Elranatamab



Care Team Contact Information:		
Pharmacy Contact Information:		
Diagnosis:		
This treatment is often use.	d for multiple myeloma, but it may be used for other diagnoses.	
Goal of Treatment:		

• Treatment may continue for a certain time period, until it no longer works, or until side effects are no longer controlled.

Treatment Regimen

Treatment Name	How the Treatment Works	How the Treatment is Given
Elranatamab (EL-rah-NA-tah-mab): Elrexfio (el-reks-fe-o)	Binds immune cells (T-cells) and cancer cells together so T-cells can more effectively attack and destroy the cancer cells	Injection given under the skin usually in the stomach area (abdomen). It may also be injected into your thigh or another area of your body.

Treatment Administration and Schedule:

Due to the risk of cytokine release syndrome (CRS), you will receive elranatamab on a "step-up dosing schedule" and may be hospitalized for 48 hours after the first "step-up" dose and for 24 hours after the second "step-up" dose.

- During the step-up dosing schedule:
 - o for your first dose, you will receive a smaller "step-up" dose on Day 1 of your treatment
 - o for your second dose, you will receive a larger "step-up" dose, which is usually given on Day 4 of your treatment
 - o for your third dose, you will receive the first full "treatment" dose, which is usually given on Day 8 of your treatment
- If your dose is delayed for any reason, you may need to repeat the step-up dosing schedule.

Weeks 1 and 2

Step-Up Dosing Schedule							
Treatment	Week 1						
Name	Day 1	Day 2	Day 3	Day 4	Day 5	Day 6	Day 7
Elranatamab	✓			~			
	First "Step-Up" Dose			Second "Step-Up" Dose			
	Week 2						
	Day 8	Day 9	Day 10	Day 11	Day 12	Day 3	Day 14
Elranatamab	✓						
	First Full "Treatment" Dose						







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Treatment Administration and Schedule (Continued)

Week 3 and Beyond

- After you receive your first full "treatment" dose, elranatamab is usually given 1 time each week through Week 24.
- Starting on Week 25, your future doses will usually be given 1 time every 2 weeks.
- Starting on Week 49, your future doses will usually be given 1 time every 4 weeks.
- Your care team will decide the time between doses and how long you will receive treatment.

Appointments: Appointments may include regular check-ups with your care team, treatment appointments, and lab and imaging visits. It's important to keep your appointments whenever you can. If you miss any appointments, call your care provider as soon as possible to reschedule your appointment.

Supportive Care to Prevent and Treat Side Effects

Description	Supportive Care Given at the Clinic or Hospital	Supportive Care Taken at Home
To help lower the risk of		
Cytokine Release Syndrome		
(CRS)		
To help lower the risk of		
infections		
Other		









Common Side Effects

Side Effect	Important Information	
Cytokine Release Syndrome	Description: CRS happens when your immune system becomes overactive. Most CR few doses. However, some CRS events can be serious and life-threatening. Symptor lightheaded, or difficulty breathing.	
(CRS) (Boxed Warning)	 Recommendations: Keep a symptom diary to record any new or worsening symptoms such as fever, chills, fatigue, or difficulty breathing. At the guidance of your care team, check vitals including temperature, blood pressure, and heart rate. Stay hydrated by drinking plenty of fluids to help manage symptoms and support overall health. Your care team may prescribe medications to help manage symptoms. 	Talk to your care team if you have: Fever of 100.4°F (38°C) or higher Trouble breathing Chills Dizziness or light-headedness Fast heartbeat Headache Increased liver enzymes in your blood
		Note: Your care team may have specific numbers for blood pressure, heart rate, and blood oxygen levels. If your numbers go beyond those limits, call your care team or get emergency help right away.
Low White Blood Cell (WBC) Count and Increased Risk of Infection	Description: WBCs help protect the body against infections. If you have a low WBC content of Recommendations: • Wash your hands and bathe regularly. • Avoid crowded places. • Stay away from people who are sick. • Your care team may prescribe a drug that promotes the growth of WBCs.	ount, you may be at a higher risk of infection. Talk to your care team if you have: Fever of 100.4°F (38°C) or higher Chills Cough Sore throat Painful urination Tiredness that is worse than normal Shortness of breath Feeling weak/generally unwell
Low Red Blood Cell (RBC) Count and Hemoglobin (Hgb)	Description: RBCs and Hgb help bring oxygen to your body's tissues and take away of tired, or look pale. Recommendations: Get 7 to 8 hours of sleep each night. Avoid operating heavy machinery when tired. Balance work and rest, staying active but resting when needed.	Talk to your care team if you have: Shortness of breath Dizziness Fast or abnormal heartbeats Severe headache









