ABOUT YOUR TECENTRIQ INFUSION



You may be receiving TECENTRIQ alone or as part of your treatment plan

HOW TECENTRIQ IS GIVEN

Your healthcare provider will give you TECENTRIQ into your vein through an intravenous (IV) line.

LENGTH OF INFUSION TIME

INITIAL INFUSION: 60 MINUTES FOLLOWING INFUSIONS: 30 MINUTES





The first infusion will be given over 60 minutes. If it's well tolerated, all following TECENTRIQ infusions may be given over 30 minutes. If TECENTRIQ is one part of your treatment, infusion time may be longer.

FLEXIBLE INFUSION SCHEDULE ERY 2 WEEKS EVERY 3 WEEKS EVERY 4 WEEK

EVERY 2 WEEKS







TECENTRIO has flexible dosing options and can be given every 2, 3, or 4 weeks. Depending on your treatment plan, you may need to see your doctor more frequently. Your doctor will decide which treatment schedule is best for you.*

STAYING COMFORTABLE DURING YOUR TREATMENT

- Some people feel cold during their infusion, so consider bringing a blanket
- Ask a companion to keep you company during your infusion
- Some appointments take longer than others, so wear comfortable clothes
- Entertain yourself with a book, tablet, or a podcast on your phone
- Take along some light snacks or a bottle of water
- A pen and treatment journal or notepad can be helpful for jotting down questions or thoughts

IMPORTANT SAFETY INFORMATION

WHAT IS THE MOST IMPORTANT INFORMATION ABOUT TECENTRIQ?

TECENTRIQ can cause your immune system to attack normal organs and tissues in any area of your body and can affect the way they work. These problems can sometimes become severe or life-threatening and can lead to death. You can have more than one of these problems at the same time. These problems may happen anytime during your treatment or even after your treatment has ended.



^{*}TECENTRIQ will be given as 840 mg every 2 weeks, 1200 mg every 3 weeks, or 1680 mg every 4 weeks.

WHAT IS TECENTRIO?

TECENTRIQ is a prescription medicine used to treat:

Adults with a type of lung cancer called non-small cell lung cancer (NSCLC).

- TECENTRIQ may be used alone as a treatment for your lung cancer:
 - to help prevent your lung cancer from coming back after your tumor(s) has been removed by surgery and you have received platinum-based chemotherapy, and
 - you have stage 2 to stage 3A NSCLC (talk to your healthcare provider about what these stages mean), and
 - your cancer tests positive for "PD-L1".
- TECENTRIQ may be used alone as your first treatment when your lung cancer:
 - has spread or grown, and
 - your cancer tests positive for "high PD-L1", and
 - your tumor does not have an abnormal "EGFR" or "ALK" gene.
- TECENTRIQ may be used with the medicines bevacizumab, paclitaxel, and carboplatin as your first treatment when your lung cancer:
 - has spread or grown, and
 - is a type called "non-squamous NSCLC", and
 - your tumor does not have an abnormal "EGFR" or "ALK" gene.
- TECENTRIQ may be used with the medicines paclitaxel protein-bound and carboplatin as your first treatment when your lung cancer:
 - o has spread or grown, and
 - is a type called "non-squamous NSCLC", and
 - your tumor does not have an abnormal "EGFR" or "ALK" gene.
- TECENTRIQ may be used alone when your lung cancer:
 - o has spread or grown, and
 - you have tried chemotherapy that contains platinum, and it did not work or is no longer working.
 - If your tumor has an abnormal "EGFR" or "ALK" gene, you should have also tried an FDA-approved therapy for tumors with these abnormal genes, and it did not work or is no longer working.

Adults with a type of lung cancer called small cell lung cancer (SCLC).

TECENTRIQ may be used with the chemotherapy medicines carboplatin and etoposide as your first treatment when your lung cancer:

• is a type called "extensive-stage small cell lung cancer," which means that it has spread or grown.

Adults with a type of liver cancer called hepatocellular carcinoma (HCC).

TECENTRIQ may be used with the medicine bevacizumab when your liver cancer:

- has spread or cannot be removed by surgery, and
- you have not received other medicines by mouth or injection through your vein (IV) to treat your cancer.

It is not known if TECENTRIQ is safe and effective when used in children for the treatment of NSCLC, SCLC or HCC.

