCER

TREATMENT GOALS

YOUR DOCTOR MAY PRESCRIBE
HERZUMA TO **TREAT YOUR**
HER2+ BREAST CANCER

APPROVED USES

HERZUMA is used for the treatment of adjuvant breast cancer and metastatic breast cancer in patients selected for therapy based on an FDA-approved test.

See complete indication on pages 2-3

IMPORTANT SAFETY INFORMATION (continued)

HERZUMA can cause serious side effects, including:

- **INFUSION REACTIONS:** Infusion reactions, sometimes serious or fatal, have occurred. Symptoms may include: fever, chills, feeling sick to your stomach (nausea), throwing up (vomiting), pain (in some cases at tumor site), headache, dizziness, and shortness of breath. These symptoms usually happen within 24 hours after receiving HERZUMA

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.



WHAT IS HER2+ BREAST CANCER?

Breast cancer happens when cells in the breast grow out of control. These cells often form a tumor that can be felt as a lump. The tumor can be benign (not cancerous) or malignant (cancerous).

When you are diagnosed with breast cancer, your doctor will order tests to learn more about how the cancer should be treated. One of those tests will assess the HER2 status of your tumor.

HER2 is a protein on the outside of all cells in the breast. HER2+ cancer cells have higher than normal levels of HER2. HER2+ cancer cells tend to grow and spread faster (metastasize) than other types of breast cancer.