Low Platelet	Platelet Description: Platelets help the blood clot and heal wounds. If you have low platelet counts, you are at a higher risk of bruising and bleed			
Count	 Recommendations: Blow your nose gently and avoid picking it. Brush your teeth gently with a soft toothbrush and maintain good oral hygiene. Use an electric razor for shaving and a nail file instead of nail clippers. Avoid over-the-counter medications that may increase the risk of bleeding, such as NSAIDs. Consult your care team or dentist before medical or dental procedures, as you may need to pause your medication. 	 Talk to your care team if you have: Nosebleed lasting over 5 minutes despite pressure Cut that continues to bleed Significant gum bleeding when flossing or brushing Severe headaches Blood in your urine or stool Blood in your spit after a cough 		
Diarrhea	Description: Diarrhea is when you have loose, watery stools. The need to use the bar Recommendations: • Keep track of how many times you go to the bathroom each day. • Drink 8 to 10 glasses of water or other fluids every day, unless your doctor tells you otherwise. • Eat small meals of mild, low-fiber foods like bananas, applesauce, potatoes, chicken, rice, and toast. • Stay away from foods with high fiber (like raw vegetables, fruits, and whole grains), foods that cause gas (like broccoli and beans), dairy foods (like yogurt and milk), and spicy, fried, and greasy foods. • Your care team might suggest a medicine for diarrhea.	throom may occur urgently. Talk to your care team if you have: • 4 or more bowel movements than normal in 24 hours • Dizziness or lightheadedness while having diarrhea • Bloody diarrhea		
Liver Problems	Description: Your liver may not work as well as it should, which may lead to symptom performed to monitor liver function. Talk to your care team if you have: • Yellowing of the skin or whites of your eyes • Dark or brown urine • Bleeding or bruising	 s like fatigue, nausea, or swelling in your belly. Lab tests might be Tiredness worse than baseline Loss of appetite Pain in the right upper stomach area 		
Fatigue	 Description: Fatigue is a constant and sometimes strong feeling of tiredness. Recommendations: Routine exercise has been shown to decrease levels of fatigue. Work with your care team to find the right type of exercise for you. Ask your family and friends for help with daily tasks and emotional support. Try healthy ways to feel better, like meditation, writing in a journal, doing yoga, and using guided imagery to lower anxiety and feel good. Make a regular sleep schedule and limit naps during the day so you can sleep better at night, aiming for 7 to 8 hours of sleep. Don't use heavy machines or do things that need your full attention if you're very tired to avoid accidents. 	Talk to your care team if you have:		









Muscle or Joint Pain or	Description: You may experience muscle or joint discomfort that makes daily activities harder. Symptoms can include aching, stiffness, swelling, and weakness, which can hinder lifting objects or walking long distances. You may also feel fatigued, making it difficult to stay active.			
Weakness	 Recommendations: Track your pain levels, areas of discomfort, and any activities that worsen or improve your symptoms. Engage in gentle exercises like walking, stretching, or yoga to maintain mobility and strength, but consult your care team before starting any new exercise routine. Apply a warm compress to relax stiff muscles or use cold packs to reduce swelling and numb pain in affected areas. Your care team may prescribe or recommend medications, including overthe-counter pain relievers. 	 Talk to your care team if you have: Persistent or worsening muscle or joint pain that does not improve with home treatments New symptoms, such as swelling, redness, or warmth in the joints Severe weakness that affects your ability to perform daily activities Side effects from pain relief medications that you are unable to manage 		
Injection Site Reactions	Description: An injection reaction is a bad response that happens during or not long a injection site are possible. Symptoms may include itching, swelling, bruising, pain, ras Talk to your care team if you have: Chills or shaking Itching, rash, or flushing Trouble breathing or wheezing; tongue-swelling			

