HOLD ON TO HOPE

COMPLETE REMISSION IS POSSIBLE

YESCARTA® is a treatment for your non-Hodgkin lymphoma. It is used when you have failed at least two other kinds of treatment. YESCARTA® is different than other cancer medicines because it is made from your own white blood cells, which have been modified to recognize and attack your lymphoma cells.
MORE THAN HALF OF PATIENTS TREATED WITH YESCARTA® ACHIEVED COMPLETE REMISSION

In a clinical study of 101 patients with non-Hodgkin lymphoma who had failed other treatments, YESCARTA® was shown to help 51% (52 out of 101) of patients achieve complete remission with ~9-month minimum follow-up.

51% OF PATIENTS EXPERIENCED PARTIAL REMISSION

72% OF PATIENTS TREATED WITH YESCARTA® ACHIEVED A MEANINGFUL RESPONSE (OBJECTIVE RESPONSE RATE)

21% OF PATIENTS EXPERIENCED PARTIAL REMISSION

RAPID AND ONGOING RESULTS

- People generally responded to treatment within 1 month (range: 0.8–6.2 months)
- For those who achieved complete remission, the response was generally ongoing

Complete Remission: The disappearance of all signs of cancer in response to treatment. This does not always mean the cancer has been cured.


IMPORTANT SAFETY INFORMATION

What is the most important information I should know about YESCARTA®?

YESCARTA® may cause side effects that are life-threatening and can lead to death. Call or see 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
- Chills or shaking chills
- Confusion
- Dizziness or lightheadedness
- Severe nausea, vomiting, or diarrhea
- Fast or irregular heartbeat
- Severe fatigue or weakness

It is important to tell your healthcare provider that you received YESCARTA® and to show them your YESCARTA® Patient Wallet Card. Your healthcare provider may give you other medicines to treat your side effects.

Please see additional Important Safety Information throughout this brochure.
AT A 2-YEAR FOLLOW-UP, OVER HALF OF PATIENTS TREATED WITH YESCARTA® WERE STILL LIVING

- YESCARTA® was studied in a clinical trial of 101 patients with non-Hodgkin lymphoma who had failed other treatments. The FDA approved this therapy based on how many patients could achieve complete remission and how long that response would last.

- A secondary goal in the study was how long they lived overall. At the 2-year follow-up, more than half of patients treated with YESCARTA® were still alive. Not all of these patients stayed in remission.
**YESCARTA® IS DIFFERENT FROM OTHER THERAPIES YOU HAVE TRIED**

**YOUR T CELLS FIGHT TO KEEP YOU HEALTHY**
T cells are a vital tool in your immune system used to fight off harmful bacteria and viruses. They also recognize and destroy cancer cells. However, certain types of cancer cells are difficult for T cells to detect.

**YESCARTA® CAR T THERAPY MODIFIES YOUR T CELLS TO FIGHT CANCER**
YESCARTA® is a therapy that modifies your own T cells in a way that enables them to recognize and destroy cancer cells. Known as a Chimeric Antigen Receptor T-cell therapy (CAR T), YESCARTA® empowers your immune system to help fight cancer.

**YESCARTA® HELPS YOUR BODY EFFECTIVELY DESTROY CANCER CELLS**

**YESCARTA® IS AVAILABLE AT AUTHORIZED TREATMENT CENTERS**
Because YESCARTA® is different than other types of cancer treatments, it is administered at specialized hospitals. These Authorized Treatment Centers have medical staff thoroughly trained and equipped to provide CAR T therapy. These treatment centers provide diligent care throughout the YESCARTA® process and can help manage any side effects that may occur. There are ~90 Authorized Treatment Centers throughout the US. If there is not a location within 2 hours of where you live, consider looking for one near family or friends. Kite Konnect®, the YESCARTA® support team, can connect patients with independent foundations for information about transportation and housing assistance.

**FIND AN ONCOLOGIST AT AN AUTHORIZED TREATMENT CENTER**
Oncologists at Authorized Treatment Centers have the most experience with CAR T therapy and will be your most informed option for discussing YESCARTA®. Visit getstartedwithYESCARTA.com to find an Authorized Treatment Center.

**IMPORTANT SAFETY INFORMATION, CONTINUED**
Tell your healthcare provider about all the medications you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

Please see additional Important Safety Information throughout this brochure.
BEGIN WITH A BLOOD DRAW
Your blood is drawn, then passed through a machine to separate your white blood cells. This process is called leukapheresis (loo-kah-fur-ee-sis).

YOUR T CELLS ARE MODIFIED
Your separated T cells are sent to a laboratory where they are modified to become CAR T cells (YESCARTA®). During this time, you’ll undergo 3 days of low-dose conditioning chemotherapy. This prepares your body to receive YESCARTA® treatment.

CAR T CELLS ARE INFUSED BACK INTO YOUR BODY
On your scheduled treatment day, you and your caregiver will go to the Authorized Treatment Center. Your healthcare team will administer a single 30-minute infusion of YESCARTA® into your vein through a central IV line.

YOUR PROGRESS IS TRACKED
After your infusion, you’ll be monitored at the Authorized Treatment Center for at least 7 days. In order for your healthcare team to continue to monitor how the treatment is working, it is necessary to stay within 2 hours of your Authorized Treatment Center for at least 4 weeks. Kite Konnect™ can help you plan for this stay.

IF YOU EXPERIENCE SIDE EFFECTS
The Authorized Treatment Center medical staff is thoroughly trained to help. You may be hospitalized for side effects, and your healthcare provider will discharge you if your side effects are under control and it is safe for you to leave the hospital.