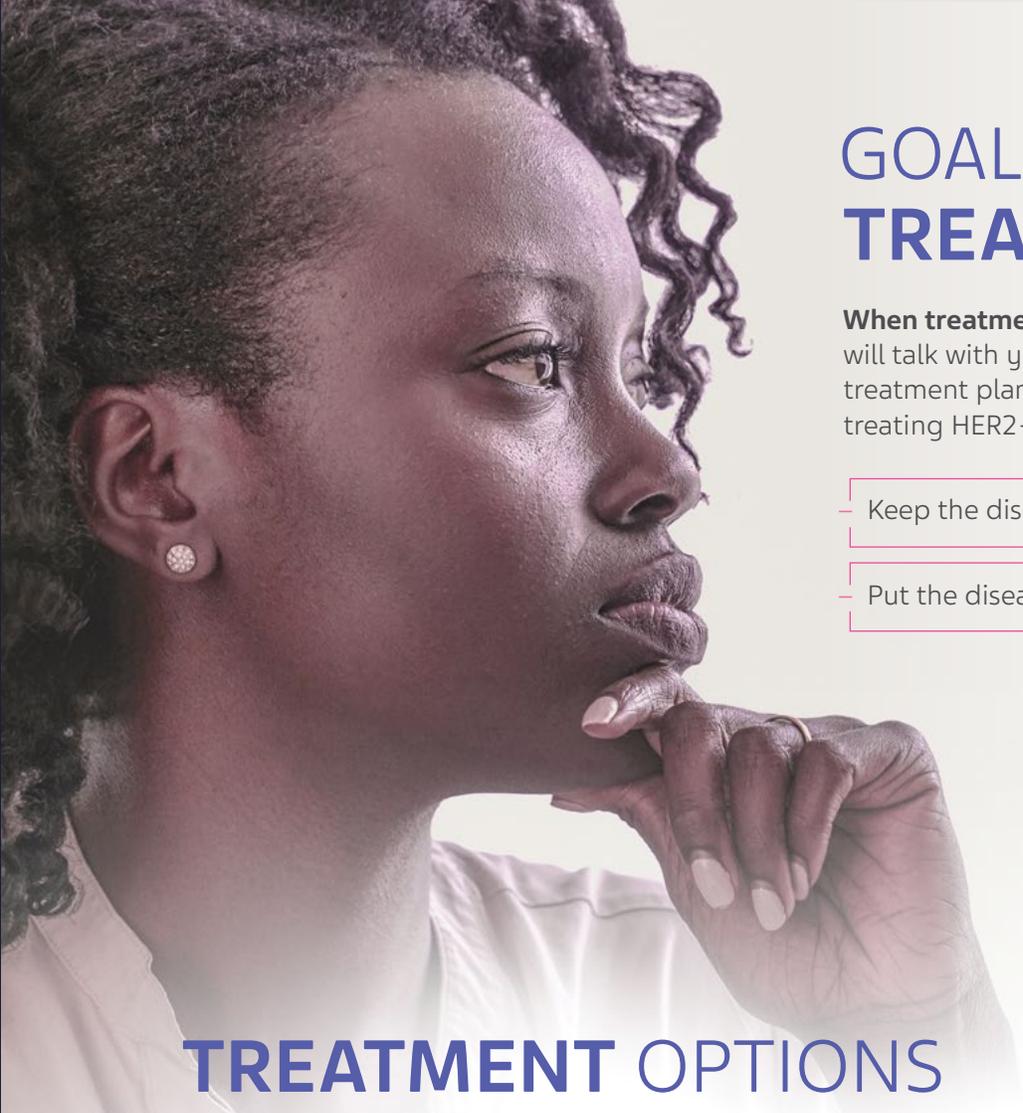
**AN ESTIMATED 1 IN 5 WOMEN WITH
BREAST CANCER WILL BE HER2+**

SYMPTOMS OF HER2+ BREAST CANCER

A new lump or mass on the breast is the most common symptom of breast cancer. They can be hard, irregular, and painless, or soft, round, and painful.

Other possible symptoms include:

- Swelling in the breast
- Skin changes on the breast or nipple including irritation, dimples, redness, scaliness, or thickening
- Painful breast or nipples
- Nipple turning inward
- Fluid discharge from the nipple
- Lump or swelling around lymph nodes under the arm or around the collar bone



GOALS FOR TREATMENT

When treatment is needed, your healthcare team will talk with you about options and come up with a treatment plan that's right for you. Some goals when treating HER2+ breast cancer may be to help:

- Keep the disease from advancing

- Put the disease into remission

TREATMENT OPTIONS

HER2+ BREAST CANCER

Treatments for breast cancer can include surgery, radiation, chemotherapy, hormone therapy, or immunotherapy. However, for HER2+ breast cancer, a therapy called targeted therapy is often recommended. Treatments for HER2+ breast cancer target the HER2+ protein.

THE APPROPRIATE TREATMENT FOR
YOU DEPENDS ON:

HOW FAST THE CANCER IS GROWING

THE STAGE OF THE CANCER

YOUR PERSONAL CHARACTERISTICS
SUCH AS AGE AND OVERALL HEALTH

PLEASE TALK WITH YOUR DOCTOR ABOUT THESE
AND OTHER TREATMENT OPTIONS

STARTING HERZUMA

STARTING TREATMENT FOR HER2+ BREAST CANCER CAN LEAVE YOU WONDERING WHAT QUESTIONS TO ASK NEXT. THE INFORMATION IN THIS SECTION MAY HELP YOU GET THE ANSWERS YOU NEED.

APPROVED USES

Patients are selected for therapy with HERZUMA based on an FDA-approved test.

Adjuvant Breast Cancer

HERZUMA is a prescription medicine used for the treatment of early-stage breast cancer that is Human Epidermal growth factor Receptor 2-positive (HER2+) and has spread into the lymph nodes, or is HER2-positive and has not spread into the lymph nodes. If it has not spread into the lymph nodes, the cancer needs to be estrogen receptor/progesterone receptor (ER/PR)-negative or have one high-risk feature. HERZUMA can be used in several different ways:

- As part of a treatment course including the chemotherapy drugs doxorubicin, cyclophosphamide, and either paclitaxel or docetaxel. This treatment course is known as “AC → TH”
- With the chemotherapy drugs docetaxel and carboplatin. This treatment course is known as “TCH”
- Alone after treatment with multiple other therapies, including an anthracycline (doxorubicin)-based therapy (a type of chemotherapy)

See Approved Uses statement continued on next page.

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.



WHAT YOU SHOULD TELL YOUR DOCTOR BEFORE STARTING HERZUMA



BEFORE RECEIVING HERZUMA TELL YOUR DOCTOR IF YOU:

- Have trouble with your heart. HERZUMA can cause heart problems in some patients
- Are pregnant or plan to become pregnant. Trastuzumab products can cause fetal harm when administered to a pregnant woman. Because trastuzumab stays in the body for a long time, you should avoid becoming pregnant for 7 months after discontinuing HERZUMA
- Are taking any other medications, including prescription and over-the-counter medicines, vitamins, and herbal supplements
- Know the medicines you take. Keep a list of them to show to your doctor and pharmacist when you get a new medicine. Do not take any new medicine without speaking with your doctor

APPROVED USES (continued)

Patients are selected for therapy with HERZUMA based on an FDA-approved test.

