

Finding the answers you need starts with asking the right questions. This guide is intended to be used as an aid in your next treatment discussion with your oncologist.

ABOUT CAR-T CELL THERAPY

- What is CAR-T cell therapy and how does it work?

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- How does CAR-T therapy work differently from other multiple myeloma treatments, such as stem-cell transplant or chemotherapy?

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- Is CAR-T cell therapy right for me?

ABOUT CARVYKTI

- If I am eligible for CARVYKTI, when may it be an appropriate next step for me?

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- What are the steps in the CARVYKTI treatment process and how long does it take to complete?

What is CARVYKTI[®] (ciltacabtagene autoleucel)?

- CARVYKTI[®] is a treatment used for adult patients who have cancer of the bone marrow called multiple myeloma. It is used when at least one other treatment has not worked or has stopped working
- CARVYKTI[®] is a medicine made from your own white blood cells, which have been changed (genetically modified) to recognize and attack your multiple myeloma cells

IMPORTANT SAFETY INFORMATION

What is the most important information I should know about CARVYKTI[®]?

CARVYKTI[®] may cause side effects that are severe or life-threatening and can lead to death. Call your healthcare provider or get emergency help right away if you get any of the following:

- fever (100.4°F/38°C or higher)
- difficulty breathing
- persistent or severe diarrhea, abdominal pain, and weight loss
- chills or shaking chills
- very low blood pressure
- fast or irregular heartbeat
- dizziness/lightheadedness

Please read full Important Safety Information on pages 5 to 6, and full [Prescribing Information](#), including Boxed Warning, for CARVYKTI.

TREATMENT EXPECTATIONS

- What can I expect from CARVYKTI treatment in the short and long term?

- What outcomes may be associated with receiving CARVYKTI after 1 other treatment has not worked or stopped working?

- What are the side effects of CARVYKTI?

- How long will I be monitored at the Activated Treatment Center for signs and symptoms of potential side effects?

- How long will my care partner(s) and I need to watch for side effects once I go home?

- How long is the entire CARVYKTI treatment process?

IMPORTANT SAFETY INFORMATION (cont'd)

What is the most important information I should know about CARVYKTI[®]? (cont'd)

- effects on your nervous system, some of which can occur days or weeks after you receive the infusion, and may initially be subtle such as:
 - feeling confused, less alert, or disoriented; having difficulty speaking or slurred speech; having difficulty reading, writing, and understanding words; memory loss
 - loss of coordination affecting movement and balance, slower movements, changes in handwriting
 - personality changes, including a reduced ability to express emotions, being less talkative, disinterested in activities, and reduced facial expression
 - tingling, numbness, and pain of hands and feet, difficulty walking, leg and/or arm weakness, and difficulty breathing
 - facial numbness, difficulty moving muscles of face and eyes

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PREPARING FOR TREATMENT

- Where would I receive the one-time infusion of CARVYKTI?

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- Do I need a care partner during and after the CARVYKTI treatment process?

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- Who can be a care partner? What do they need to know about being a care partner?

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- How long would my care partner and I need to stay at or near an Activated Treatment Center, and do I have to come back for a follow-up?

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- Are there any patient support programs available?

IMPORTANT SAFETY INFORMATION (cont'd)

What is the most important information I should know about CARVYKTI[®]? (cont'd)

It is important that you tell your healthcare providers that you have received CARVYKTI[®] and to show them your CARVYKTI[®] Patient Wallet Card. Your healthcare providers may give you other medicines to treat your side effects.

Before you receive CARVYKTI[®], tell your healthcare provider about all your medical conditions, including if you have:

- Current or past neurologic problems (such as seizures, stroke, new or worsening memory loss)
- Lung or breathing problems
- Kidney problems
- Heart problems
- A recent or active infection
- Liver problems
- Low blood counts

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TOPICS FOR A HELPFUL CHAT WITH YOUR DOCTOR ABOUT CARVYKTI (cont'd)



Traveling away from home for medical care can be financially and logistically challenging. Support is available for eligible patients and their care partners.



Patients prescribed CARVYKTI may be eligible for the MyCARVYKTI Patient Support Program sponsored by Johnson & Johnson and Legend Biotech. MyCARVYKTI is a patient support program designed to support eligible patients and their care partner(s) with travel-related expenses during CARVYKTI treatment at an Activated Treatment Center.

The program may offer support with:



Transportation



Lodging



Meals and Out-of-Pocket Travel Expenses

You may receive this support for leukapheresis, infusion, and bridging therapy if administered at the Activated Treatment Center where you are receiving CARVYKTI treatment.* To participate in MyCARVYKTI, you must have been prescribed CARVYKTI and meet distance and income eligibility criteria. Participants must reside in the United States or its territories.

MyCARVYKTI may be an option for you if you do not have health insurance. The program is available to eligible patients regardless of insurance coverage.

To find out if you are eligible, contact the MyCARVYKTI program directly at:



800-559-7875, option 1

Monday through Friday, 8:00 AM – 8:00 PM ET

Additional information available at [MyCARVYKTIPatientSupport.com](https://www.mycarvykti.com/patient-support)

*MyCARVYKTI support is available only when you and your care partner(s) remain at or near the Activated Treatment Center during the above treatment trips.

To view more patient resources, visit [CARVYKTI.com/resources-and-support/](https://www.carvykti.com/resources-and-support/)

IMPORTANT SAFETY INFORMATION (cont'd)

What is the most important information I should know about CARVYKTI®? (cont'd)

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

How will I receive CARVYKTI®?

