

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use ELREXFIO safely and effectively. See full prescribing information for ELREXFIO.

ELREXFIO™ (elranatamab-bcmm) injection, for subcutaneous use
Initial U.S. Approval: 2023

WARNING: CYTOKINE RELEASE SYNDROME and NEUROLOGIC TOXICITY including IMMUNE EFFECTOR CELL-ASSOCIATED NEUROTOXICITY SYNDROME

See full prescribing information for complete boxed warning.

- Cytokine Release Syndrome (CRS), including life-threatening or fatal reactions, can occur in patients receiving ELREXFIO. Initiate treatment with ELREXFIO step-up dosing schedule to reduce risk of CRS. Withhold ELREXFIO until CRS resolves or permanently discontinue based on severity. (2.2, 2.5, 5.1)
- Neurologic toxicity, including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS), and serious and life-threatening reactions, can occur in patients receiving ELREXFIO. Monitor patients for signs and symptoms of neurologic toxicity, including ICANS, during treatment. Withhold ELREXFIO until the neurologic toxicity resolves or permanently discontinue based on severity. (2.5, 5.2)
- ELREXFIO is available only through a restricted program called the ELREXFIO Risk Evaluation and Mitigation Strategy (REMS). (5.3)

----- **INDICATIONS AND USAGE** -----

ELREXFIO is a bispecific B-cell maturation antigen (BCMA)-directed CD3 T-cell engager indicated for the treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy including a proteasome inhibitor, an immunomodulatory agent, and an anti-CD38 monoclonal antibody.

This indication is approved under accelerated approval based on response rate and durability of response. Continued approval for this indication may be contingent upon verification of clinical benefit in a confirmatory trial(s). (1)

----- **DOSAGE AND ADMINISTRATION** -----

| ELREXFIO Dosing Schedule (2.2) | | | |
|--------------------------------|---|----------------------------|-------|
| Dosing Schedule | Day | ELREXFIO Dose | |
| Step-up Dosing Schedule | Day 1 | Step-up dose 1 | 12 mg |
| | Day 4 | Step-up dose 2 | 32 mg |
| | Day 8 | First treatment dose | 76 mg |
| Weekly Dosing Schedule | One week after first treatment dose and weekly thereafter through week 24 | Subsequent treatment doses | 76 mg |

| ELREXFIO Dosing Schedule (2.2) | | | |
|---|--------------------------------------|----------------------------|-------|
| Dosing Schedule | Day | ELREXFIO Dose | |
| Biweekly (Every 2 Weeks) Dosing Schedule* | Week 25 and every 2 weeks thereafter | Subsequent treatment doses | 76 mg |
| *Responders only week 25 onward. | | | |

- Patients should be hospitalized for 48 hours after administration of the first step-up dose, and for 24 hours after administration of the second step-up dose. (2.1)
- For subcutaneous injection only. (2.2)
- Administer pre-treatment medications as recommended. (2.3)
- See Full Prescribing Information for instructions on preparation and administration. (2.6)

----- **DOSAGE FORMS AND STRENGTHS** -----

- Injection:
- 76 mg/1.9 mL (40 mg/mL) in a single-dose vial. (3)
 - 44 mg/1.1 mL (40 mg/mL) in a single-dose vial. (3)

----- **CONTRAINDICATIONS** -----

None. (4)

----- **WARNINGS AND PRECAUTIONS** -----

- **Infections:** Can cause severe, life-threatening, or fatal infections. Monitor patients for signs and symptoms of infection and treat appropriately. Do not initiate treatment in patients with active infections. (5.4)
- **Neutropenia:** Monitor complete blood cell counts at baseline and periodically during treatment. (5.5)
- **Hepatotoxicity:** Can cause elevated ALT, AST, and bilirubin. Monitor liver enzymes and bilirubin at baseline and during treatment as clinically indicated. (5.6)
- **Embryo-Fetal Toxicity:** May cause fetal harm. Advise females of reproductive potential of the potential risk to the fetus and to use effective contraception. (5.7, 8.1, 8.3)

----- **ADVERSE REACTIONS** -----

Most common adverse reactions (incidence ≥20%) are CRS, fatigue, injection site reaction, diarrhea, upper respiratory tract infection, musculoskeletal pain, pneumonia, decreased appetite, rash, cough, nausea, and pyrexia. The most common Grade 3 to 4 laboratory abnormalities (≥30%) are decreased lymphocytes, decreased neutrophils, decreased hemoglobin, decreased white blood cells, and decreased platelets. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Pfizer Inc. at 1-800-438-1985 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

----- **USE IN SPECIFIC POPULATIONS** -----

Lactation: Advise not to breastfeed. (8.2)

See 17 for PATIENT COUNSELING INFORMATION and Medication Guide.

Revised: 8/2023

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FULL PRESCRIBING INFORMATION

WARNING: CYTOKINE RELEASE SYNDROME and NEUROLOGIC TOXICITY including IMMUNE EFFECTOR CELL-ASSOCIATED NEUROTOXICITY SYNDROME

- Cytokine Release Syndrome (CRS), including life-threatening or fatal reactions, can occur in patients receiving ELREXFIO. Initiate treatment with ELREXFIO step-up dosing schedule to reduce the risk of CRS. Withhold ELREXFIO until CRS resolves or permanently discontinue based on severity [see *Dosage and Administration* (2.2, 2.5), *Warnings and Precautions* (5.1)].
- Neurologic toxicity, including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS), and serious and life-threatening reactions, can occur in patients receiving ELREXFIO. Monitor patients for signs and symptoms of neurologic toxicity, including ICANS, during treatment. Withhold ELREXFIO until the neurologic toxicity resolves or permanently discontinue based on severity [see *Dosage and Administration* (2.5), *Warnings and Precautions* (5.2)].
- Because of the risk of CRS and neurologic toxicity, including ICANS, ELREXFIO is available only through a restricted program under a Risk Evaluation and Mitigation Strategy (REMS) called ELREXFIO REMS [see *Warnings and Precautions* (5.3)].

1 INDICATIONS AND USAGE

ELREXFIO is indicated for the treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy, including a proteasome inhibitor, an immunomodulatory agent, and an anti-CD38 monoclonal antibody.

This indication is approved under accelerated approval based on response rate and durability of response [see *Clinical Studies* (14)]. Continued approval for this indication may be contingent upon verification of clinical benefit in a confirmatory trial(s).

2 DOSAGE AND ADMINISTRATION

2.1 Important Dosing Information

Administer ELREXFIO subcutaneously according to the step-up dosing schedule to reduce the incidence and severity of cytokine release syndrome (CRS).

Administer pre-treatment medications prior to each dose in the ELREXFIO step-up dosing schedule, which includes step-up dose 1, step-up dose 2, and the first treatment dose as recommended [see *Dosage and Administration* (2.2, 2.3)].

ELREXFIO should only be administered by a qualified healthcare professional with appropriate medical support to manage severe reactions such as CRS and neurologic toxicity, including ICANS [see *Warnings and Precautions* (5.1, 5.2)].

Due to the risk of CRS, patients should be hospitalized for 48 hours after administration of the first step-up dose, and for 24 hours after administration of the second step-up dose.

2.2 Recommended Dosage

For subcutaneous injection only.

The recommended dosing schedule for ELREXFIO is provided in Table 1. The recommended dosages of ELREXFIO subcutaneous injection are: step-up dose 1 of 12 mg on Day 1, step-up dose 2 of 32 mg on Day 4, followed by the first treatment dose of 76 mg on Day 8, and then 76 mg weekly thereafter through week 24.

For patients who have received at least 24 weeks of treatment with ELREXFIO and have achieved a response [partial response (PR) or better] and maintained this response for at least 2 months, the dose interval should transition to an every two-week schedule.

Continue treatment with ELREXFIO until disease progression or unacceptable toxicity.

Administer pre-treatment medications prior to each dose in the ELREXFIO step-up dosing schedule, which includes step-up dose 1, step-up dose 2, and the first treatment dose as recommended [see *Dosage and Administration (2.3)*].

Table 1. ELREXFIO Dosing Schedule

| Dosing Schedule | Day | ELREXFIO Dose | |
|--|--|----------------------------|-------|
| | | | |
| Step-up Dosing Schedule | Day 1 ^a | Step-up dose 1 | 12 mg |
| | Day 4 ^{a,b} | Step-up dose 2 | 32 mg |
| | Day 8 ^{a,c} | First treatment dose | 76 mg |
| Weekly Dosing Schedule | One week after first treatment dose and weekly thereafter ^d through week 24 | Subsequent treatment doses | 76 mg |
| Biweekly (Every 2 Weeks) Dosing Schedule *Responders only week 25 onward | Week 25 and every 2 weeks thereafter ^d | Subsequent treatment doses | 76 mg |

- a. Administer pre-treatment medications prior to each dose in the ELREXFIO step-up dosing schedule, which includes step-up dose 1, step-up dose 2, and the first treatment dose [see *Dosage and Administration (2.3)*].
- b. A minimum of 2 days should be maintained between step-up dose 1 (12 mg) and step-up dose 2 (32 mg).
- c. A minimum of 3 days should be maintained between step-up dose 2 (32 mg) and the first treatment (76 mg) dose.
- d. A minimum of 6 days should be maintained between treatment doses.

Note: See Table 2 for recommendations on restarting ELREXFIO after dose delays.

2.3 Recommended Pre-treatment Medications

Administer the following pre-treatment medications approximately 1 hour before the first three doses of ELREXFIO in the step-up dosing schedule, which includes step-up dose 1, step-up dose 2, and the first treatment dose as described in Table 1 to reduce the risk of CRS [see *Warnings and Precautions (5.1)*]:

- acetaminophen (or equivalent) 650 mg orally
- dexamethasone (or equivalent) 20 mg orally or intravenously
- diphenhydramine (or equivalent) 25 mg orally

2.4 Restarting ELREXFIO After Dosage Delay

If a dose of ELREXFIO is delayed, restart therapy based on the recommendations listed in Table 2 and resume the dosing schedule accordingly [see *Dosage and Administration (2.2)*]. Administer pre-treatment medications as indicated in Table 2.

Table 2. Recommendation for Restarting Therapy with ELREXFIO After Dosage Delay

| Last Dose Administered | Time Since the Last Dose Administered | Action for Next Dose |
|----------------------------|--|--|
| Step-up dose 1 (12 mg) | 2 weeks or less (≤ 14 days) | Restart ELREXFIO at step-up dose 2 (32 mg). ^a If tolerated, increase to 76 mg 4 days later. |
| | Greater than 2 weeks (> 14 days) | Restart ELREXFIO step-up dosing schedule at step-up dose 1 (12 mg). ^a |
| Step-up dose 2 (32 mg) | 2 weeks or less (≤ 14 days) | Restart ELREXFIO at 76 mg. ^a |
| | Greater than 2 weeks to less than or equal to 4 weeks (15 days to ≤ 28 days) | Restart ELREXFIO at step-up dose 2 (32 mg). ^a If tolerated, increase to 76 mg 1 week later. |
| | Greater than 4 weeks (> 28 days) | Restart ELREXFIO step-up dosing schedule at step-up dose 1 (12 mg). ^a |
| Any treatment dose (76 mg) | 6 weeks or less (≤ 42 days) | Restart ELREXFIO at 76 mg. |
| | Greater than 6 weeks to less or equal to 12 weeks (43 days to ≤ 84 days) ^b | Restart ELREXFIO at step-up dose 2 (32 mg). ^a If tolerated, increase to 76 mg 1 week later. |
| | Greater than 12 weeks (> 84 days) ^b | Restart ELREXFIO step-up dosing schedule at step-up dose 1 (12 mg). ^a |

a. Administer pre-treatment medications prior to the ELREXFIO dose [see *Dosage and Administration (2.3)*].

b. Consider benefit-risk of restarting ELREXFIO in patients who require a dose delay of more than 42 days due to an adverse reaction.

2.5 Dosage Modifications for Adverse Reactions

Dosage reductions of ELREXFIO are not recommended.

Dosage delays may be required to manage toxicities related to ELREXFIO [see *Warnings and Precautions (5)*]. Recommendations on restarting ELREXFIO after a dose delay are provided in Table 2.

See Table 3 and Table 4 for recommended actions for adverse reactions of CRS and ICANS, respectively. See Table 5 for recommended actions for neurologic toxicity excluding ICANS and Table 6 for recommended actions for other adverse reactions following administration of ELREXFIO. Consider further management per current practice guidelines.

Management of CRS, Neurologic Toxicity Including ICANS

Cytokine Release Syndrome (CRS)

Management recommendations for CRS are summarized in Table 3.

Identify CRS based on clinical presentation [see *Warnings and Precautions (5.1)*]. Evaluate and treat other causes of fever, hypoxia, and hypotension.

If CRS is suspected, withhold ELREXFIO until CRS resolves. Manage CRS according to the recommendations in Table 3 and consider further management per current practice guidelines. Administer supportive therapy for CRS, which may include intensive care for severe or life-threatening CRS. Consider laboratory testing to monitor for disseminated intravascular coagulation (DIC), hematology parameters, as well as pulmonary, cardiac, renal, and hepatic function.