IMPORTANT SAFETY INFORMATION (CONTINUED)

Call or see your healthcare provider right away if you develop any new or worse signs or symptoms, including:

Lung problems

- cough
- shortness of breath
- chest pain

Intestinal problems

- diarrhea (loose stools) or more frequent bowel movements than usual
- stools that are black, tarry, sticky, or have blood or mucus
- severe stomach-area (abdomen) pain or tenderness

Liver problems

- yellowing of your skin or the whites of your eyes
- severe nausea or vomiting
- pain on the right side of your stomach area (abdomen)
- dark urine (tea colored)
- bleeding or bruising more easily than normal

• vour voice gets deeper

dizziness or fainting

changes in mood or

behavior, such as

decreased sex drive,

Hormone gland problems

- headaches that will not go away or unusual headaches
- eye sensitivity to light
- eye problems
- rapid heartbeat
- increased sweating extreme tiredness

Kidney problems

 decrease in your amount of urine

Skin problems

- rash
- itching
- skin blistering or peeling

Genentech

- weight gain or weight loss
- feeling more hungry or thirsty than usual
- urinating more often than usual
- hair loss
- feeling cold
- blood in your urine
- swelling of your ankles

or genital area

- loss of appetite

irritability, or

forgetfulness

constipation

- painful sores or ulcers in mouth or nose, throat,
- fever or flu-like symptoms
- swollen lymph nodes

© 2023 Genentech USA, Inc. All rights reserved. M-US-00013041(v3.0)

Problems can also happen in other organs.

These are not all of the signs and symptoms of immune system problems that can happen with TECENTRIQ. Call or see your healthcare provider right away for any new or worse signs or symptoms, including:

- Chest pain, irregular heartbeat, shortness of breath, or swelling of ankles
- Confusion, sleepiness, memory problems, changes in mood or behavior, stiff neck, balance problems, tingling or numbness of the arms or legs
- Double vision, blurry vision, sensitivity to light, eye pain, changes in eyesight
- Persistent or severe muscle pain or weakness, muscle cramps
- Low red blood cells, bruising

Infusion reactions that can sometimes be severe or life-threatening. Signs and symptoms of infusion reactions may include:

- chills or shaking
- · shortness of breath or wheezing
- feeling like passing out
- fever
- itching or rash flushing dizziness back or neck pain

Complications, including graft-versus-host disease (GVHD), in people who have received a bone marrow (stem cell) transplant that uses donor stem cells (allogeneic). These complications can be serious and can lead to death. These complications may happen if you underwent transplantation either before or after being treated with TECENTRIQ. Your healthcare provider will monitor you for these complications.

Getting medical treatment right away may help keep these problems from becoming more serious. Your healthcare provider will check you for these problems during your treatment with TECENTRIQ. Your healthcare provider may treat you with corticosteroid or hormone replacement medicines. Your healthcare provider may also need to delay or completely stop treatment with TECENTRIQ if you have severe side effects.

Before you receive TECENTRIQ, tell your healthcare provider about all of your medical conditions, including if you:

- have immune system problems such as Crohn's disease, ulcerative colitis, or lupus
- have received an organ transplant
- have received or plan to receive a stem cell transplant that uses donor stem cells (allogeneic)
- have received radiation treatment to your chest area
- have a condition that affects your nervous system, such as myasthenia gravis or Guillain-Barré syndrome
- are pregnant or plan to become pregnant. TECENTRIQ can harm your unborn baby. Tell your healthcare provider right away if you become pregnant or think you may be pregnant during treatment with TECENTRIQ. Females who are able to become pregnant:
- Your healthcare provider should do a pregnancy test before you start treatment with TECENTRIQ.
- You should use an effective method of birth control during your treatment and for at least 5 months after the last dose of TECENTRIQ.
- are breastfeeding or plan to breastfeed. It is not known if TECENTRIQ passes into your breast milk. Do not breastfeed during treatment and for at least 5 months after the last dose of TECENTRIQ.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

The most common side effects of TECENTRIQ when used alone include:

- feeling tired or weak decreased appetite
- nausea cough
- · shortness of breath

The most common side effects of TECENTRIQ when used in lung cancer with other anti-cancer medicines include:

- feeling tired or weak
- diarrhea
- nausea constipation
- decreased appetite

The most common side effects of TECENTRIQ when used in hepatocellular carcinoma (HCC) with bevacizumab include:

 high blood pressure
 feeling tired or weak • too much protein in the urine TECENTRIQ may cause fertility problems in females, which may affect the ability to have children. Talk to your healthcare provider if you have concerns about fertility. These are not all the possible side effects of TECENTRIQ. Ask your healthcare provider or pharmacist for more information about the benefits and side effects of TECENTRIQ. You may report side effects to the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

You may also report side effects to Genentech at 1-888-835-2555. Please see full Prescribing Information and Medication Guide for additional Important Safety Information.