Metastatic Breast Cancer

HERZUMA has 2 approved uses in metastatic breast cancer:

- HERZUMA in combination with the chemotherapy drug paclitaxel is approved for the first-line treatment of Human Epidermal growth factor Receptor 2-positive (HER2+) metastatic breast cancer
- HERZUMA alone is approved for the treatment of HER2-positive breast cancer in patients who have received one or more chemotherapy courses for metastatic disease

IMPORTANT SAFETY INFORMATION (continued)

HERZUMA can cause serious side effects, including:

- **SEVERE LUNG PROBLEMS:** Trastuzumab product use can result in serious and fatal lung problems. Symptoms may include: severe shortness of breath, fluid in or around the lungs, weakening of the valve between the heart and lungs, not enough oxygen in the body, swelling of the lungs, and scarring of the lungs. Your healthcare provider may check for signs of severe lung problems

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.

WHY IS HEART MONITORING NECESSARY WHILE RECEIVING HERZUMA THERAPY?



Your healthcare team will monitor your heart for potential cardiac issues to catch them early on so they can be properly treated.

Your heart will be monitored at the start of treatment, every 3 months while on HERZUMA therapy, and every 6 months for at least 2 years following completion of HERZUMA as a component of adjuvant therapy. Heart function will be tested with an ECHO or MUGA scan.*†

DURING HERZUMA THERAPY

– Pausing HERZUMA

In case of reduced heart function, your doctor may hold HERZUMA therapy up to 3 times

– Restarting HERZUMA

Your doctor may restart HERZUMA therapy up to 3 times if your heart function improves and returns to normal limits after HERZUMA has been held

– Discontinuing HERZUMA

HERZUMA will be discontinued if you experience congestive heart failure or significantly reduced heart function (when the heart can't keep up with the amount of blood flowing through it)

AFTER HERZUMA THERAPY

Your heart should be monitored every 6 months for at least 2 years

*ECHO scan is an ultrasound image of the heart.

†MUGA scan takes a moving picture of your heart pumping blood following an injection of a nontoxic radioactive substance.

IMPORTANT SAFETY INFORMATION (continued)

HERZUMA can cause serious side effects, including:

- **HARM TO UNBORN BABIES OR BIRTH DEFECTS:** Trastuzumab products may result in birth defects or the death of an unborn baby. Contraception should be used while receiving HERZUMA and for **7 months** after your last dose of HERZUMA. If you are or become pregnant while receiving HERZUMA or **within 7 months** after your last dose of HERZUMA, you should immediately contact your doctor. Patients should contact their healthcare professional with a known or suspected pregnancy

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.

HOW WILL YOU RECEIVE HERZUMA?



- HERZUMA is given by infusion through a needle placed in a vein (intravenous infusion) in your arm, or through a port
- Talk to your doctor to find out whether you will receive HERZUMA in an infusion center or in your doctor's office
- The first dose of HERZUMA takes about 90 minutes. After that, the other doses of HERZUMA only take about 30 minutes

WILL YOU HAVE TO TAKE ANY MEDICINE BEFORE YOU RECEIVE HERZUMA?



- Your healthcare provider may prescribe medicines before each infusion of HERZUMA to reduce infusion side effects such as fever and chills
- Taking the suggested medication before treatment may reduce the chance of having a severe reaction during the first HERZUMA infusion
- Be sure to ask your doctor or nurse about what you should take before HERZUMA treatment

IMPORTANT SAFETY INFORMATION (continued)

HERZUMA can cause serious side effects, including:

- **LOW WHITE BLOOD CELL COUNTS:** Low white blood cell counts (which may be life-threatening) were seen more often in patients receiving trastuzumab plus chemotherapy than in patients receiving chemotherapy alone. Your healthcare provider may check for signs of low white blood cell counts

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.

HOW WILL YOUR DOCTOR CHECK UP ON YOU DURING TREATMENT?



- Your doctor should do blood tests and monitor your heart regularly to check for side effects to HERZUMA
- Before each HERZUMA treatment, your healthcare provider will ask you questions about your general health. Tell your healthcare provider about any new symptoms



Herzuma[®]
(trastuzumab-pkrb)
For Injection • 420 mg/vial • 150 mg/vial



HOW CAN YOU GET READY FOR YOUR HERZUMA INFUSION?

Use this list to help prepare yourself before every infusion:

1 PLAN TRANSPORTATION TO AND FROM YOUR INFUSION APPOINTMENT

You may feel exhausted after your infusion, so having someone else drive you home after treatments is a good idea.