Table 3. Recommendations for Management of CRS

| Grade ^a | Presenting Symptoms | Actions |
|----------------------------|--|---|
| Grade 1 | Temperature ≥ 100.4 °F (38 °C) ^b | <ul style="list-style-type: none"> Withhold ELREXFIO until CRS resolves.^c Administer pretreatment medications prior to next dose of ELREXFIO. |
| Grade 2 | Temperature ≥ 100.4 °F (38 °C) with either: <ul style="list-style-type: none"> Hypotension responsive to fluid and not requiring vasopressors, and/or Oxygen requirement of low-flow nasal cannula^d or blow-by | <ul style="list-style-type: none"> Withhold ELREXFIO until CRS resolves.^c Monitor patients daily for 48 hours following the next dose of ELREXFIO. Instruct patients to remain within proximity of a healthcare facility, and consider hospitalization. Administer pretreatment medications prior to next dose of ELREXFIO. |
| Grade 3 (First occurrence) | Temperature ≥ 100.4 °F (38 °C) with either: <ul style="list-style-type: none"> Hypotension requiring one vasopressor with or without vasopressin, and/or Oxygen requirement of high-flow nasal cannula^d, facemask, non-rebreather mask, or Venturi mask | <ul style="list-style-type: none"> Withhold ELREXFIO until CRS resolves.^c Provide supportive therapy, which may include intensive care. Patients should be hospitalized for 48 hours following the next dose of ELREXFIO. Administer pretreatment medications prior to next dose of ELREXFIO. |
| Grade 3 (Recurrent) | Temperature ≥ 100.4 °F (38 °C) with either: <ul style="list-style-type: none"> Hypotension requiring one vasopressor with or without vasopressin, and/or Oxygen requirement of high-flow nasal cannula^d, facemask, non-rebreather mask, or Venturi mask | <ul style="list-style-type: none"> Permanently discontinue therapy with ELREXFIO. Provide supportive therapy, which may include intensive care. |
| Grade 4 | Temperature ≥ 100.4 °F (38 °C) with either: <ul style="list-style-type: none"> Hypotension requiring multiple vasopressors (excluding vasopressin), and/or Oxygen requirement of positive pressure (e.g., continuous positive airway pressure [CPAP], bilevel positive airway pressure [BiPAP], intubation, and mechanical ventilation) | <ul style="list-style-type: none"> Permanently discontinue therapy with ELREXFIO. Provide supportive therapy, which may include intensive care. |

- a. Based on American Society for Transplantation and Cellular Therapy (ASTCT) 2019 grading criteria for CRS.
b. Attributed to CRS. Fever may not always be present concurrently with hypotension or hypoxia as it may be masked by interventions such as antipyretics or anti-cytokine therapy.
c. See Table 2 for recommendations on restarting ELREXFIO after dose delays.
d. Low-flow nasal cannula is ≤ 6 L/min, and high-flow nasal cannula is >6 L/min.

Neurologic Toxicity Including ICANS

Management recommendations for ICANS and neurologic toxicity are summarized in Table 4 and Table 5.

At the first sign of neurologic toxicity, including ICANS, withhold ELREXFIO and consider neurology evaluation. Rule out other causes of neurologic symptoms. Provide supportive therapy, which may include intensive care, for severe or life-threatening neurologic toxicities, including ICANS [see *Warnings and Precautions (5.2)*]. Manage ICANS according to the recommendations in Table 4 and consider further management per current practice guidelines.

Table 4. Recommendations for Management of ICANS

| Grade ^a | Presenting Symptoms ^b | Actions |
|-------------------------------|---|--|
| Grade 1 | ICE score 7-9 ^c Or depressed level of consciousness ^d : awakens spontaneously. | <ul style="list-style-type: none"> • Withhold ELREXFIO until ICANS resolves.^e • Monitor neurologic symptoms and consider consultation with a neurologist and other specialists for further evaluation and management. • Consider non-sedating, anti-seizure medications (e.g., levetiracetam) for seizure prophylaxis. |
| Grade 2 | ICE score 3-6 ^c Or depressed level of consciousness ^d : awakens to voice. | <ul style="list-style-type: none"> • Withhold ELREXFIO until ICANS resolves.^e • Administer dexamethasone^f 10 mg intravenously every 6 hours. Continue dexamethasone use until resolution to Grade 1 or less, then taper. • Monitor neurologic symptoms and consider consultation with a neurologist and other specialists for further evaluation and management. • Consider non-sedating, anti-seizure medications (e.g., levetiracetam) for seizure prophylaxis. • Monitor patients daily for 48 hours following the next dose of ELREXFIO. Instruct patients to remain within proximity of a healthcare facility, and consider hospitalization. |
| Grade 3 (First occurrence) | ICE score 0-2 ^c or depressed level of consciousness ^d : awakens only to tactile stimulus, or seizures ^d , either: <ul style="list-style-type: none"> • any clinical seizure, focal or generalized, that resolves rapidly, or • non-convulsive seizures on electroencephalogram (EEG) that resolve with intervention, or raised intracranial pressure: focal/local edema on neuroimaging ^d | <ul style="list-style-type: none"> • Withhold ELREXFIO until ICANS resolves.^e • Administer dexamethasone^f 10 mg intravenously every 6 hours. Continue dexamethasone use until resolution to Grade 1 or less, then taper. • Monitor neurologic symptoms and consider consultation with a neurologist and other specialists for further evaluation and management. • Consider non-sedating, anti-seizure medications (e.g., levetiracetam) for seizure prophylaxis. • Provide supportive therapy, which may include intensive care. • Patients should be hospitalized for 48 hours following the next dose of ELREXFIO. |
| Grade 3 (recurrent) | ICE score 0-2 ^c or depressed level of consciousness ^d : awakens only to tactile stimulus, or seizures ^d , either: <ul style="list-style-type: none"> • any clinical seizure, focal or generalized, that resolves rapidly, or • non-convulsive seizures on electroencephalogram (EEG) that resolve with intervention, or raised intracranial pressure: focal/local edema on neuroimaging ^d | <ul style="list-style-type: none"> • Permanently discontinue ELREXFIO. • Administer dexamethasone^f 10 mg intravenously every 6 hours. Continue dexamethasone use until resolution to Grade 1 or less, then taper. • Monitor neurologic symptoms and consider consultation with a neurologist and other specialists for further evaluation and management. • Consider non-sedating, anti-seizure medications (e.g., levetiracetam) for seizure prophylaxis. • Provide supportive therapy, which may include intensive care. |

Table 4. Recommendations for Management of ICANS

| Grade ^a | Presenting Symptoms ^b | Actions |
|--------------------|--|---|
| Grade 4 | <p>ICE score 0^c</p> <p>Or, depressed level of consciousness^d either:</p> <ul style="list-style-type: none"> • patient is unarousable or requires vigorous or repetitive tactile stimuli to arouse, or • stupor or coma, <p>or seizures^d, either:</p> <ul style="list-style-type: none"> • life-threatening prolonged seizure (>5 minutes), or • repetitive clinical or electrical seizures without return to baseline in between, <p>or motor findings^d:</p> <ul style="list-style-type: none"> • deep focal motor weakness such as hemiparesis or paraparesis, <p>or raised intracranial pressure/cerebral edema^d, with signs/symptoms such as:</p> <ul style="list-style-type: none"> • diffuse cerebral edema on neuroimaging, or • decerebrate or decorticate posturing, or • cranial nerve VI palsy, or • papilledema, or • Cushing’s triad | <ul style="list-style-type: none"> • Permanently discontinue ELREXFIO. • Administer dexamethasone^f 10 mg intravenously every 6 hours. Continue dexamethasone use until resolution to Grade 1 or less, then taper. • Alternatively, consider administration of methylprednisolone 1,000 mg per day intravenously for 3 days. • Monitor neurologic symptoms and consider consultation with a neurologist and other specialists for further evaluation and management. • Consider non-sedating, anti-seizure medications (e.g., levetiracetam) for seizure prophylaxis. • Provide supportive therapy, which may include intensive care. |

- a. Based on American Society for Transplantation and Cellular Therapy (ASTCT) 2019 grading criteria for ICANS.
- b. Management is determined by the most severe event, not attributable to any other cause.
- c. If patient is arousable and able to perform Immune Effector Cell-Associated Encephalopathy (ICE) Assessment, assess: Orientation (oriented to year, month, city, hospital = 4 points); Naming (name 3 objects, e.g., point to clock, pen, button = 3 points); Following Commands (e.g., “show me 2 fingers” or “close your eyes and stick out your tongue” = 1 point); Writing (ability to write a standard sentence = 1 point); and Attention (count backwards from 100 by ten = 1 point). If patient is unarousable and unable to perform ICE Assessment (Grade 4 ICANS) = 0 points.
- d. Not attributable to any other cause.
- e. See Table 2 for recommendations on restarting ELREXFIO after dose delays.
- f. All references to dexamethasone administration are dexamethasone or equivalent medications.

Table 5. Recommendations for Management of Neurologic Toxicity, Excluding ICANS

| Adverse Reaction | Severity | Actions |
|---------------------------------------|---------------------------------------|---|
| Neurologic Toxicity (excluding ICANS) | Grade 1 | <ul style="list-style-type: none"> • Withhold ELREXFIO until neurologic toxicity symptoms resolve or stabilize. |
| | Grade 2 Grade 3 (First occurrence) | <ul style="list-style-type: none"> • Withhold ELREXFIO until neurologic toxicity symptoms improve to Grade 1 or less. • Provide supportive therapy. |
| | Grade 3 (Recurrent) Grade 4 | <ul style="list-style-type: none"> • Permanently discontinue ELREXFIO. • Provide supportive therapy, which may include intensive care. |

Table 6. Recommended Dosage Modifications for Other Adverse Reactions

| Adverse Reactions | Severity | Actions |
|--|---|---|
| Hematologic Adverse Reactions [see Warnings and Precautions (5.5)] | Absolute neutrophil count less than $0.5 \times 10^9/L$ | <ul style="list-style-type: none"> Withhold ELREXFIO until absolute neutrophil count is $0.5 \times 10^9/L$ or higher.^b |
| | Febrile neutropenia | <ul style="list-style-type: none"> Withhold ELREXFIO until absolute neutrophil count is $1 \times 10^9/L$ or higher and fever resolves.^b |
| | Hemoglobin less than 8 g/dL | <ul style="list-style-type: none"> Withhold ELREXFIO until hemoglobin is 8 g/dL or higher.^b |
| | Platelet count less than 25,000/mcL Platelet count between 25,000/mcL and 50,000/mcL with bleeding | <ul style="list-style-type: none"> Withhold ELREXFIO until platelet count is 25,000/mcL or higher and no evidence of bleeding.^b |
| Infections and Other Non-hematologic Adverse Reactions ^a [see Warnings and Precautions (5.4, 5.6) and Adverse Reactions (6.1)] | Grade 3 | <ul style="list-style-type: none"> Withhold ELREXFIO until adverse reaction improves to \leqGrade 1 or baseline.^b |
| | Grade 4 | <ul style="list-style-type: none"> Consider permanent discontinuation of ELREXFIO. If ELREXFIO is not permanently discontinued, withhold subsequent treatment doses of ELREXFIO (e.g., doses administered after ELREXFIO step-up dosing schedule) until adverse reaction improves to Grade 1 or less. |

a. Based on National Cancer Institute Common Terminology Criteria for Adverse Events (NCI-CTCAE), Version 5.0.

b. See Table 2 for recommendations on restarting ELREXFIO after dose delays.

2.6 Preparation and Administration Instructions

ELREXFIO is intended for subcutaneous use by a healthcare provider only.

ELREXFIO should be administered by a healthcare provider with adequate medical personnel and appropriate medical equipment to manage severe reactions, including CRS and neurologic toxicity, including ICANS [see Warnings and Precautions (5.1, 5.2)].

ELREXFIO 76 mg/1.9 mL (40 mg/mL) vial and 44 mg/1.1 mL (40 mg/mL) vial are supplied as ready-to-use solution that do not need dilution prior to administration.

ELREXFIO is a clear to slightly opalescent, and colorless to pale brown liquid solution. Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. Do not administer if solution is discolored or contains particulate matter.

Use aseptic technique to prepare and administer ELREXFIO.

Preparation

ELREXFIO vials are for one-time use in a single patient and do not contain any preservatives.

Prepare ELREXFIO following the instructions below (see Table 7) depending on the required dose. Use a 44 mg/1.1 mL (40 mg/mL) single-dose vial for step-up dose 1 or step-up dose 2.

Table 7. Injection Volumes

| Total Dose (mg) | Volume of Injection |
|-----------------|---------------------|
| 12 mg | 0.3 mL |
| 32 mg | 0.8 mL |
| 76 mg | 1.9 mL |

Remove the appropriate strength ELREXFIO vial from refrigerated storage 2 °C to 8 °C (36 °F to 46 °F)]. Once removed from refrigerated storage, equilibrate ELREXFIO to ambient temperature [15 °C to 30 °C (59 °F to 86 °F)]. Do not warm ELREXFIO in any other way.

Withdraw the required injection volume of ELREXFIO from the vial into an appropriately sized syringe with stainless steel injection needles (30G or wider) and polypropylene or polycarbonate syringe material. Discard unused portion.

Administration

Inject the required volume of ELREXFIO into the subcutaneous tissue of the abdomen (preferred injection site). Alternatively, ELREXFIO may be injected into the subcutaneous tissue at other sites (e.g., thigh).

Do not inject into tattoos or scars or areas where the skin is red, bruised, tender, hard or not intact.

Storage of Prepared Syringe

If the prepared dosing syringe is not used immediately, store syringe between 2 °C (36 °F) to 30 °C (86 °F) for a maximum of 4 hours.

3 DOSAGE FORMS AND STRENGTHS

ELREXFIO injection is a clear to slightly opalescent, and colorless to pale brown liquid solution available as:

- 76 mg/1.9 mL (40 mg/mL) in a single-dose vial
- 44 mg/1.1 mL (40 mg/mL) in a single-dose vial

4 CONTRAINDICATIONS

None.

5 WARNINGS AND PRECAUTIONS

5.1 Cytokine Release Syndrome (CRS)

ELREXFIO can cause CRS, including life-threatening or fatal reactions [see *Adverse Reactions (6.1)*].

In the clinical trial, CRS occurred in 58% of patients who received ELREXFIO at the recommended dosing schedule [see *Dosage and Administration (2.2)*], with Grade 1 CRS in 44% of patients, Grade 2 CRS in 14% of patients, and Grade 3 CRS in 0.5% of patients. Recurrent CRS occurred in 13% of patients. Most patients experienced CRS after the first step-up dose (43%) or the second step-up dose (19%), with 7% of patients having CRS after the first treatment dose and 1.6% of patients after a subsequent dose. The median time to onset of CRS was 2 (range: 1 to 9) days after the most recent dose, with a median duration of 2 (range: 1 to 19) days.

Clinical signs and symptoms of CRS may include, but are not limited to, fever, hypoxia, chills, hypotension, tachycardia, headache, and elevated liver enzymes.

Initiate therapy according to the ELREXFIO step-up dosing schedule to reduce risk of CRS and monitor patients following administration of ELREXFIO accordingly [see *Dosage and Administration (2.2, 2.5)*]. Administer pre-treatment medications prior to each dose in the step-up dosing schedule to reduce risk of CRS [see *Dosage and Administration (2.3)*].

Counsel patients to seek medical attention should signs or symptoms of CRS occur. At the first sign of CRS, evaluate patients immediately for hospitalization. Manage CRS according to the recommendations and consider further management per current practice guidelines. Withhold or permanently discontinue ELREXFIO based on severity [see *Dosage and Administration (2.5)*].