FOLLOW-UP VISITS WITH YOUR LOCAL HEALTHCARE PROVIDER
Your local healthcare team will perform blood tests to track your progress and help with any side effects that may occur. If you miss an appointment, it’s important to reschedule as soon as possible.
What are the possible or reasonably likely side effects of YESCARTA®?

The most common side effects of YESCARTA® include:

- Fever (100.4°F/38°C or higher)
- Low white blood cells (can occur with a fever)
- Low red blood cells
- Low blood pressure (dizziness or lightheadedness, headache, feeling tired, short of breath)
- Fast heartbeat
- Confusion
- Difficulty speaking or slurred speech
- Nausea
- Diarrhea

These are not all the possible side effects of YESCARTA®. Call your healthcare provider about any side effects that concern you. You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch or call 1-800-FDA-1088.

Please see additional Important Safety Information throughout this brochure.

When is CAR T an Option?
The treatment that’s right for you can depend on the type and stage of your cancer, your overall health, and your personal preferences. In some cases, you may need multiple phases and types of therapy. It’s important to know that even if your cancer requires more than an initial treatment, you have options.

YESCARTA® CAR T therapy is a clinically approved option when primary and secondary treatments have failed. Discussing YESCARTA® proactively with your healthcare team can help prepare you to begin therapy in the event that you need a third-line treatment.

Learn About the Treatment Journey
WHEN OTHER TREATMENTS HAVE FAILED, YESCARTA® CAR T THERAPY IS AN OPTION

1 PRIMARY TREATMENTS
The first treatment recommended based on type and stage of cancer

- Up to 6 cycles of chemotherapy with rituximab; with or without radiation therapy
- Supportive care to address any side effects or health issues that arise

SCREEN FOR RESULTS
Once chemotherapy cycles are complete, the results of a PET/CT scan are assessed to determine the next steps in treatment

CANCER IS IN REMISSION
- Signs and symptoms of cancer appear to be gone
- Routine follow-up testing with healthcare team

CANCER RESISTS OR RETURNS
- Cancer does not respond to therapy or returns after a period of improvement
- Healthcare team will consider third-line treatments

Talk to your oncologist about what treatment options may be right for you.

2 SECONDARY TREATMENTS
Secondary treatments are typically pursued if your cancer persists or returns after your primary treatment

- Additional rounds of chemotherapy; with or without rituximab
- High-dose radiation
- Stem-cell transplant with chemotherapy
- Clinical trial
- Supportive care

3 THIRD-LINE TREATMENTS
Third-line treatments are typically pursued if your cancer persists or returns after 2 or more lines of therapy

- CAR T therapy
- Clinical trial
- Supportive care
- Alternative second-line therapies, such as chemotherapy and radiation

YESCARTA® CAN PROVIDE A CHANCE FOR COMPLETE REMISSION EVEN WHEN YOUR CANCER HAS RESISTED OTHER THERAPIES

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

Please see additional Important Safety Information throughout this brochure.
IS HERE TO HELP

CONNECT WITH AN AUTHORIZED TREATMENT CENTER ONCOLOGIST
YESCARTA® is only available at Authorized Treatment Centers with medical staff thoroughly trained and equipped to provide this treatment. Kite Konnect™ can direct you to an oncologist who has experience treating with YESCARTA®.

REIMBURSEMENT SUPPORT
Benefit investigation, claims appeals information, and support for eligible and underinsured patients.

LOGISTICS SUPPORT
Providing information about potential options for transportation and housing assistance.

ONGOING COMMITMENT
Telephone support before, during, and after your treatment.

Learn more about Kite Konnect™ and YESCARTA®
1-844-454-KITE (5483), Monday-Friday, 5 am-6 pm PT

IMPORTANT SAFETY INFORMATION, CONTINUED

How will I receive YESCARTA®?

• Since YESCARTA® is made from your own white blood cells, your blood will be collected by a process called “leukapheresis” (loo-kah-fur-ee-sis), which will concentrate your white blood cells.
• Your blood cells will be sent to a manufacturing center to make your YESCARTA®.
• Before you get YESCARTA®, you will get 3 days of chemotherapy to prepare your body.
• When your YESCARTA® is ready, your healthcare provider will give it to you through a catheter placed into your vein (intravenous infusion). The infusion usually takes less than 30 minutes.
• You will be monitored where you received your treatment daily for at least 7 days after the infusion.
• You should plan to stay close to the location where you received your treatment for at least 4 weeks after getting YESCARTA®. Your healthcare provider will help you with any side effects that may occur.
• You may be hospitalized for side effects and your healthcare provider will discharge you if 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 do 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 YESCARTA®?

• Do not drive, operate heavy machinery, or do other dangerous things for 8 weeks after you get YESCARTA® because the treatment can cause sleepiness, confusion, weakness, temporary memory and coordination problems.
• Do not donate blood, organs, tissues, and cells for transplantation.

Please see additional Important Safety Information throughout this brochure.
PREPARE FOR TOMORROW

FIND AN AUTHORIZED TREATMENT CENTER ONCOLOGIST TODAY

Whether you need to make a treatment decision today or would just like information for the future, an oncologist with YESCARTA® experience is the best person to answer your questions.

GET STARTED AT

getstartedwithYESCARTA.com

• Download a CAR T Discussion Tool to help determine if YESCARTA® is an option for you
• Search for an Authorized Treatment Center to find an oncologist with YESCARTA® experience

Please see full Prescribing Information, including IMPORTANT WARNING, and Medication Guide.