2 BRING SOMETHING TO PASS THE TIME

A day at the clinic can be long. Reading magazines, completing a word search, or enjoying a similar activity can help you occupy the time.

3 BRING FOOD AND BEVERAGES

You may be at the clinic for most of the day, so pack some snacks or a light meal and bring a water bottle.

4 TELL YOUR DOCTOR OR NURSE ABOUT MEDICINES YOU ARE TAKING

If you take any other medicines, tell your doctor or nurse. Do not start any new medications without talking to your doctor. Your doctor may give you special instructions for your infusion day.

5 SPEAK UP

Tell your doctor or nurse about any concerns you have.

QUESTIONS FOR YOUR DOCTOR

IT CAN BE DIFFICULT TO KNOW THE RIGHT QUESTIONS to ask your doctor after being diagnosed with HER2+ breast cancer or being prescribed HERZUMA. While you may have already talked with your doctor about some of these topics, the questions that follow can be a good way to start or continue those conversations.

IMPORTANT SAFETY INFORMATION (continued)

Your healthcare provider will stop treatment with HERZUMA if you have severe, serious, or life-threatening side effects.

The most common side effects for patients receiving HERZUMA for breast cancer include: headache, diarrhea, feeling sick to your stomach (nausea), chills, fever, infection, weakened heart muscle, unable to sleep, cough, and rash.

These are not all of the possible side effects with HERZUMA.

Please see the HERZUMA full Prescribing Information, including **BOXED WARNINGS**.

Talk to your doctor about any side effects you may experience.

You may report side effects to FDA at 1-800-FDA-1088. You may also report side effects to Teva at 1-888-483-8279.

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.

EXAMPLE QUESTIONS TO ASK YOUR DOCTOR



QUESTIONS ABOUT HER2+ BREAST CANCER:

WHAT STAGE OF HER2+ BREAST CANCER DO I HAVE?

WHAT ARE MY TREATMENT OPTIONS?

WHAT ARE THE POSSIBLE RISKS AND SIDE EFFECTS OF
HER2+ BREAST CANCER TREATMENT OPTIONS?

WHAT IS MY TREATMENT PLAN?

WHERE CAN I GET MORE INFORMATION ABOUT MY
TREATMENT OPTIONS AND HER2+ BREAST CANCER?

QUESTIONS ABOUT HERZUMA:

WHY HAVE I BEEN PRESCRIBED HERZUMA?

WHAT ARE SIDE EFFECTS OF HERZUMA?

WHAT ARE POSSIBLE RISKS AND BENEFITS OF
HERZUMA?

HOW LONG WILL I NEED TO TAKE HERZUMA?

WHICH PROGRAMS CAN HELP ME SAVE ON THE
COST OF HERZUMA?

APPROVED USES

Patients are selected for therapy with HERZUMA based on an FDA-approved test.

Adjuvant Breast Cancer

HERZUMA is a prescription medicine used for the treatment of early-stage breast cancer that is Human Epidermal growth factor Receptor 2-positive (HER2+) and has spread into the lymph nodes, or is HER2-positive and has not spread into the lymph nodes. If it has not spread into the lymph nodes, the cancer needs to be estrogen receptor/progesterone receptor (ER/PR)-negative or have one high-risk feature. HERZUMA can be used in several different ways:

- As part of a treatment course including the chemotherapy drugs doxorubicin, cyclophosphamide, and either paclitaxel or docetaxel. This treatment course is known as “AC → TH”
- With the chemotherapy drugs docetaxel and carboplatin. This treatment course is known as “TCH”
- Alone after treatment with multiple other therapies, including an anthracycline (doxorubicin)-based therapy (a type of chemotherapy)

See Approved Uses statement continued on next page.

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.

SUPPORT AND SAVINGS

FOR HERZUMA

OUR TEVA SHARED SOLUTIONS[®]
PROGRAM CAN HELP YOU
GET STARTED

APPROVED USES (continued)

Patients are selected for therapy with HERZUMA based on an FDA-approved test.