ELREXFIO is available only through a restricted program under a REMS [see *Warnings and Precautions (5.3)*].

5.2 Neurologic Toxicity, Including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS)

ELREXFIO can cause serious or life-threatening neurologic toxicity, including ICANS [see *Adverse Reactions (6.1)*].

In the clinical trial, neurologic toxicity occurred in 59% of patients who received ELREXFIO at the recommended dosing schedule [see *Dosage and Administration (2.2)*], with Grade 3 or 4 neurologic toxicity occurring in 7% of patients. Neurologic toxicities included headache (18%), encephalopathy (15%), motor dysfunction (13%), sensory neuropathy (13%), and Guillain-Barré Syndrome (0.5%).

In the clinical trial, ICANS occurred in 3.3% of patients who received ELREXFIO at the recommended dosing schedule [see *Dosage and Administration (2.2)*]. Most patients had ICANS after the first step-up dose (2.7%), 1 (0.5%) patient had ICANS after the second step-up dose and 1 (0.5%) patient had ICANS after subsequent dose(s). Recurrent ICANS occurred in 1.1% of patients. The median time to onset was 3 (range: 1 to 4) days after the most recent dose, with a median duration of 2 (range: 1 to 18) days. The most frequent clinical manifestations of ICANS included a depressed level of consciousness and Grade 1 or Grade 2 Immune Effector Cell-Associated Encephalopathy (ICE) scores. The onset of ICANS can be concurrent with CRS, following resolution of CRS, or in the absence of CRS.

Counsel patients to seek medical attention should signs or symptoms of neurologic toxicity occur. Monitor patients for signs and symptoms of neurologic toxicities during treatment with ELREXFIO. At the first sign of neurologic toxicity, including ICANS, evaluate and treat patients immediately based on severity. Withhold or permanently discontinue ELREXFIO based on severity per recommendations [see *Dosage and Administration (2.5)*] and consider further management per current practice guidelines.

Due to the potential for neurologic toxicity including ICANS, patients receiving ELREXFIO are at risk of depressed level of consciousness. Advise patients not to drive or operate heavy or potentially dangerous machinery for 48 hours after completing each of the 2 step-up doses and the first treatment dose within the ELREXFIO step-up dosing schedule and in the event of new onset of any neurological toxicity symptoms until symptoms resolve [see *Dosage and Administration (2.2)*].

ELREXFIO is available only through a restricted program under a REMS [see *Warnings and Precautions (5.3)*].

5.3 ELREXFIO REMS

ELREXFIO is available only through a restricted program under a REMS called the ELREXFIO REMS because of the risks of CRS and neurologic toxicity, including ICANS [see *Warnings and Precautions (5.1, 5.2)*].

Notable requirements of the ELREXFIO REMS include the following:

- Prescribers must be certified with the program by enrolling and completing training.
- Prescribers must counsel patients receiving ELREXFIO about the risk of CRS and neurologic toxicity, including ICANS, and provide patients with ELREXFIO Patient Wallet Card.
- Pharmacies and healthcare settings that dispense ELREXFIO must be certified with the ELREXFIO REMS program and must verify prescribers are certified through the ELREXFIO REMS program.
- Wholesalers and distributors must only distribute ELREXFIO to certified pharmacies or healthcare settings.

Further information about the ELREXFIO REMS program is available at www.ELREXFIOREMS.com or by telephone at 1-844-923-7845.

5.4 Infections

ELREXFIO can cause severe, life-threatening, or fatal infections. In the clinical trial, in patients who received ELREXFIO according to the recommended dosing schedule, serious infections, including opportunistic infections, occurred in 42% of patients, with Grade 3 or 4 infections in 31%, and fatal infections in 7%. The most common serious infections reported ($\geq 5\%$) were pneumonia and sepsis [see *Adverse Reactions (6.1)*].

Do not initiate treatment with ELREXFIO in patients with active infections. Monitor patients for signs and symptoms of infection prior to and during treatment with ELREXFIO and treat appropriately. Withhold or permanently discontinue ELREXFIO based on severity [see *Dosage and Administration (2.5)*]. Administer prophylactic antimicrobial and anti-viral medications according to current practice guidelines. Consider treatment with subcutaneous or intravenous immunoglobulin (IVIG) as appropriate.

5.5 Neutropenia

ELREXFIO can cause neutropenia and febrile neutropenia. In patients who received ELREXFIO at the recommended dose in the clinical trial, decreased neutrophils occurred in 62% of patients, with Grade 3 or 4 decreased neutrophils in 51%. Febrile neutropenia occurred in 2.2% of patients [see *Adverse Reactions (6.1)*].

Monitor complete blood cell counts at baseline and periodically during treatment. Provide supportive care according to current practice guidelines. Monitor patients with neutropenia for signs of infection. Withhold ELREXFIO based on severity [see *Dosage and Administration (2.5)*].

5.6 Hepatotoxicity

ELREXFIO can cause hepatotoxicity. In the clinical trial, elevated ALT occurred in 36% of patients, with Grade 3 or 4 ALT elevation occurring in 3.8%; elevated AST occurred in 40% of patients, with Grade 3 or 4 AST elevation occurring in 6%. Grade 3 or 4 total bilirubin elevations occurred in 0.5% of patients [see *Adverse Reactions (6.1)*]. Liver enzyme elevation can occur with or without concurrent CRS.

Monitor liver enzymes and bilirubin at baseline and during treatment as clinically indicated. Withhold ELREXFIO or consider permanent discontinuation of ELREXFIO based on severity [see *Dosage and Administration (2.5)*].

5.7 Embryo-Fetal Toxicity

Based on its mechanism of action, ELREXFIO may cause fetal harm when administered to a pregnant woman. Advise pregnant women of the potential risk to the fetus. Advise females of reproductive potential to use effective contraception during treatment with ELREXFIO and for 4 months after the last dose [see *Use in Specific Populations (8.1, 8.3)*].

6 ADVERSE REACTIONS

The following adverse reactions are discussed elsewhere in labeling:

- Cytokine Release Syndrome [see *Warnings and Precautions (5.1)*].
- Neurologic Toxicity, Including ICANS [see *Warnings and Precautions (5.2)*].
- Infections [see *Warnings and Precautions (5.4)*].
- Neutropenia [see *Warnings and Precautions (5.5)*].
- Hepatotoxicity [see *Warnings and Precautions (5.6)*].

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

Relapsed/Refractory Multiple Myeloma

MagnetisMM-3

The safety of ELREXFIO was evaluated in MagnetisMM-3 [see *Clinical Studies (14)*]. The safety population described (n = 183) includes patients who received the recommended dosage regimen of 12 mg subcutaneously on Day 1, 32 mg on Day 4, and 76 mg once weekly starting on Day 8. Among patients who received ELREXFIO, 42% were exposed for 6 months or longer and 9% were exposed for one year or longer.

The median age of patients who received ELREXFIO was 68 years (range: 36 to 88 years); 48% were female; 61% were White, 10% were Hispanic/Latino, 9% were Asian, and 6% were Black or African American.

Serious adverse reactions occurred in 68% of patients who received ELREXFIO at the recommended dosing schedule. Serious adverse reactions in >2% of patients included pneumonia (25%), sepsis (13%), CRS (13%), upper respiratory tract infection (4.4%), acute kidney injury (3.8%), urinary tract infection (3.3%), COVID-19 (3.3%), encephalopathy (3.3%), pyrexia (2.2%), and febrile neutropenia (2.2%). Fatal adverse reactions occurred in 10% of patients including pneumonia (3.3%), sepsis (2.7%), acute respiratory distress syndrome (0.5%), cardio-respiratory arrest (0.5%), cardiogenic shock (0.5%), cardiopulmonary failure (0.5%), COVID-19 (0.5%), failure to thrive (0.5%), and pulmonary embolism (0.5%).

Permanent discontinuations of ELREXFIO due to an adverse reaction occurred in 17% of patients. Adverse reactions which resulted in permanent discontinuation of ELREXFIO in >2% of patients included septic shock (2.2%).

Dosage interruptions of ELREXFIO due to an adverse reaction occurred in 73% of patients. Adverse reactions which resulted in dose interruptions of ELREXFIO in >5% of patients included neutropenia, pneumonia, COVID-19, upper respiratory tract infection, thrombocytopenia, and anemia.

The most common adverse reactions (≥20%) were CRS, fatigue, injection site reaction, diarrhea, upper respiratory tract infection, musculoskeletal pain, pneumonia, decreased appetite, rash, cough, nausea, and pyrexia. The most common Grade 3 to 4 laboratory abnormalities (≥30%) were decreased lymphocytes, decreased neutrophils, decreased hemoglobin, decreased white blood cells, and decreased platelets.

Table 8 summarizes adverse reactions in MagnetisMM-3.

Table 8. Adverse Reactions (≥10%) in Patients with Relapsed or Refractory Multiple Myeloma Who Received ELREXFIO in MagnetisMM-3

| System Organ Class Preferred Term | ELREXFIO N = 183 | |
|---|---------------------|---------------------|
| | All Grades (%) | Grade 3 or 4 (%) |
| Immune system disorders | | |
| Cytokine release syndrome | 58 | 0.5 [#] |
| Hypogammaglobulinemia* | 13 | 2.2 [#] |
| General disorders and site administration conditions | | |
| Fatigue* | 43 | 6 [#] |
| Injection site reaction* | 37 | 0 |
| Pyrexia | 21 | 2.7 [#] |
| Edema* | 18 | 1.1 [#] |
| Gastrointestinal disorders | | |
| Diarrhea | 36 | 1.1 [#] |
| Nausea | 22 | 0 |
| Constipation | 15 | 0 |
| Vomiting | 14 | 0 |
| Infections | | |
| Upper respiratory tract infection* | 34 | 4.9 |

Table 8. Adverse Reactions (≥10%) in Patients with Relapsed or Refractory Multiple Myeloma Who Received ELREXFIO in MagnetisMM-3

| System Organ Class Preferred Term | ELREXFIO N = 183 | |
|--|---------------------|------------------|
| Pneumonia ^a | 32 | 19 |
| Sepsis ^b | 15 | 11 |
| Urinary tract infection [*] | 12 | 4.4 [#] |
| Musculoskeletal and connective tissue disorders | | |
| Musculoskeletal pain [*] | 34 | 2.7 [#] |
| Metabolism and nutrition disorders | | |
| Decreased appetite | 26 | 1.1 [#] |
| Skin and Subcutaneous Tissue disorders | | |
| Rash ^c | 25 | 0 |
| Dry skin | 13 | 0 |
| Skin exfoliation [*] | 10 | 0 |
| Respiratory, thoracic and mediastinal disorders | | |
| Cough [*] | 24 | 0 |
| Dyspnea [*] | 15 | 3.3 [#] |
| Nervous system disorders | | |
| Headache | 18 | 0.5 |
| Encephalopathy ^d | 15 | 2.7 |
| Sensory neuropathy ^e | 13 | 0.5 [#] |
| Motor dysfunction ^f | 13 | 2.2 [#] |
| Cardiac disorders | | |
| Cardiac arrhythmia [*] | 16 | 2.2 |
| Vascular disorders | | |
| Hemorrhage [*] | 13 | 1.6 |
| Psychiatric disorders | | |
| Insomnia | 13 | 0 |
| Injury, poisoning and procedural complications | | |
| Fall | 10 | 0.5 [#] |

Adverse reactions were graded based on CTCAE Version 5.0, with the exception of CRS, which was graded based on the ASTCT 2019 criteria.

* Includes other related terms.

Only grade 3 adverse reactions occurred.

- a. Pneumonia includes COVID-19 pneumonia, lower respiratory tract infection, lower respiratory tract infection viral, pneumocystis jirovecii pneumonia, pneumonia, pneumonia adenoviral, pneumonia bacterial, pneumonia cytomegaloviral, pneumonia fungal, pneumonia influenzal, pneumonia pseudomonal, pneumonia viral.
- b. Sepsis includes bacteremia, device related bacteremia, device related sepsis, escherichia bacteremia, escherichia sepsis, klebsiella sepsis, pseudomonal sepsis, sepsis, septic shock, staphylococcal bacteremia, staphylococcal sepsis, streptococcal sepsis, urosepsis.
- c. Rash includes erythema, palmar-plantar erythrodysesthesia syndrome, rash, rash erythematous, rash macular, rash maculo-papular, rash pustular, symmetrical drug-related intertriginous and flexural exanthema.
- d. Encephalopathy includes agitation, altered state of consciousness, cognitive disorder, confusional state, delirium, depressed level of consciousness, disorientation, hallucination, lethargy, memory impairment, mental status changes, metabolic encephalopathy, somnolence, toxic encephalopathy.
- e. Sensory neuropathy includes burning sensation, dysesthesia, hypoesthesia, neuropathy peripheral, paresthesia, parosmia, peripheral sensorimotor neuropathy, peripheral sensory neuropathy, polyneuropathy, sensory loss.
- f. Motor dysfunction includes ataxia, balance disorder, gait disturbance, motor dysfunction, muscle contracture, muscle spasms, muscular weakness, peripheral motor neuropathy, peroneal nerve palsy, tremor.

Clinically relevant adverse reactions in <10% of patients who received ELREXFIO included ICANS, febrile neutropenia, Guillain-Barré syndrome, abdominal pain, acute kidney injury, COVID-19, cardiac failure, congestion, and thrombosis.

Table 9 summarizes laboratory abnormalities in MagnetisMM-3.

Table 9. Select Laboratory Abnormalities (≥30%) That Worsened from Baseline in Patients with Relapsed or Refractory Multiple Myeloma Who Received ELREXFIO in MagnetisMM-3^a

| Laboratory Abnormality | ELREXFIO ^b | |
|--------------------------------|-----------------------|------------------|
| | All Grades (%) | Grade 3 or 4 (%) |
| Hematology | | |
| Lymphocyte count decreased | 91 | 84 |
| White blood cell decreased | 69 | 40 |
| Hemoglobin decreased | 68 | 43 |
| Neutrophil count decreased | 62 | 51 |
| Platelet count decreased | 61 | 32 |
| Chemistry | | |
| Albumin decreased | 55 | 6 |
| AST increase | 40 | 6 |
| Creatinine increased | 38 | 3.3 |
| Potassium decreased | 36 | 8 |
| ALT increase | 36 | 3.8 |
| Alkaline phosphatase increased | 34 | 1.1 |
| Creatinine clearance decreased | 32 | 10 |

- a. Laboratory tests were graded according to NCI-CTCAE Version 5.0.
- b. The denominator used to calculate the rate varied from 181 to 183 based on the number of patients with a baseline value and at least one post-treatment value.