Select Rare and Serious Side Effects

Side Effect	Talk to Your Care Team if You Have Any of These Signs or Symptoms		
Neurologic Problems (Boxed Warning) This includes Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS)	 Headache Agitation, trouble staying awake, confusion or disorientation, seeing or hearing things that are not real (hallucinations) Trouble speaking, thinking, remembering things, paying attention, or understanding things 	 Problems walking, muscle weakness, shaking (tremors), loss of balance, or muscle spasms Numbness and tingling (feeling like "pins and needles") Burning, throbbing, or stabbing pain Changes in your handwriting 	
Low Immunoglobulin Levels	 Getting sick often (like colds or pneumonia) Taking longer to feel better after being sick Feeling tired or weak 	 Having skin infections or rashes Getting upset stomachs or diarrhea more frequently Possibly having allergies or other immune problems 	

Before starting treatment, ask your care team when to call 9-1-1 or seek emergency help.

If you experience any new, worsening, or uncontrolled side effects, contact your care team immediately.









Intimacy, Pregnancy, and Breastfeeding

- Treatment may **change how you feel about intimacy and your body.** However, physical closeness—such as holding hands and hugging—remains safe. It is common to have questions about intimacy. If needed, talk to your care team for guidance.
- Treatment may harm an unborn baby.
 - o If you might get pregnant, take a pregnancy test before starting treatment.
 - Use an effective method of birth control during treatment and for 4 months after your last dose.
 - If you think you might be pregnant or if you become pregnant, tell your care team right away.
- Do NOT breastfeed during treatment and for 4 months after your last dose.

Additional Information

- Tell your care team about all the medicines you take.
 - This includes prescriptions, over-the-counter drugs, vitamins, and herbal products. Before starting any new medicine, supplement, or vaccine, ask your care team first.
- The most common severe changes in laboratory test results are decreased white blood cells, red blood cells, and platelets.
- Your care team will monitor you for **signs and symptoms of CRS and neurologic problems** during treatment with elranatamab, as well as other side effects and treat you if needed. Your care provider may temporarily stop or completely stop your treatment with elranatamab if you develop CRS, neurologic problems, or any other side effects that are severe.
- Do not drive or operate heavy or dangerous machinery during and for 48 hours after completing each of the 2 doses of elranatamab that are part of the "step-up dosing schedule" and your first full treatment dose, and at any time during treatment with elranatamab if you develop new neurologic symptoms until the symptoms go away.
- Elranatamab is available only through a special FDA program called the ELREXFIO Risk Evaluation and Mitigation Strategy (REMS). You will receive a Patient Wallet Card from your healthcare provider. Carry the Patient Wallet Card with you at all times and show it to all of your healthcare providers. The Patient Wallet Card lists signs and symptoms of CRS and neurologic problems.
- This Patient Education Sheet may not describe all possible side effects.
 Call your healthcare provider for medical advice about side effects. You may report side effects to the FDA at 1-800-FDA-1088.

Notes

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Scan the QR code below to access this education sheet.









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Important notice: The Association of Cancer Care Centers (ACCC), Hematology/Oncology Pharmacy Association (HOPA), Network for Collaborative Oncology Devèlopment & Advancement, Inc. (NCODA), and Oncology Nursing Society (ONS) have collaborated in gathering information for and developing this patient education guide. This guide represents a brief summary of the medication derived from information provided by the drug manufacturer and other resources.

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