Metastatic Breast Cancer

HERZUMA has 2 approved uses in metastatic breast cancer:

- HERZUMA in combination with the chemotherapy drug paclitaxel is approved for the first-line treatment of Human Epidermal growth factor Receptor 2-positive (HER2+) metastatic breast cancer
- HERZUMA alone is approved for the treatment of HER2-positive breast cancer in patients who have received one or more chemotherapy courses for metastatic disease

IMPORTANT SAFETY INFORMATION

HERZUMA can cause serious side effects, including:

- **HEART PROBLEMS:** Trastuzumab products can cause heart problems—such as congestive heart failure or reduced heart function—with or without symptoms. The risk for and seriousness of these heart problems were highest in people who received both a trastuzumab product and a certain type of chemotherapy (anthracycline). In a study of adjuvant (early) breast cancer, one patient died of significantly weakened heart muscle. Your healthcare provider will check for signs of heart problems before, during, and after treatment with HERZUMA
- **INFUSION REACTIONS:** Infusion reactions, sometimes serious or fatal, have occurred. Symptoms may include: fever, chills, feeling sick to your stomach (nausea), throwing up (vomiting), pain (in some cases at tumor site), headache, dizziness, and shortness of breath. These symptoms usually happen within 24 hours after receiving HERZUMA

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.



Herzuma[®]
(trastuzumab-pkrb)
For Injection • 420 mg/vial • 150 mg/vial

HAVING AN ILLNESS IS HARD.

Figuring out insurance benefits and financial assistance can make it harder. With our Teva **Shared Solutions**[®], we can help you understand your insurance benefits and may be able to help you find financial assistance for your treatment.

IMPORTANT SAFETY INFORMATION (continued)

HERZUMA can cause serious side effects, including:

- **SEVERE LUNG PROBLEMS:** Trastuzumab product use can result in serious and fatal lung problems. Symptoms may include: severe shortness of breath, fluid in or around the lungs, weakening of the valve between the heart and lungs, not enough oxygen in the body, swelling of the lungs, and scarring of the lungs. Your healthcare provider may check for signs of severe lung problems
- **HARM TO UNBORN BABIES OR BIRTH DEFECTS:** Trastuzumab products may result in birth defects or the death of an unborn baby. Contraception should be used while receiving HERZUMA and for **7 months** after your last dose of HERZUMA. If you are or become pregnant while receiving HERZUMA or **within 7 months** after your last dose of HERZUMA, you should immediately contact your doctor. Patients should contact their healthcare professional with a known or suspected pregnancy
- **LOW WHITE BLOOD CELL COUNTS:** Low white blood cell counts (which may be life-threatening) were seen more often in patients receiving trastuzumab plus chemotherapy than in patients receiving chemotherapy alone. Your healthcare provider may check for signs of low white blood cell counts

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.

teva

teva | Shared Solutions[®] for Biosimilars

- Benefits verification and coverage determination

- Support for precertification and prior authorization

Eligible patients
pay as little as



TERMS AND CONDITIONS

The HERZUMA Cost Support Program helps commercially insured patients in the United States (including the United States territories) who are prescribed HERZUMA pay for their eligible out-of-pocket costs for the drug. The Program does not cover the costs of physician office visits or evaluations, blood work or other testing, or transportation. See complete Terms and Conditions at [HERZUMASavingsOffer.com](https://www.hurzumasavingsoffer.com). Eligible Patients must have commercial insurance coverage for HERZUMA. Uninsured and cash-paying patients are NOT eligible for this Program. Patients enrolled in any state or federally funded healthcare program are NOT eligible for this Program, nor are patients with commercial insurance coverage that does not provide coverage for HERZUMA. Call for more information: **1-888-587-3263**.

SEE IF YOU ARE ELIGIBLE TO SAVE ON
HERZUMA, AND GET INSURANCE AND
REIMBURSEMENT INFORMATION.