7 DRUG INTERACTIONS

For certain CYP substrates, minimal changes in the concentration may lead to serious adverse reactions. Monitor for toxicity or drug concentrations of such CYP substrates when co-administered with ELREXFIO.

ELREXFIO causes release of cytokines [see *Clinical Pharmacology (12.2)*] that may suppress activity of cytochrome P450 (CYP) enzymes, resulting in increased exposure of CYP substrates. Increased exposure of CYP substrates is more likely to occur after the first dose of ELREXFIO Day 1 and up to 14 days after the 32 mg dose on Day 4 and during and after CRS [see *Warnings and Precautions (5.1)*].

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Based on the mechanism of action, ELREXFIO may cause fetal harm when administered to a pregnant woman [see *Clinical Pharmacology (12.1)*]. There are no available data on the use of ELREXFIO in pregnant women to evaluate for a drug associated risk. No animal reproductive or developmental toxicity studies have been conducted with ELREXFIO. Elranatamab-bcmm causes T-cell activation and cytokine release; immune activation may compromise pregnancy maintenance. In addition, based on the finding of B-cell depletion in non-pregnant animals, elranatamab-bcmm can cause B-cell lymphocytopenia in infants exposed to elranatamab-bcmm in-utero. Human immunoglobulin (IgG) is known to cross the placenta after the first trimester of pregnancy; therefore, elranatamab-bcmm has the potential to be transmitted from the mother to the developing fetus. Advise women of the potential risk to the fetus.

ELREXFIO is associated with hypogammaglobulinemia, therefore, assessment of immunoglobulin levels in newborns of mothers treated with ELREXFIO should be considered.

In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

8.2 Lactation

Risk Summary

There are no data on the presence of elranatamab-bcmm in human milk, the effects on the breastfed child, or the effects on milk production. Maternal IgG is known to be present in human milk.

Because of the potential for serious adverse reactions in a breastfed child, advise women not to breastfeed during treatment with ELREXFIO and for 4 months after the last dose.

8.3 Females and Males of Reproductive Potential

ELREXFIO may cause fetal harm when administered to a pregnant woman [*see Use in Specific Populations (8.1)*].

Pregnancy Testing

Verify the pregnancy status of females of reproductive potential prior to initiating treatment with ELREXFIO.

Contraception

Females

Advise females of reproductive potential to use effective contraception during treatment and for 4 months after the last dose of ELREXFIO.

8.4 Pediatric Use

The safety and effectiveness of ELREXFIO in pediatric patients have not been established.

8.5 Geriatric Use

Of the 183 patients with relapsed or refractory multiple myeloma treated with ELREXFIO in MagnetisMM-3 at the recommended dosage, 62% were 65 years of age or older, and 19% were 75 years of age or older. No overall differences in safety or effectiveness were observed in patients 65-74 years of age compared to younger patients. Clinical studies did not include sufficient numbers of patients 75 years of age or older to determine whether they respond differently from younger patients.

11 DESCRIPTION

Elranatamab-bcmm is a bispecific B-cell maturation antigen (BCMA)-directed CD3 T-cell engager. It is a bispecific, humanized immunoglobulin 2-alanine (IgG2Δa) kappa antibody derived from two monoclonal antibodies (mAbs), an anti-BCMA mAb and an anti-CD3 mAb. Each of these mAbs contributes one distinct heavy (H) chain and one distinct light (L) chain to the bispecific elranatamab-bcmm. The resulting 4-chain bispecific antibody is covalently linked via five inter-chain disulfide bonds. Elranatamab-bcmm is produced using two recombinant Chinese hamster ovary (CHO) cell lines, one that contains the DNA encoding the sequence for anti-BCMA monoclonal antibody (mAb) and one that contains the sequence for anti-CD3 mAb, which are grown separately in suspension culture using chemically-defined (CD), animal-derived component-free (ACF) media. The molecular weight of elranatamab-bcmm is approximately 148.5 kDa.

ELREXFIO™ (elranatamab-bcmm) injection is a sterile, preservative-free, clear to slightly opalescent, and colorless to pale brown liquid solution for subcutaneous administration. ELREXFIO (elranatamab-bcmm) is supplied at a concentration of 40 mg/mL in either 76 mg/1.9 mL or 44 mg/1.1 mL single-dose vials. Each mL of solution contains 40 mg elranatamab-bcmm, edetate disodium (0.045 mg), histidine (1.12 mg), L-histidine hydrochloride monohydrate (2.67 mg), polysorbate 80 (0.2 mg), sucrose (85 mg) and Water for Injection. The pH is 5.8.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Elranatamab-bcmm is a bispecific B-cell maturation antigen (BCMA)-directed T-cell engaging antibody that binds BCMA on plasma cells, plasmablasts, and multiple myeloma cells and CD3 on T-cells leading to cytolysis of the BCMA-expressing cells. Elranatamab-bcmm activated T-cells, caused proinflammatory cytokine release, and resulted in multiple myeloma cell lysis.

12.2 Pharmacodynamics

Cytokine Concentrations

Transient elevation of circulating cytokines IL-2, IL-6, IL-8, IL-10, TNF- α , and IFN- γ was observed at dosage levels of 30 μ g/kg (0.03 times the approved recommended dosage) and above. After administration of the approved recommended dosage of ELREXFIO, the highest elevation of cytokines was generally observed within 72 hours after first elranatamab-bcmm dose at 12 mg on Day 1, and generally returned to baseline prior to the administration of the first full dose 76 mg on Day 8.

12.3 Pharmacokinetics

Pharmacokinetic parameters are presented as geometric mean (coefficient of variation [CV]%) and are based upon subcutaneously administered unless otherwise specified.

Elranatamab-bcmm exhibits dose proportional pharmacokinetics over dose range from 6 to 76 mg (0.079 to 1 times the approved recommended dosage). Elranatamab-bcmm maximum concentration [33.6 mcg/mL (48%)] is achieved at the end of weekly dosing regimen (i.e., at week 24 of 76 mg weekly dosing). Pharmacokinetic exposures are summarized for the recommended dosage of ELREXFIO in Table 10.

Table 10. Pharmacokinetic Parameters of Elranatamab-bcmm in Subjects with Relapsed or Refractory Multiple Myeloma

| Timepoint | Parameters | | |
|---|------------------------------|------------------------------|---------------------------------|
| | C _{avg} (mcg/mL) | C _{max} (mcg/mL) | C _{trough} (mcg/mL) |
| First full 76 mg dose | 3.1 (94%) | 3.8 (94%) | 3.3 (102%) |
| End of weekly dose (week 24) ^a | 32.7 (49%) | 33.6 (48%) | 31.2 (50%) |
| Steady state (biweekly dosing) ^{a,b} | 18.4 (57%) | 20.1 (55%) | 15.9 (64%) |

a. In patients who have achieved a response.

b. Steady state exposure of elranatamab biweekly dose is approximated at week 48.

Absorption

The mean bioavailability of elranatamab-bcmm was 56.2% when administered subcutaneously. The median (min, max) T_{max} after elranatamab SC administration was 7 (3 to 7) days.

Distribution

The steady state volume of distribution of elranatamab-bcmm was 7.76 L (33%).

Elimination

The half-life of elranatamab-bcmm is 22 (64%) days at the 76 mg dosage, with clearance of 0.324 L/day (100%) following 24 weeks dosing.

Metabolism

Elranatamab-bcmm is expected to be metabolized into small peptides by catabolic pathways.

Specific Populations

No clinically significant differences in the pharmacokinetics of elranatamab-bcmm were observed based on age (36 to 89 years), sex, race (White, Asian, or Black), body weight (37 to 160 kg), mild or moderate renal impairment (estimated glomerular filtration rate [eGFR] by Modification of Diet in Renal Disease [MDRD] method: 30 to 89 mL/min), or mild hepatic impairment (total bilirubin 1 to $\leq 1.5 \times$ ULN or any AST greater than ULN).

The effects of severe renal impairment (eGFR 15 to 29 mL/min), end-stage renal disease (eGFR < 15 mL/min), or moderate to severe hepatic impairment (total bilirubin > 1.5 times ULN and any AST) on the PK of elranatamab-bcmm are unknown.

12.6 Immunogenicity

The observed incidence of anti-drug antibodies is highly dependent on the sensitivity and specificity of the assay. Differences in assay methods preclude meaningful comparisons of the incidence of anti-drug antibodies in the studies described below with the incidence of anti-drug antibodies in other studies, including those of elranatamab-bcmm or of other elranatamab products.

In the MagnetisMM-3 study, of the 168 participant who received recommended step-up and full dosage of ELREXFIO for up to 24 month and are evaluable for presence of ADA against elranatamab-bcmm, 8.9% (15/168) of patients tested positive for anti-elranatamab-bcmm-antibodies. Among the 15 patients who tested positive for ADAs, 60% (9/15) tested positive for neutralizing antibodies against elranatamab-bcmm. The effect of these antibodies on the pharmacokinetics, pharmacodynamics, safety, and/or effectiveness of ELREXFIO products is unknown.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

No carcinogenicity or genotoxicity studies have been conducted with elranatamab-bcmm.

No animal studies have been performed to evaluate the effects of elranatamab-bcmm on fertility.

14 CLINICAL STUDIES

14.1 Relapsed or Refractory Multiple Myeloma

The efficacy of ELREXFIO monotherapy was evaluated in patients with relapsed or refractory multiple myeloma in an open-label, single arm, multi-center study (MagnetisMM-3, NCT04649359). The study included patients who were refractory to at least one proteasome inhibitor (PI), one immunomodulatory agent (IMiD), and one anti-CD38 monoclonal antibody. MagnetisMM-3 included 123 patients naïve to prior BCMA-directed therapy (pivotal Cohort A) and 64 patients with prior BCMA-directed antibody drug conjugate (ADC) or chimeric antigen receptor (CAR) T-cell therapy (supportive Cohort B). Patients had measurable disease by International Myeloma Working Group (IMWG) criteria at enrollment. The study included patients with an Eastern Cooperative Oncology Group (ECOG) score of ≤ 2 , adequate baseline bone marrow (absolute neutrophil count $\geq 1.0 \times 10^9/L$, platelet count $\geq 25 \times 10^9/L$, hemoglobin level ≥ 8 g/dL), renal (CrCL ≥ 30 mL/min), and hepatic (AST and ALT $\leq 2.5 \times$ ULN, total bilirubin $\leq 2 \times$ ULN) function, and left-ventricular ejection fraction $\geq 40\%$. Patients with a stem cell transplant within 12 weeks prior to enrollment and active infections were excluded from the study.

Eligible patients received subcutaneous administration of ELREXFIO at step-up doses of 12 mg on Day 1 and 32 mg on Day 4 of treatment, followed by the first treatment dose of ELREXFIO (76 mg) on Day 8 of treatment. Thereafter, patients received 76 mg once weekly. After 24 weeks, in patients who achieved an IMWG response category of partial response or better with responses persisting for at least 2 months, the dose interval was changed from every week to every 2 weeks.

The 123 patients enrolled in pivotal Cohort A had received a median of 5 prior lines of therapy (range: 2 to 22). Ninety-seven patients who were not exposed to prior BCMA-directed therapy and received at least four prior lines of therapy comprised the efficacy population. Among the 97 patients in the efficacy population, the median age was 69

(range: 46 to 89) years with 18.6% of patients ≥ 75 years of age. Forty percent were female; 59.8% were White, 13.4% were Asian, 7.2% were Hispanic/Latino, 5.2% were Black or African American. Disease stage (R-ISS) at study entry was 20.6% in Stage I, 53.6% in Stage II, and 17.5% in Stage III. The median time since initial diagnosis of multiple myeloma to enrollment was 79.6 (range: 16 to 228) months. 96.9% were triple-class refractory, and 94.8% were refractory to their last line of therapy. 69.1% received prior autologous stem cell transplantation, and 7.2% received prior allogenic stem cell transplantation. High-risk cytogenetics [t(4;14), t(14;16), or del(17p)] were present in 22.7% of patients. 34.0% of patients had extramedullary disease at baseline by BICR.

Efficacy was based on response rate and duration of response (DOR), as assessed by BICR based on IMWG criteria. Efficacy results from BCMA-directed therapy naïve patients are shown in Table 11.

The median (range) time to first response (TTR) was 1.22 (0.9 to 6.5) months. With a median follow-up of 11.1 months (95% CI: 10.6, 12.0) among responders, the DOR rate at 6 months was 90.4% (95% CI: 78.4%, 95.9%) and at 9 months was 82.3% (95% CI: 67.1%, 90.9%).

Table 11. Efficacy Results from BCMA-directed Therapy Naïve Patients

| | N = 97 |
|--|----------------------------|
| Objective Response Rate (ORR: sCR+CR+VGPR+PR), n (%) (95% CI) | 56 (57.7%) (47.3, 67.7) |
| Complete response (CR) or better ^a | 25 (25.8%) |
| Very good partial response (VGPR) | 25 (25.8%) |
| Partial response (PR) | 6 (6.2%) |
| Duration of Response (DOR) (months) | |
| Median (95% CI) | NR (12.0, NE) |

Abbreviations: CI = Confidence interval; NR = Not reached; NE = Not estimable.

a. Complete response or better = Stringent complete response (sCR) + complete response (CR).

Among the 64 patients enrolled in Cohort B who previously received a PI, an IMiD, an anti-CD38 monoclonal antibody, and a BCMA-directed therapy, 63 patients received at least four prior lines of therapy. Patients had received a median of 8 prior lines of therapy (range: 4 to 19); 73% and 32% received prior BCMA-directed ADC and CAR T-cell therapy, respectively.

Confirmed ORR by BICR was 33.3% (95% CI: 22.0, 46.3). After a median (95% CI) follow-up of 10.2 (9.9, 11.0) months among responders, median DOR was not reached (95% CI: NE, NE) and the DOR rate at 9 months was 84.3% (95% CI: 58.7, 94.7).

16 HOW SUPPLIED/STORAGE AND HANDLING

How Supplied

ELREXFIO™ (elranatamab-bcmm) injection is a sterile, preservative-free, clear to slightly opalescent, and colorless to pale brown liquid solution supplied as follows:

- One 76 mg/1.9 mL (40 mg/mL) single-dose vial in a carton. NDC: 0069-4494-02
- One 44 mg/1.1 mL (40 mg/mL) single-dose vial in a carton. NDC: 0069-2522-02

ELREXFIO is supplied in a single-dose glass vial sealed with a rubber stopper (not made of natural rubber latex) and an aluminum seal with a flip-off cap.