- CARVYKTI® is made from your own white blood cells, so your blood will be collected by a process called “leukapheresis” (loo-kah-fur-ee-sis). The procedure can take 3 to 6 hours and may need to be repeated.
- Your white blood cells are sent to a manufacturing center to make CARVYKTI®. It takes about 4-5 weeks from the time your cells are received at the manufacturing site and are available to be shipped back to your healthcare provider, but the time may vary.

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- Your white blood cells are sent to a manufacturing center to make CARVYKTI[®]. It takes about 4-5 weeks from the time your cells are received at the manufacturing site and are available to be shipped back to your healthcare provider, but the time may vary.
- While CARVYKTI[®] is being made, you may get other medicines to treat the multiple myeloma. This is so your multiple myeloma does not get worse.

Before you get CARVYKTI[®], your healthcare provider will give you chemotherapy for 3 days to prepare your body.

Thirty to 60 minutes before you are given CARVYKTI[®], you may be given other medicines. These may include:

- medicines for an allergic reaction (antihistamines)
- medicines for fever (such as acetaminophen)

When your CARVYKTI[®] is ready, your healthcare provider will give CARVYKTI[®] to you through a catheter (tube) placed into your vein (intravenous infusion). Your dose of CARVYKTI[®] will be given in one infusion bag. The infusion usually takes approximately 30-60 minutes.

After getting CARVYKTI[®], you will be monitored at the certified healthcare facility where you received your treatment for at least 7 days after the infusion.

IMPORTANT SAFETY INFORMATION (cont'd)

How will I receive CARVYKTI[®]? (cont'd)

You should plan to stay close to the location where you received your treatment for at least 2 weeks. Your healthcare provider will check to see that your treatment is working and help you with any side effects that may occur. You may be hospitalized if you develop serious side effects until your side effects are under control and it is safe for you to leave the hospital.

Your healthcare provider will want to do blood tests to follow your progress. It is important that you have your blood tested. If you miss an appointment, call your healthcare provider as soon as possible to reschedule.

What should I avoid after receiving CARVYKTI[®]?

- Avoid driving for at least 2 weeks after you get CARVYKTI[®]
- You must not be given certain vaccines called live vaccines for some time before and after CARVYKTI[®] treatment. Talk to your healthcare provider if you need to have any vaccinations
- Do not donate blood, organs, tissues, or cells for transplantation

What are the possible or reasonably likely side effects of CARVYKTI[®]?

The most common side effects of CARVYKTI[®] include:

- fever (100.4°F/38°C or higher), chills
- dizziness/lightheadedness
- headache, muscle or joint pain, feeling very tired
- altered mental state, confusion
- infections
- low levels of antibodies (immunoglobulins) in the blood
- cough, being short of breath
- diarrhea, nausea, decreased appetite, constipation
- fast or irregular heartbeat
- problems with blood clotting

In a study comparing CARVYKTI[®] to standard therapy, there was a higher rate of death in the first 10 months in the CARVYKTI[®] arm (14%) compared to the standard therapy arm (12%). The increased rate of deaths occurred before receiving CARVYKTI[®] and after treatment with CARVYKTI[®]. The reasons for death were progression of multiple myeloma and side effects of the treatment.

CARVYKTI[®] can cause a very common side effect called cytokine release syndrome, or CRS, which can be severe or fatal. Symptoms of CRS include fever, difficulty breathing, dizziness or lightheadedness, nausea,

headache, fast heartbeat, low blood pressure, or fatigue. Tell your healthcare provider right away if you develop fever or any of these other symptoms after receiving CARVYKTI[®].

CARVYKTI[®] can increase the risk of life-threatening infections, including COVID-19, that may lead to death. Tell your healthcare provider right away if you develop fever, chills, or any signs or symptoms of an infection.

CARVYKTI[®] can cause various neurologic side effects, some of which may be severe or fatal. Symptoms include but are not limited to confusion, disorientation, loss of consciousness, seizures, difficulty speaking, reading or writing, tremor, slower movements, changes in personality, depression, tingling and numbness of hands and feet, leg and arm weakness, and facial numbness.

CARVYKTI[®] can lower one or more types of your blood cells (red blood cells, white blood cells, or platelets [cells that help blood to clot]), which may make you feel weak or tired, or increase your risk of severe infection or bleeding that may lead to death. After treatment, your healthcare provider will test your blood to check for this. Tell your healthcare provider right away if you get a fever, chills, or any signs or symptoms of an infection, are feeling tired, or have bruising or bleeding.

CARVYKTI[®] can cause serious gastrointestinal side effects, including severe or persistent diarrhea or ruptured bowel, which can be life-threatening and may lead to death. Tell your healthcare provider right away if you develop diarrhea, abdominal pain, weight loss, fever, chills, or any signs or symptoms of an infection.

CARVYKTI[®] may increase your risk of getting cancers, including certain types of blood cancers. Your healthcare provider should monitor you for this.

Having CARVYKTI[®] in your blood may cause some commercial Human Immunodeficiency Virus (HIV) tests to incorrectly give you an HIV-positive result even though you may be HIV-negative.

These are not all the possible side effects of CARVYKTI[®]. Call your healthcare provider if you have any side effects.

You may report side effects to FDA at 1-800-FDA-1088.

Please read full [Prescribing Information](#), including [Boxed Warning](#), for CARVYKTI[®].

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