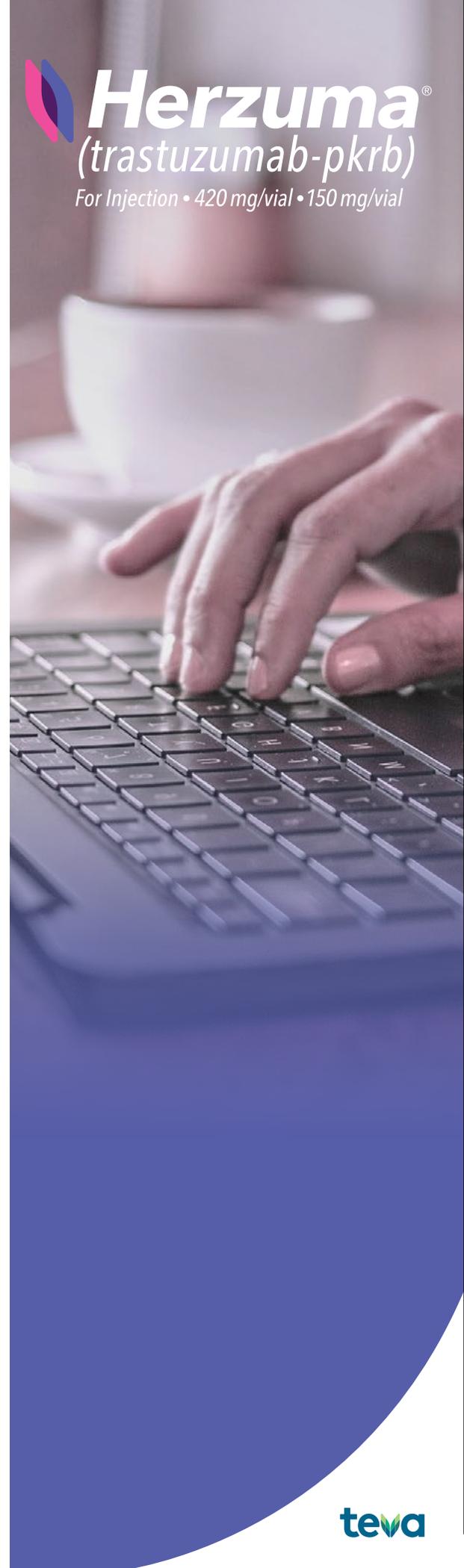
To learn more

CALL

1-888-587-3263

Monday–Friday, 9 AM–7 PM (EST)

 **Herzuma[®]**
(trastuzumab-pkrb)
For Injection • 420 mg/vial • 150 mg/vial



HELPFUL RESOURCES

CANCER ORGANIZATIONS

AMERICAN CANCER SOCIETY

1-800-ACS-2345 (1-800-227-2345)
www.cancer.org

CANCERCARE

1-800-813-HOPE (1-800-813-4673)
www.cancercare.org

NATIONAL CANCER INSTITUTE

1-800-4-CANCER (1-800-422-6237)
www.cancer.gov

BREAST CANCER ORGANIZATIONS

LIVING BEYOND BREAST CANCER

1-855-807-6386
www.lbbc.org

METASTATIC BREAST CANCER NETWORK

1-888-500-370
www.mbcn.org

SUPPORT ORGANIZATIONS

CANCER HOPE NETWORK

1-877-HOPENET (1-877-467-3638)
www.cancerhopenetwork.org

PATIENT ADVOCATE FOUNDATION

1-800-532-5274
www.patientadvocate.org

This is a list of sources that you may find helpful. Please note that this information was accurate at the time of publication, but is subject to change without notice. Ask your healthcare team to recommend additional resources.

IMPORTANT SAFETY INFORMATION (continued)

Your healthcare provider will stop treatment with HERZUMA if you have severe, serious, or life-threatening side effects.

The most common side effects for patients receiving HERZUMA for breast cancer include: headache, diarrhea, feeling sick to your stomach (nausea), chills, fever, infection, weakened heart muscle, unable to sleep, cough, and rash.

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.



Herzuma[®]
(trastuzumab-pkrb)
For Injection • 420 mg/vial • 150 mg/vial

YOUR GUIDE TO TREATMENT WITH HERZUMA

THIS BROCHURE CONTAINS
IMPORTANT INFORMATION FOR
YOUR TREATMENT.

BE SURE TO:

Check out [AboutHerzuma.com](https://www.aboutherzuma.com) for potential savings on HERZUMA

Models appearing in all images within this brochure depict an actor portrayal.

IMPORTANT SAFETY INFORMATION (continued)

These are not all of the possible side effects with HERZUMA.

Please see the HERZUMA full Prescribing Information, including **BOXED WARNINGS**.

Talk to your doctor about any side effects you may experience.

You may report side effects to FDA at 1-800-FDA-1088. You may also report side effects to Teva at 1-888-483-8279.

Please see Important Safety Information throughout and full [Prescribing Information](#), including **BOXED WARNINGS**, for HERZUMA to discuss with your doctor.

teva

HERZUMA[®] is a registered trademark of Celltrion Inc., used under license.
Herceptin[®] is a registered trademark of Genentech USA, Inc.
©2022 Teva Pharmaceuticals USA, Inc. TRA-40203 December 2022