Storage and Handling

Store refrigerated at 2 °C to 8 °C (36 °F to 46 °F) in the original carton until time of use to protect from light. Do not freeze or shake the vial or carton.

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved patient labeling (Medication Guide).

Cytokine Release Syndrome (CRS)

Discuss the signs and symptoms associated with CRS, including fever, hypoxia, chills, hypotension, tachycardia, and elevated liver enzymes. Advise patients to immediately contact their healthcare provider if they experience any signs or symptoms of CRS. Advise patients that they will be hospitalized for 48 hours after administration of the first step-up dose, and for 24 hours after administration of the second step-up dose [see *Dosage and Administration (2.5)*, *Warnings and Precautions (5.1)*].

Neurologic Toxicity, Including Immune Effector Cell-associated Neurotoxicity Syndrome (ICANS)

Discuss the signs and symptoms associated with neurologic toxicity, including ICANS, including headache, encephalopathy, motor dysfunction, sensory neuropathy, and Guillain-Barré Syndrome. Advise patients to immediately contact their healthcare provider if they experience any signs or symptoms of neurologic toxicity. Advise patients to refrain from driving or operating heavy or potentially dangerous machinery for 48 hours after completing each of the 2 step-up doses and the first treatment dose within the ELREXFIO step-up dosing schedule and in the event of new onset of any neurological toxicity symptoms until symptoms resolve [see *Dosage and Administration (2.5)*, *Warnings and Precautions (5.2)*].

ELREXFIO REMS

ELREXFIO is available only through a restricted program called ELREXFIO REMS. Inform patients that they will be given an ELREXFIO Patient Wallet Card that they should carry with them at all times and show to all of their healthcare providers. This card describes signs and symptoms of CRS and neurologic toxicity, including ICANS which, if experienced, should prompt the patient to immediately seek medical attention [see *Warnings and Precautions (5.3)*].

Infections

Discuss the signs and symptoms of infection [see *Dosage and Administration (2.5)*, *Warnings and Precautions (5.4)*].

Neutropenia

Discuss the signs and symptoms associated with neutropenia and febrile neutropenia [see *Dosage and Administration (2.5)*, *Warnings and Precautions (5.5)*].

Hepatotoxicity

Advise patients that liver enzyme elevations may occur and that they should report symptoms that may indicate liver toxicity, including fatigue, anorexia, right upper abdominal discomfort, dark urine, or jaundice [see *Warnings and Precautions (5.6)*].

Embryo-Fetal Toxicity

Advise pregnant women and females of reproductive potential of the potential risk to a fetus. Advise females of reproductive potential to inform their healthcare provider if they are pregnant or become pregnant. Advise females of reproductive potential to use effective contraception during treatment with ELREXFIO and for 4 months after the last dose [see *Warnings and Precautions (5.7)*, *Use in Specific Populations (8.1, 8.3)*].

Lactation

Advise women not to breastfeed during treatment with ELREXFIO and for 4 months after the last dose [see *Use in Specific Populations (8.2)*].

Manufactured by:

Pfizer Inc.

NY, NY 10001

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New York, NY 10001

LAB-1518-0.8

MEDICATION GUIDE
ELREXFIO™ (el-reks-fe-o)
(elranatamab-bcmm)
injection, for subcutaneous use

What is the most important information I should know about ELREXFIO?

ELREXFIO may cause serious side effects, including:

- **Cytokine Release Syndrome (CRS).** CRS is common during treatment with ELREXFIO and can also be serious, life-threatening, or can lead to death. Tell your healthcare provider or get medical help right away if you develop any signs or symptoms of CRS, including:
 - fever of 100.4 °F (38 °C) or higher
 - headache
 - trouble breathing
 - increased liver enzymes in your blood. See **“What are the possible side effects of ELREXFIO?”** for more information about the signs and symptoms of liver problems.
 - chills
 - dizziness or light-headedness
 - fast heartbeat

Due to the risk of CRS, you will receive ELREXFIO on a “step-up dosing schedule” and should be hospitalized for 48 hours after the first “step-up” dose and for 24 hours after the second “step-up” dose of ELREXFIO.

- During the step-up dosing schedule:
 - for your first dose, you will receive a smaller “step-up” dose of ELREXFIO on Day 1 of your treatment
 - for your second dose, you will receive a larger “step-up” dose of ELREXFIO, which is usually given on Day 4 of your treatment
 - for your third dose, you will receive the first full “treatment” dose of ELREXFIO, which is usually given on Day 8 of your treatment
- If your dose of ELREXFIO is delayed for any reason, you may need to repeat the step-up dosing schedule.
- Before each dose of ELREXFIO you receive during the step-up dosing schedule, you will receive medicines to help reduce your risk of CRS. Your healthcare provider will decide if you need to receive medicines to help reduce your risk of CRS with future doses.
- See **“How will I receive ELREXFIO?”** for more information about how you will receive ELREXFIO.
- **Neurologic problems.** ELREXFIO can cause neurologic problems that can be serious or life-threatening. Tell your healthcare provider or get medical help right away if you develop any signs or symptoms of neurologic problems, including:
 - headache
 - problems walking, muscle weakness, shaking (tremors), loss of balance, or muscle spasms
 - agitation, trouble staying awake, confusion or disorientation, seeing or hearing things that are not real (hallucinations)
 - numbness and tingling (feeling like “pins and needles”)
 - trouble speaking, thinking, remembering things, paying attention, or understanding things
 - burning, throbbing, or stabbing pain
 - changes in your handwriting

- **ELREXFIO is available only through the ELREXFIO Risk Evaluation and Mitigation Strategy (REMS) due to the risk of CRS and neurologic problems.** You will receive an ELREXFIO Patient Wallet Card from your healthcare provider. **Carry the ELREXFIO Patient Wallet Card with you at all times and show it to all of your healthcare providers.** The ELREXFIO Patient Wallet Card lists symptoms of CRS and neurologic problems. **Get medical help right away if you develop any of the symptoms listed on the ELREXFIO Patient Wallet Card.** You may need to be treated in a hospital.

Your healthcare provider will monitor you for signs and symptoms of CRS and neurologic problems during treatment with ELREXFIO, as well as other side effects, and will treat you if needed. Your healthcare provider may temporarily stop or completely stop your treatment with ELREXFIO if you develop CRS, neurologic problems, or any other side effects that are severe.

If you have any questions about ELREXFIO, ask your healthcare provider.

See “What are possible side effects of ELREXFIO?” for more information about side effects.

What is ELREXFIO?

ELREXFIO is a prescription medicine used to treat adults with multiple myeloma who:

- have already received at least 4 treatment regimens, including a proteasome inhibitor, an immunomodulatory agent and an anti-CD38 monoclonal antibody to treat their multiple myeloma, **and**
- their cancer has come back or did not respond to prior treatment.

It is not known if ELREXFIO is safe and effective in children.

Before receiving ELREXFIO, tell your healthcare provider about all of your medical conditions, including if you:

- have an infection.
- are pregnant or plan to become pregnant. ELREXFIO may harm your unborn baby.
Females who are able to become pregnant:
 - Your healthcare provider should do a pregnancy test before you start treatment with ELREXFIO.
 - You should use effective birth control (contraception) during treatment and for 4 months after your last dose of ELREXFIO.
 - Tell your healthcare provider right away if you become pregnant or think that you may be pregnant during treatment with ELREXFIO.
- are breastfeeding or plan to breastfeed. It is not known if ELREXFIO passes into your breast milk. Do not breastfeed during treatment and for 4 months after your last dose of ELREXFIO.

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 ELREXFIO?

- ELREXFIO will be given to you by your healthcare provider as an injection under your skin (subcutaneous injection), usually in your stomach-area (abdomen). Your thigh or another area of your body may also be used.
- See **“What is the most important information I should know about ELREXFIO?”** for more information about how you will receive ELREXFIO.
- After you receive your first full “treatment” dose, ELREXFIO 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.

If you miss any appointments, call your healthcare provider as soon as possible to reschedule your appointment. It is important for you to be monitored closely for side effects during treatment with ELREXFIO.

What should I avoid while receiving ELREXFIO?

Do not drive, operate heavy or potentially dangerous machinery, or do other dangerous activities during treatment with ELREXFIO:

- for 48 hours after completing each of the 2 doses of ELREXFIO that are part of the “step-up dosing schedule” and your first full treatment dose, **and**
- at any time during treatment with ELREXFIO if you develop any new neurologic symptoms such as dizziness, confusion, shaking (tremors), sleepiness, or any other symptom that impairs consciousness, until the symptoms go away.

See **“What is the most important information I should know about ELREXFIO?”** for more information about signs and symptoms of neurologic problems.

What are the possible side effects of ELREXFIO?

ELREXFIO may cause serious side effects, including:

- See **“What is the most important information I should know about ELREXFIO?”**
- **Infections.** Upper respiratory tract infections and pneumonia are common during treatment with ELREXFIO. ELREXFIO can cause bacterial and viral infections that are severe, life-threatening, or that may lead to death.
 - Your healthcare provider may prescribe medicines for you to help prevent infections and treat you as needed if you develop an infection during treatment with ELREXFIO.
 - Tell your healthcare provider right away if you develop any signs or symptoms of an infection during treatment with ELREXFIO, including:

| | |
|---------------------------------------|------------------------------------|
| ▪ fever of 100.4 °F (38 °C) or higher | ▪ chest pain |
| ▪ chills | ▪ sore throat |
| ▪ cough | ▪ pain during urination |
| ▪ shortness of breath | ▪ feeling weak or generally unwell |
- **Decreased white blood cell counts.** Decreased white blood cell counts are common during treatment with ELREXFIO and can also be severe. Fever can happen with low white blood cell counts and may be a sign that you have an infection. Your healthcare provider will treat you as needed.
- **Liver problems.** ELREXFIO can cause increased liver enzymes and bilirubin in your blood. These increases can happen with or without you also having CRS. Tell your healthcare provider if you develop any of the following signs or symptoms of liver problems:

- tiredness
- loss of appetite
- pain in your right upper stomach-area (abdomen)
- dark urine
- yellowing of your skin or the white part of your eyes

Your healthcare provider will check your blood and monitor you for signs and symptoms of these serious side effects before you start and during treatment with ELREXFIO and may temporarily or completely stop treatment with ELREXFIO if you develop certain side effects.

The most common side effects of ELREXFIO include:

- tiredness
- injection site reaction, such as redness, itching, pain, bruising, rash, swelling, tenderness
- diarrhea
- muscle and bone pain
- decreased appetite
- rash
- cough
- nausea
- fever

The most common severe abnormal lab test results with ELREXFIO include decreased white blood cells, red blood cells, and platelets.

These are not all of the possible side effects of ELREXFIO.

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

General information about the safe and effective use of ELREXFIO.

Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide. You can ask your pharmacist or healthcare provider for more information about ELREXFIO that is written for health professionals.

What are the ingredients in ELREXFIO?

Active ingredient: elranatamab-bcmm

Inactive ingredients: edetate disodium, histidine, L-histidine hydrochloride monohydrate, polysorbate 80, sucrose, and Water for Injection

Manufactured by: Pfizer Inc., NY, NY 10001

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Division of Pfizer Inc.

New York, NY 10001

LAB-1551-0.4

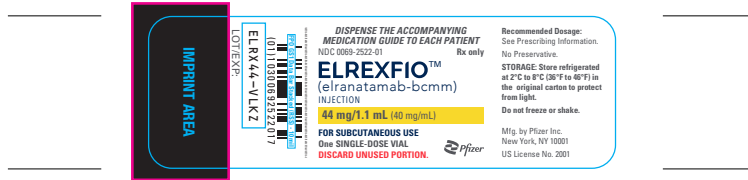
For more information on ELREXFIO, go to www.ELREXFIO.com

For more information on Pfizer, go to www.Pfizer.com or call 1-800-438-1985

This Medication Guide has been approved by the U.S. Food and Drug Administration.

Issued: August 2023

ELREXFIO elranatamab 44mg Vial Label OTD LOGO 18P June 29, 2023 am
Black PMS 032 PMS 604 PMS 655 PMS cool gray 11



NOTE: expiry format:
EXP: MM/YYYY

090177e19de5742b\Approved\Approved On: 29-Jun-2023 20:25 (GMT)

090177e19de57d8b\Approved\Approved On: 29-Jun-2023 20:35 (GMT)

NOTE: expiry format:
EXP: MM/YYYY

ELRX44-FCKZ

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NO VARNISH
NO INK


LOT/EXP/SN: GTIN: 00300692522027

One SINGLE-DOSE VIAL

44 mg/1.1 mL (40 mg/mL)

INJECTION
ELREXFIO™
(elranatamab-bcmm)

ELREXFIO elranatamab 44mg vial carton OTD LOGO 18P June 28, 2023 am
 PMS 032 PMS 604 PMS 655 PMS cool gray 2 PMS cool gray 11




DISPENSE THE ACCOMPANYING
MEDICATION GUIDE TO EACH PATIENT
NDC 0069-2522-02

ELREXFIO™
(elranatamab-bcmm)
INJECTION

44 mg/1.1 mL (40 mg/mL)

For Subcutaneous Injection
by a Healthcare Provider Only
One SINGLE-DOSE VIAL
DISCARD UNUSED PORTION.
Rx only



Recommended Dosage:
See Prescribing Information.
No Preservative.

STORAGE: Store refrigerated at
2°C to 8°C (36°F to 46°F) in the original
carton to protect from light.
Do not freeze or shake.

Manufactured by
Pfizer Inc.
New York, NY 10001
US License No. 2001


Distributed by
Pfizer Labs
Division of Pfizer Inc.
New York, NY 10001

CONTENTS: Each carton contains one
preservative-free 44 mg (40 mg/mL)
single-dose vial of ELREXFIO™.
Each mL of solution contains 40 mg
elranatamab-bcmm, edetate disodium
(0.045 mg), histidine (1.12 mg), L-histidine
hydrochloride monohydrate (2.67 mg),
polysorbate 80 (0.2 mg), sucrose (85 mg)
and Water for Injection.

ELREXFIO™
(elranatamab-bcmm)
INJECTION

44 mg/1.1 mL (40 mg/mL)

For Subcutaneous Injection
by a Healthcare Provider Only
One SINGLE-DOSE VIAL
DISCARD UNUSED PORTION.



300692522027

ELREXFIO elranatamab 44mg Vial Label SAMPLE ELRXSMPL-VL 9P June 28, 2023 am
Black PMS 032 PMS 604 PMS 655 PMS cool gray 11

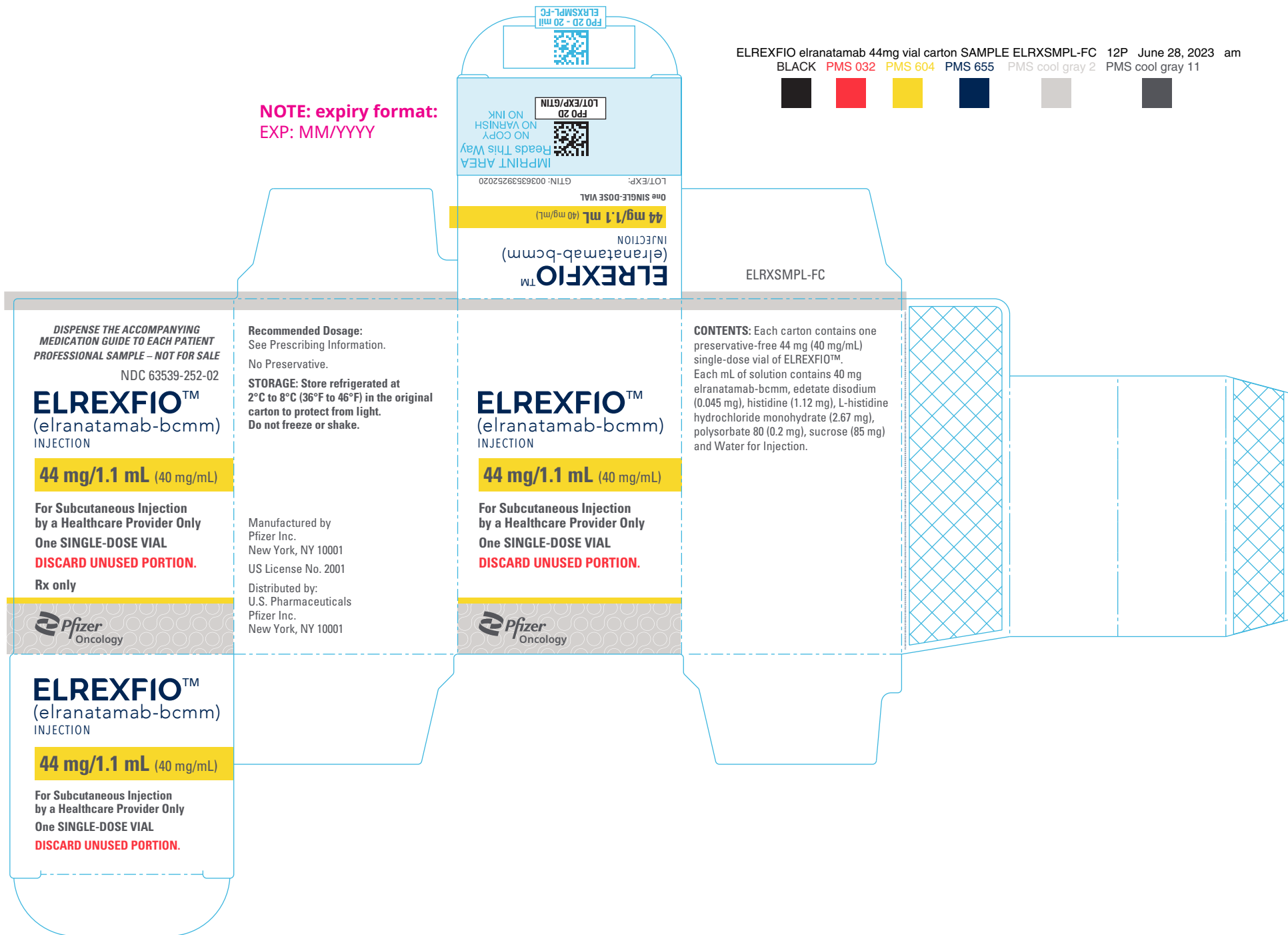


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**NOTE: expiry format:
 EXP: MM/YYYY**

ELREXFIO elranatamab 44mg vial carton SAMPLE ELRXSMPL-FC 12P June 28, 2023 am
 BLACK PMS 032 PMS 604 PMS 655 PMS cool gray 2 PMS cool gray 11



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ELREXFIO elranatamab 76mg Vial Label OTD LOGO 10P June 28, 2023 am

Black PMS 032 PMS 655 PMS cool gray 11



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NOTE: expiry format:
EXP: MM/YYYY

ELRX76-FCKZ


ELREXFIO™
(elranatamab-bcmm)
INJECTION

76 mg/1.9 mL (40 mg/mL)

One SINGLE-DOSE VIAL

For Subcutaneous Injection
by a Healthcare Provider Only

One SINGLE-DOSE VIAL
DISCARD UNUSED PORTION.
Rx only




ELREXFIO™
(elranatamab-bcmm)
INJECTION

76 mg/1.9 mL (40 mg/mL)


For Subcutaneous Injection
by a Healthcare Provider Only

One SINGLE-DOSE VIAL
DISCARD UNUSED PORTION.



ELREXFIO elranatamab 76mg vial carton OTD LOGO 11P June 28, 2023 am

PMS 032 PMS 655 PMS cool gray 2 PMS cool gray 11




ELRX76-FCKZ


ELREXFIO™
(elranatamab-bcmm)
INJECTION

76 mg/1.9 mL (40 mg/mL)

For Subcutaneous Injection
by a Healthcare Provider Only

One SINGLE-DOSE VIAL
DISCARD UNUSED PORTION.




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 300694494025

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LOT/EXP/SN: GTIN: 00300694494025

One SINGLE-DOSE VIAL
76 mg/1.9 mL (40 mg/mL)

ELREXFIO™
(elranatamab-bcmm)
INJECTION

ELRX76-FCKZ


ELRX76-FCKZ

ELREXFIO™
(elranatamab-bcmm)
INJECTION

76 mg/1.9 mL (40 mg/mL)

For Subcutaneous Injection
by a Healthcare Provider Only

One SINGLE-DOSE VIAL
DISCARD UNUSED PORTION.




ELREXFIO™
(elranatamab-bcmm)
INJECTION

76 mg/1.9 mL (40 mg/mL)

For Subcutaneous Injection
by a Healthcare Provider Only

One SINGLE-DOSE VIAL
DISCARD UNUSED PORTION.



DISPENSE THE ACCOMPANYING
MEDICATION GUIDE TO EACH PATIENT


NDC 0069-4494-02

ELREXFIO™
(elranatamab-bcmm)
INJECTION

76 mg/1.9 mL (40 mg/mL)

For Subcutaneous Injection
by a Healthcare Provider Only

One SINGLE-DOSE VIAL
DISCARD UNUSED PORTION.
Rx only



Recommended Dosage:
See Prescribing Information.

No Preservative.

STORAGE: Store refrigerated at
2°C to 8°C (36°F to 46°F) in the original
carton to protect from light.
Do not freeze or shake.

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CONTENTS: Each carton contains one
preservative-free 76 mg (40 mg/mL)
single-dose vial of ELREXFIO™.

Each mL of solution contains 40 mg
elranatamab-bcmm, edetate disodium
(0.045 mg), histidine (1.12 mg), L-histidine
hydrochloride monohydrate (2.67 mg),
polysorbate 80 (0.2 mg), sucrose (85 mg)
and Water for Injection.


ELRX76-FCKZ

ELREXFIO™
(elranatamab-bcmm)
INJECTION

76 mg/1.9 mL (40 mg/mL)

For Subcutaneous Injection
by a Healthcare Provider Only

One SINGLE-DOSE VIAL
DISCARD UNUSED PORTION.



Risk Evaluation and Mitigation Strategy (REMS) Document ELREXFIO (elranatamab-bcmm) REMS

I. Administrative Information

Risk: Cytokine Release Syndrome, neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome

Application Number: BLA 761345

Application Holder: Pfizer Inc.

Initial REMS Approval: 08/2023

II. REMS Goal

The goal of the ELREXFIO REMS is to mitigate the risks of Cytokine Release Syndrome (CRS), and neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) by:

- Ensuring prescribers are aware of the importance of monitoring for signs and symptoms of CRS and neurologic toxicity including ICANS in patients exposed to ELREXFIO

III. REMS Requirements

Pfizer Inc. must ensure that healthcare providers, patients, pharmacies, healthcare settings and wholesalers-distributors comply with the following requirements:

1. Healthcare providers who prescribe ELREXFIO must:

To become certified to prescribe

1. Review the drug's Prescribing Information.
2. Review the [Prescriber Training Program](#) and [Adverse Reaction Management Guide](#).
3. Successfully complete the [Knowledge Assessment](#) and submit it to the REMS.
4. Enroll in the REMS by completing and submitting the [Prescriber Enrollment Form](#) to the REMS.

Before treatment initiation (first step-up dose)

5. Counsel the patient on:
 - a) how to recognize and respond to signs and symptoms of CRS and neurologic toxicity including ICANS,
 - b) the need to report all symptoms suggestive of CRS and neurologic toxicity including ICANS to their healthcare provider or emergency room provider immediately,
 - c) the need to carry the [Patient Wallet Card](#) at all times.
 6. Complete the [Patient Wallet Card](#) and provide the [Patient Wallet Card](#) to the patient.
-

| | |
|---|---|
| At all times | 7. Report serious adverse events of CRS and neurologic toxicity including ICANS to the REMS. |
| 2. Patients who are prescribed ELREXFIO: | |
| Before treatment | 1. Receive counseling from the prescriber using the Patient Wallet Card . |
| At all times | 2. Have the Patient Wallet Card with you and inform other healthcare providers about treatment with ELREXFIO. |
| 3. Pharmacies and healthcare settings that dispense ELREXFIO must: | |
| To become certified to dispense | <ol style="list-style-type: none">1. Designate an Authorized Representative to carry out the certification process and oversee implementation and compliance with the REMS requirements on behalf of the pharmacy and/or healthcare setting.2. Have the Authorized Representative review the Pharmacy and Healthcare Setting Training Program.3. Have the Authorized Representative enroll in the REMS by completing and submitting the Pharmacy and Healthcare Setting Enrollment Form to the REMS.4. Train all relevant staff involved in dispensing ELREXFIO on the REMS requirements using the Pharmacy and Healthcare Setting Training Program. |
| Before dispensing | 5. Obtain authorization to dispense each prescription by contacting the REMS to verify the prescriber is certified. |
| To maintain certification to dispense | 6. Have a new Authorized Representative enroll in the REMS by completing and submitting the Pharmacy and Healthcare Setting Enrollment Form to the REMS if the Authorized Representative changes. |
| At all times | <ol style="list-style-type: none">7. Report serious adverse events of CRS and neurologic toxicity including ICANS to the REMS.8. Not distribute, transfer, loan or sell ELREXFIO except to certified pharmacies and healthcare settings.9. Maintain records of staff training.10. Maintain records that processes and procedures are in place and are being followed.11. Maintain records of all ELREXFIO dispenses and provide data to the REMS and Wholesalers-Distributors, as requested.12. Comply with audits carried out by Pfizer Inc. or a third party acting on behalf of Pfizer Inc. to ensure that all training, processes, and procedures are in place and are being followed. |
| 4. Wholesalers-distributors that distribute ELREXFIO must: | |
| To be able to distribute | <ol style="list-style-type: none">1. Establish processes and procedures to ensure that ELREXFIO is distributed only to certified pharmacies and healthcare settings.2. Train all relevant staff involved in distribution on the REMS requirements. |
| At all times | 3. Distribute only to certified pharmacies and healthcare settings. |

4. Maintain records that all processes and procedures are in place and are being followed.
5. Maintain and submit records of drug distribution to the REMS at least on a monthly basis.
6. Comply with audits carried out by Pfizer Inc. or a third party acting on behalf of Pfizer Inc. to ensure that all processes and procedures are in place and are being followed.

Pfizer Inc. must provide training to healthcare providers who prescribe ELREXFIO. The training includes the following educational materials: [Prescriber Training Program](#), [Adverse Reaction Management Guide](#), and [Knowledge Assessment](#). The training must be available online.

Pfizer Inc. must provide training to pharmacies and healthcare settings that dispense ELREXFIO. The training includes the following educational material: [Pharmacy and Healthcare Setting Training Program](#). The training must be available online.

To inform healthcare providers about the REMS and the risks and safe use of ELREXFIO, Pfizer Inc. must disseminate REMS communication materials according to the table below:

| Target Audience | Communication Materials & Dissemination Plans |
|---|---|
| Healthcare providers including oncologists, oncology physician assistants, oncology nurse practitioners, hematologists, and oncology nurses who are likely to prescribe and care for patients treated with ELREXFIO | <p>REMS Letters: Healthcare Provider REMS Letter, Professional Society REMS Letter with attachment REMS Fact Sheet</p> <ol style="list-style-type: none"> 1. E-mail within 30 calendar days of the date ELREXFIO is first commercially distributed and again 12 months later. <ol style="list-style-type: none"> a. Send by mail within 30 calendar days of the date the first e-mail was sent if a healthcare provider’s e-mail address is not available, or the e-mail is undeliverable. b. Send a second e-mail within 30 calendar days of the date the first e-mail was sent if the first e-mail is marked unopened. c. Send by mail within 30 calendar days of the date the second e-mail was sent if the second e-mail is marked as unopened. 2. Disseminate through field-based sales and medical representatives for 12 months from the date ELREXFIO is first commercially distributed. 3. Disseminate within 30 calendar days of the date ELREXFIO is first commercially distributed and again 12 months later through the following professional societies and request the letter or content be provided to their members. <ol style="list-style-type: none"> a. American Society of Clinical Oncology (ASCO); American Society of Hematology (ASH); Advanced Practitioner Society for Hematology and Oncology (APSHO); Oncology Nursing Society (ONS); National Comprehensive Cancer Network (NCCN); Society of Hematologic Oncology (SOHO); Hematology Oncology Pharmacy Association (HOPA); American Pharmacists Association (APhA); American Society of Health-System Pharmacists (ASHP) 4. Disseminate at Professional Meetings where Pfizer Inc. has a presence for 12 months from the date ELREXFIO is first commercially distributed. <p>REMS Fact Sheet</p> <ol style="list-style-type: none"> 1. Disseminate at Professional Meetings where Pfizer Inc. has a presence for 12 months from the date ELREXFIO is first commercially distributed. 2. Disseminate through field-based sales and medical representatives during the initial and/or follow-up discussion with healthcare providers for 12 months after ELREXFIO is first commercially distributed. Field-based sales and/or medical representatives to orally review the key risk messages contained in the REMS Fact Sheet during the visit with the healthcare provider. |

To support REMS operations, Pfizer Inc. must:

1. Authorize dispensing for each prescription based on verifying the prescriber is certified.
2. Establish and maintain the [REMS Website](http://www.ELREXFIOREMS.com), www.ELREXFIOREMS.com. The [REMS Website](http://www.ELREXFIOREMS.com) must include the capability to enroll prescribers, pharmacies and healthcare settings, complete training online, maintain records of that training, and an option to print the Prescribing Information (PI), Medication Guide, and REMS materials. All product websites for consumers and healthcare providers must include prominent REMS- specific links to the [REMS Website](http://www.ELREXFIOREMS.com).
3. Make the [REMS Website](http://www.ELREXFIOREMS.com) fully operational and all REMS materials available through the website and the REMS Coordinating Center at the time ELREXFIO first becomes commercially available.
4. Establish and maintain a REMS Coordinating Center for REMS participants at 1-844-923-7845.
5. Establish and maintain a validated, secure database of all REMS participants who are enrolled and/or certified in the REMS.
6. Ensure prescribers, pharmacies and healthcare settings can enroll in the REMS online, by fax and e-mail.
7. Ensure pharmacies and healthcare settings are able to obtain authorization to dispense by phone and online.
8. Notify prescribers, pharmacies and healthcare settings within 1 calendar day after they become certified in the REMS.
9. Provide certified prescribers access to the database of certified pharmacies and healthcare settings.
10. Provide certified pharmacies and healthcare settings access to the database of certified prescribers.
11. Provide authorized wholesalers-distributors access to the list of certified pharmacies and healthcare settings.
12. Report serious CRS and neurologic toxicity including ICANS as soon as possible to the FDA but no later than 15 calendar days from the initial receipt of the information by the applicant. This requirement does not affect the applicant's other reporting and follow-up requirements under applicable FDA regulations.

To ensure REMS participants' compliance with the REMS, Pfizer Inc. must:

13. Verify annually that the designated authorized representative for certified pharmacies and healthcare settings remains the same. If different, the pharmacy/healthcare setting must re-certify with a new authorized representative.
14. Maintain adequate records to demonstrate that REMS requirements have been met, including, but not limited to records of: ELREXFIO distribution and dispensing; certification of pharmacies and healthcare settings; authorized wholesalers-distributors; and audits of REMS participants. These records must be readily available for FDA inspections.
15. Monitor pharmacies, healthcare settings, and wholesalers-distributors on an ongoing basis to ensure the requirements of the REMS are being met. Take corrective action if noncompliance is identified, including decertification or deauthorization.
16. Establish and maintain a plan for addressing noncompliance with REMS requirements.
17. Audit all certified pharmacies and healthcare settings within 180 calendar days after the pharmacy or healthcare setting receives their first shipment of ELREXFIO and annually thereafter to ensure that all REMS processes and procedures are in place, functioning, and support the REMS requirements.
18. Audit wholesalers-distributors that have distributed ELREXFIO within 180 calendar days of being authorized to distribute ELREXFIO and annually thereafter to ensure that all REMS processes and procedures are in place, functioning, and support the REMS requirements. Take corrective action if noncompliance is identified.
19. Take reasonable steps to improve operations of and compliance with the requirements in the ELREXFIO REMS based on monitoring and evaluation of the ELREXFIO REMS.

IV. REMS Assessment Timetable

Pfizer Inc. must submit REMS Assessments to the FDA annually from the date of the initial approval of the REMS (08/14/2023). To facilitate inclusion of as much information as possible while allowing reasonable time to prepare the submission, the reporting interval covered by each assessment should conclude no earlier than 60 calendar days before the submission date for that assessment. Pfizer Inc. must submit each assessment so that it will be received by the FDA on or before the due date.

V. REMS Materials

The following materials are part of the ELREXFIO REMS:

Enrollment Forms

Prescriber:

1. [Prescriber Enrollment Form](#)

Pharmacy and Healthcare Setting:

2. [Pharmacy and Healthcare Setting Enrollment Form](#)

Training and Educational Materials

Prescriber:

3. [Prescriber Training Program](#)
4. [Adverse Reaction Management Guide](#)
5. [Knowledge Assessment](#)

Patient:

6. [Patient Wallet Card](#)

Pharmacy and Healthcare Setting:

7. [Pharmacy and Healthcare Setting Training Program](#)

Communication Materials

8. [Healthcare Provider REMS Letter](#)
9. [Professional Society REMS Letter](#)
10. [REMS Fact Sheet](#)

Other Materials

11. [REMS Website](#)

VI. Statutory Elements

This REMS is required under section 505-1 of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 355-1) and consists of the following elements:

1. Communication Plan
2. Elements to Assure Safe Use
 - Health care providers who prescribe ELREXFIO are specially certified under 505-1(f)(3)(A)
 - Pharmacies and health care settings that dispense ELREXFIO are specially certified under 505-1(f)(3)(B)
3. Implementation System
4. Timetable for Submission of Assessments

ELREXFIO™ REMS Prescriber Enrollment Form



Instructions

To become a certified prescriber in the ELREXFIO REMS and prescribe ELREXFIO:

- Review the Prescribing Information, the **Prescriber Training Program** and **Adverse Reaction Management Guide**.
- Successfully complete and submit the **Knowledge Assessment** and this form online at www.ELREXFIOREMS.com
- If not enrolling online,
 - Complete all required fields below and submit this form to the REMS Coordinating Center by e-mail at ELREXFIOREMS@ubc.com, or via fax to 1-800-349-5131.
 - Complete the **Knowledge Assessment** and submit it to the REMS Coordinating Center by e-mail at ELREXFIOREMS@ubc.com, or via fax to 1-800-349-5131.
 - The ELREXFIO REMS will verify both the **Knowledge Assessment** and **Prescriber Enrollment Form** are complete and provide confirmation of certification via e-mail after processing.

If ELREXFIO will be dispensed and administered in the same location, an Authorized Representative must complete the Pharmacy and Healthcare Setting certification.

Certified ELREXFIO Prescribers cannot be designated as the Authorized Representative for a Pharmacy or Healthcare Setting.

PRESCRIBER INFORMATION (Fields marked with an * are REQUIRED)

| | | | |
|---|---------|---------------------|---------------------|
| First Name*: | | Last Name*: | |
| Credentials*: <input type="checkbox"/> MD <input type="checkbox"/> DO <input type="checkbox"/> NP <input type="checkbox"/> PA <input type="checkbox"/> Other (please specify) _____ | | | |
| Specialty* (Select one): <input type="checkbox"/> Oncology <input type="checkbox"/> Hematology <input type="checkbox"/> Internal Medicine/Family Medicine <input type="checkbox"/> Other (please specify) _____ | | | |
| National Provider Identifier (NPI)#*: | | State License #*: | State of Practice*: |
| Practice/Facility Name*: | | | |
| Address Line 1*: | | | |
| Address Line 2: | | | |
| City*: | State*: | Zip Code*: | |
| E-mail*: | Phone*: | Fax: | |
| I am a prescriber becoming certified in the ELREXFIO REMS. By signing this form, I agree to comply with all REMS requirements. | | | |
| To become certified to prescribe, I must: | | | |
| <ul style="list-style-type: none">• Review the ELREXFIO Prescribing Information.• Review the Prescriber Training Program and the Adverse Reaction Management Guide.• Successfully complete the Knowledge Assessment and submit it to the REMS.• Enroll in the REMS by completing and submitting the Prescriber Enrollment Form to the REMS. | | | |
| Before treatment initiation (first step-up dose) of ELREXFIO, I must: | | | |
| <ul style="list-style-type: none">• Counsel the patient on:<ul style="list-style-type: none">• how to recognize and respond to symptoms of Cytokine Release Syndrome (CRS) and neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS)• the need to report all symptoms suggestive of CRS and neurologic toxicity including ICANS to their healthcare provider or emergency room healthcare provider immediately,• the need to carry the Patient Wallet Card at all times.• Complete the Patient Wallet Card and provide the Patient Wallet Card to the patient. | | | |
| At all times, I must: | | | |
| <ul style="list-style-type: none">• Report serious adverse events of CRS and neurologic toxicity including ICANS to the REMS. | | | |
| Signature*: | | Date (MM/DD/YYYY)*: | |



ELREXFIO™ REMS Pharmacy and Healthcare Setting Enrollment Form



Instructions

To become certified in the ELREXFIO REMS and dispense ELREXFIO, a pharmacy/healthcare setting must designate an ***Authorized Representative to:**

- Review the **Pharmacy and Healthcare Setting Training Program**
- Complete this **Pharmacy and Healthcare Setting Enrollment Form:**
 - online at www.ELREXFIOREMS.com for immediate enrollment
 - by e-mail at ELREXFIOREMS@ubc.com or
 - via fax to 1-800-349-5131.
- The ELREXFIO REMS will verify that the **Pharmacy and Healthcare Setting Enrollment Form** is complete and provide confirmation of certification via e-mail after processing.
- ELREXFIO cannot be dispensed until the Pharmacy and Healthcare Setting certification is complete.

***Certified ELREXFIO Prescribers cannot be designated as the Authorized Representatives for a Pharmacy or Healthcare Setting.**

PHARMACY AND HEALTHCARE SETTING INFORMATION (Fields marked with an * are REQUIRED)

| | | |
|--|---------|--|
| Enrollment Type (Please Check One*) <input type="checkbox"/> New Certification <input type="checkbox"/> Change in Authorized Representative | | |
| Pharmacy/Healthcare Setting Name*: | | |
| Pharmacy/Healthcare Setting National Provider Identifier (NPI)#*: (If you do not have an NPI, please reach out to the ELREXFIO REMS) | | DEA# (On file with distributor account): |
| Pharmacy/Healthcare Setting Type (select one)*: <input type="checkbox"/> Inpatient Hospital Pharmacy <input type="checkbox"/> Outpatient Hospital Pharmacy <input type="checkbox"/> Oncology Infusion Center <input type="checkbox"/> Community Oncology Physician Office <input type="checkbox"/> Specialty Pharmacy <input type="checkbox"/> Other (please specify) _____ | | |
| Address Line 1*: | | Address Line 2: |
| City*: | State*: | Zip Code*: |
| Phone*: | | Fax*: |

SHIP TO INFORMATION (Fields marked with an * are REQUIRED)

| | | |
|--|---------|-----------------|
| Ship To Address <input type="checkbox"/> Same as Above Ship To Contact Name*: | | |
| Address Line 1*: | | Address Line 2: |
| City*: | State*: | Zip Code*: |
| Phone*: | | Fax*: |

ELREXFIO™ REMS Pharmacy and Healthcare Setting Enrollment Form



AUTHORIZED REPRESENTATIVE INFORMATION (Fields marked with an * are REQUIRED)

| | |
|---|-------------|
| E-mail*: | |
| First Name*: | Last Name*: |
| Credentials* (select one): <input type="checkbox"/> PharmD <input type="checkbox"/> R.Ph <input type="checkbox"/> RN <input type="checkbox"/> Other (please specify): (e.g., MD, DO, Office Administrator, Practice Manager): | |
| Phone*: | Fax*: |

Your pharmacy or healthcare setting's information will be shared with Pfizer Inc.'s wholesaler-distributor partners, to allow your pharmacy or healthcare setting to purchase product.

AUTHORIZED REPRESENTATIVE RESPONSIBILITIES

As the Authorized Representative, I must:

- Review the **Pharmacy and Healthcare Setting Training Program**.
- Enroll in the REMS by completing and submitting this **Pharmacy and Healthcare Setting Enrollment Form** to the REMS.
- Train all relevant staff involved in dispensing ELREXFIO on the REMS requirements using the **Pharmacy and Healthcare Setting Training Program**.
- Oversee implementation and compliance with the REMS requirements on behalf of the pharmacy and/or healthcare setting.

Before dispensing, staff must:

- Obtain authorization to dispense each prescription by contacting the REMS to verify the prescriber is certified.

At all times, staff must:

- Report serious adverse events of Cytokine Release Syndrome (CRS) and neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) to the REMS.
- Not distribute, transfer, loan or sell ELREXFIO except to certified pharmacies and healthcare settings.
- Maintain records of staff training.
- Maintain records that processes and procedures are in place and are being followed.
- Maintain records of all ELREXFIO dispenses and provide data to the REMS and Wholesalers-Distributors, as requested.
- Comply with audits carried out by Pfizer Inc. or a third party acting on behalf of Pfizer Inc. to ensure that all training, processes, and procedures are in place and are being followed.
- Have a new Authorized Representative enroll in the REMS by completing and submitting the **Pharmacy and Healthcare Setting Enrollment Form** to the REMS if the Authorized Representative changes.

| | |
|---------------------------------------|---------------------|
| Authorized Representative Signature*: | Date (MM/DD/YYYY)*: |
|---------------------------------------|---------------------|



PRESCRIBER TRAINING PROGRAM

Risk Evaluation and Mitigation Strategy (REMS)



ELREXFIO Prescriber Training Module

- This educational module contains information on adverse reactions associated with ELREXFIO including Cytokine Release Syndrome (CRS) and neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS). The adverse reactions listed within this module are not all inclusive of adverse reactions associated with ELREXFIO.
- Please refer to the Prescribing Information for additional information.

This training module provides details related to prescriber requirements of the ELREXFIO REMS.

1 ELREXFIO Prescriber Training Module



Training Outline

- **Prescriber REMS Requirements** (Slides 6-7)
- **Risk of Cytokine Release Syndrome (CRS)** (Slides 9-14)
- **Risk of Neurologic Toxicity including Effector Cell-Associated Neurotoxicity Syndrome (ICANS)** (Slides 16-18)

2 ELREXFIO Prescriber Training Module

 **ELREXFIO™**
(elranatamab-bcmm)
INJECTION FOR SUBCUTANEOUS USE | 44 mg/1 mL | 76 mg/1.9 mL

What is a REMS?

- A Risk Evaluation and Mitigation Strategy (REMS) is a program required by the FDA to manage known or potential serious risks associated with a drug product. The FDA has determined that a REMS is necessary to ensure that the benefits of ELXREXFIO outweigh its risks.
- The goal of the ELXREXFIO REMS is to mitigate the risks of Cytokine Release Syndrome (CRS) and neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) by ensuring prescribers are aware of the importance of monitoring for signs and symptoms of CRS and neurologic toxicity including ICANS in patients exposed to ELXREXFIO.

3 ELXREXFIO Prescriber Training Module



ELREXFIO Indication

- ELREXFIO is a B-cell maturation antigen (BCMA)-directed and CD3-directed bispecific antibody indicated for the treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy, including a proteasome inhibitor, an immunomodulatory agent, and an anti-CD38 monoclonal antibody.

4 ELREXFIO Prescriber Training Module



/01

Prescriber REMS Requirements

 **ELREXFIO**[™]
(elranatamab-bcmm)
INJECTION FOR SUBCUTANEOUS USE | 44 mg/11 mL
75 mg/19 mL

Prescriber REMS Certification Requirements

Requirements to become certified to prescribe ELREXFIO are as follows:

- Review the Prescribing Information, this **Prescriber Training Program**, and the **Adverse Reaction Management Guide**
- Successfully complete the **Knowledge Assessment** and submit it to the REMS
- Enroll in the REMS by completing and submitting the **Prescriber Enrollment Form** to the REMS
- Before treatment initiation (first step-up dose), complete the **Patient Wallet Card** and provide the **Patient Wallet Card** to the patient

At all times

- Report serious adverse events of CRS and neurologic toxicity including ICANS to the REMS

ELREXFIO Prescribers cannot be designated as the Authorized Representative for a certified Pharmacy or Healthcare Setting*

*If ELREXFIO will be dispensed and administered in the same location, an Authorized Representative must complete the Pharmacy and Healthcare Setting certification.

Prescriber Counseling

- Counsel the patient or caregiver on how to recognize and respond to signs and symptoms of CRS and neurologic toxicity including ICANS using the **Patient Wallet Card**, and instruct the patient to carry it with them at all times.
- Instruct the patient that they should be hospitalized for 48 hours after administration of the first step-up dose, and for 24 hours after administration of the second step-up dose.
- Instruct the patient of the need to report all symptoms suggestive of CRS and/or neurologic toxicity including ICANS to their healthcare provider or emergency room provider immediately.
- Advise patients not to drive or operate heavy or potentially dangerous machinery for 48 hours after completing each of the 2 step-up doses and the first treatment dose within the ELREXFIO step-up dosing schedule and in the event of new onset of any neurological toxicity symptoms until symptoms resolve.

/02

Cytokine Release Syndrome (CRS)

 **ELREXIO™**
(elranatamab-bcmm)
INJECTION FOR SUBCUTANEOUS USE | 44 mg/11 mL
76 mg/19 mL

ELREXFIO Boxed Warning

Cytokine Release Syndrome (CRS), including life-threatening or fatal reactions, can occur in patients receiving ELREXFIO. Initiate treatment with ELREXFIO step-up dosing schedule to reduce risk of CRS. Withhold ELREXFIO until CRS resolves or permanently discontinue based on severity.

Neurologic toxicity, including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS), and serious and life-threatening reactions, can occur in patients receiving ELREXFIO. Monitor patients for signs and symptoms of neurologic toxicity, including ICANS, during treatment. Withhold ELREXFIO until the neurologic toxicity resolves or permanently discontinue based on severity.

ELREXFIO is available only through a restricted program called the ELREXFIO Risk Evaluation and Mitigation Strategy (REMS).



Cytokine Release Syndrome

- CRS, including life-threatening or fatal reactions, can occur following treatment with ELREXFIO
- CRS was reported in 58% of patients who received the recommended dosage in the clinical trial (n=183)
- Most patients experienced CRS after the first step-up dose (43%) or the second step-up dose (19%), with 7% of patients having CRS after the first treatment dose and 1.6% of patients after a later dose
- Recurrent CRS occurred in 13% of patients
- Patients developed mostly Grade 1 (44%) and Grade 2 (14%) CRS.
- 1 (0.5%) patient developed Grade 3 CRS
- The median time to onset of CRS was 2 (range: 1 to 9) days after the most recent dose of ELREXFIO
- The median duration of CRS was 2 (range: 1 to 19) days

Clinical Signs and Symptoms of CRS

- Patients should be closely monitored for signs or symptoms of CRS
- Potentially life-threatening complications of CRS may include:
 - Cardiac dysfunction
 - Adult respiratory distress syndrome
 - Neurologic toxicity
 - Renal and/or hepatic failure
 - Disseminated intravascular coagulation (DIC)

Signs and Symptoms

- Fever
- Chills
- Hypoxia
- Hypotension
- Sinus Tachycardia
- Headache
- Elevated Liver Enzymes

Lee DW, Gardner R, Porter DL, et al. Current concepts in the diagnosis and management of cytokine release syndrome. Blood 2014; 124(2): 188-95. Errata in Blood: 2015;126(8):1048. And 2016;128(11):1533

ELREXFIO Dosing Schedule

| Dosing Schedule | Day | ELREXFIO Dose | |
|--|--|----------------------------|-------|
| | | | |
| Step-up Dosing Schedule | Day 1 ^a , | Step-up dose 1 | 12 mg |
| | Day 4 ^{a,b} | Step-up dose 2 | 32 mg |
| | Day 8 ^{a,c} | First treatment dose | 76 mg |
| Weekly Dosing Schedule | One week after first treatment dose and weekly thereafter ^d through week 24 | Subsequent treatment doses | 76 mg |
| Biweekly (Every 2 Weeks) Dosing Schedule *Responders only week 25 onward | Week 25 and every 2 weeks thereafter ^d | Subsequent treatment doses | 76 mg |

^a Administer pre-treatment medications prior to each dose in the ELREXFIO step-up dosing schedule, which includes step-up dose 1, step-up dose 2, and the first treatment dose

^b A minimum of 2 days should be maintained between step-up dose 1 (12 mg) and step-up dose 2 (32 mg).

^c A minimum of 3 days should be maintained between step-up dose 2 (32 mg) and the first treatment (76 mg) dose.

^d A minimum of 6 days should be maintained between treatment doses.

Note: See Table 2 in Prescribing Information for recommendations on restarting ELREXFIO after dose delays.

Important Dosing Information

- Administer the following pretreatment medications approximately 1 hour before the first three doses of ELREXFIO in the step-up dosing schedule, which includes step-up dose 1, step-up dose 2, and the first treatment dose to reduce the risk of CRS:
 - Acetaminophen (or equivalent) 650 mg orally
 - Dexamethasone (or equivalent) 20 mg orally or intravenously
 - Diphenhydramine (or equivalent) 25 mg orally
- Due to the risk of CRS, patients should be hospitalized for 48 hours after administration of the first step-up dose, and for 24 hours after administration of the second step-up dose.

CRS Management

- Counsel patients to seek medical attention should signs or symptoms of CRS occur.
- At the first sign of CRS, immediately evaluate patients for hospitalization.
- Withhold or permanently discontinue ELREXFIO based on severity and manage CRS according to the recommendations in the Prescribing Information and in the **Adverse Reaction Management Guide**. Consider further management per current practice guidelines.
- Administer supportive therapy for CRS based on severity, which may include intensive care for severe or life-threatening CRS.
- Consider laboratory testing to monitor for disseminated intravascular coagulation (DIC), hematology parameters, as well as pulmonary, cardiac, renal, and hepatic function.

/03

Neurologic Toxicity Including ICANS

 **ELREXFIO**[™]
(elranatamab-bcmm)
INJECTION FOR SUBCUTANEOUS USE | 44 mg/11 mL
76 mg/19 mL

Neurologic Toxicity Including ICANS

- Neurologic toxicity, including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS), and serious and life-threatening reactions, can occur in patients receiving ELREXFIO.
- In the clinical trial, neurologic toxicity was reported in 59% of patients receiving ELREXFIO.
- Grade 3 or 4 neurologic toxicity occurred in 7% of patients. Neurologic toxicities included headache (18%), encephalopathy (14%), motor dysfunction (14%), sensory neuropathy (13%), and Guillain-Barre Syndrome (0.5%)
- In the clinical trial, ICANS was reported in 3.3% of patients receiving ELREXFIO.
- The most frequent clinical manifestations of ICANS included a depressed level of consciousness and Grade 1 or Grade 2 Immune Effector Cell-Associated Encephalopathy (ICE) scores.
- The onset of ICANS can be concurrent with CRS, following resolution of CRS, or in the absence of CRS.
- ICANS occurred in 2.7% of patients after the first step-up dose (2.7%),
- One (1; 0.5%) patient had ICANS after the second step-up dose, and
- One (1; 0.5%) patient had ICANS after subsequent dose(s).
- Recurrent ICANS occurred in 1.1% of patients.
- The median time to onset was 3 (range: 1 to 4) days after the most recent dose, with a median duration of 2 (range: 1 to 18) days.

Neurologic Toxicity Management

- Counsel patients to seek medical attention should signs or symptoms of neurologic toxicity occur.
- Monitor patients for signs or symptoms of neurologic toxicity during treatment.
- At the first sign of neurologic toxicity including ICANS, evaluate and treat patients immediately based on severity.
- Withhold or permanently discontinue ELREXFIO and manage neurologic toxicity according to the recommendations in the Prescribing Information and in the table below. Consider further management per current practice guidelines.

| Recommendations for Management of Neurologic Toxicity, excluding ICANS | | |
|--|---------------------------------------|--|
| Adverse Reaction | Severity | Actions |
| Neurologic Toxicity (excluding ICANS) | Grade 1 | <ul style="list-style-type: none">● Withhold ELREXFIO until neurologic toxicity symptoms resolve or stabilize. |
| | Grade 2 Grade 3 (First occurrence) | <ul style="list-style-type: none">● Withhold ELREXFIO until neurologic toxicity symptoms improve to Grade 1 or less.● Provide supportive therapy. |
| | Grade 3 (Recurrent) Grade 4 | <ul style="list-style-type: none">● Permanently discontinue ELREXFIO.● Provide supportive therapy, which may include intensive care. |

ICANS Management

- Counsel patients to seek medical attention should signs or symptoms of ICANS occur.
- Monitor patients for signs or symptoms of ICANS during treatment.
- At the first sign of ICANS, evaluate and treat patients immediately based on severity.
- Withhold or permanently discontinue ELREXFIO and manage ICANS according to the recommendations in the Prescribing Information and **Adverse Reaction Management Guide**. Consider further management per current practice guidelines.

Adverse Event Reporting

- Reporting of suspected adverse events following administration of therapy is vital for the continued monitoring of the risk/benefit balance of therapy.
- Healthcare providers must report serious adverse events of CRS and neurologic toxicity including ICANS to the REMS Coordinating Center at 1-844-923-7845. Report all other suspected adverse events associated with ELREXFIO to Pfizer Inc. at 1-800-438-1985 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

This label may not be the latest approved by FDA.
For current labeling information, please visit <https://www.fda.gov/drugsatfda>

 **ELREXFIO**[™]
(elranatamab-bcmm)
INJECTION FOR SUBCUTANEOUS USE | 44 mg/1.1 mL
76 mg/1.9 mL

For further information, please visit
www.ELREXFIOREMS.com or
call 1-844-923-7845



This label may not be the latest approved by FDA.
For current labeling information, please visit <https://www.fda.gov/drugsatfda>

 **ELREXFIO**TM
(elranatamab-bcmm)

INJECTION FOR | 44 mg/1.1 mL
SUBCUTANEOUS USE | 76 mg/1.9 mL

ELREXFIO

Adverse Reaction Management Guide

This is supplemental to the ELREXFIO US Prescribing Information (USPI).



Management of Cytokine Release Syndrome (CRS)

Identify CRS based on clinical presentation. Evaluate and treat other causes of fever, hypoxia, and hypotension.

If CRS is suspected, withhold ELREXFIO until CRS resolves. Clinical signs and symptoms of CRS may include, but are not limited to, fever, hypoxia, chills, hypotension, tachycardia, headache, and elevated liver enzymes. At first sign of CRS, evaluate patients immediately for hospitalization. Manage according to the recommendations in Table 1 below and consider further management per current practice guidelines. Withhold or permanently discontinue ELREXFIO based on severity. Administer supportive therapy for CRS, which may include intensive care for severe or life-threatening CRS. Consider laboratory testing to monitor for disseminated intravascular coagulation (DIC), hematology parameters, as well as pulmonary, cardiac, renal, and hepatic function.

Table 1: Recommendations for Management of CRS

| Grade ^a | Presenting Symptoms | Actions |
|-------------------------------|---|--|
| Grade 1 | Temperature $\geq 100.4^{\circ}\text{F}$ (38°C) ^b | <ul style="list-style-type: none"> Withhold ELREXFIO until CRS resolves.^c Administer pretreatment medications prior to next dose of ELREXFIO. |
| Grade 2 | <ul style="list-style-type: none"> Temperature $\geq 100.4^{\circ}\text{F}$ (38°C) with either: Hypotension responsive to fluid and not requiring vasopressors, and/or Oxygen requirement of low-flow nasal cannula^d or blow-by | <ul style="list-style-type: none"> Withhold ELREXFIO until CRS resolves.^c Monitor patients daily for 48 hours following the next dose of ELREXFIO. Instruct patients to remain within proximity of a healthcare facility and consider hospitalization. Administer pretreatment medications prior to next dose of ELREXFIO. |
| Grade 3 (First occurrence) | <ul style="list-style-type: none"> Temperature $\geq 100.4^{\circ}\text{F}$ (38°C) with either: Hypotension requiring one vasopressor with or without vasopressin, and/or Oxygen requirement of high-flow nasal cannula^d, facemask, non-rebreather mask, or Venturi mask | <ul style="list-style-type: none"> Withhold ELREXFIO until CRS resolves.^c Provide supportive therapy, which may include intensive care. Patients should be hospitalized for 48 hours following the next dose of ELREXFIO. Administer pretreatment medications prior to next dose of ELREXFIO. |
| Grade 3 (Recurrent) | <ul style="list-style-type: none"> Temperature $\geq 100.4^{\circ}\text{F}$ (38°C) with either: Hypotension requiring one vasopressor with or without vasopressin, and/or Oxygen requirement of high-flow nasal cannula^d, facemask, non-rebreather mask, or Venturi mask. | <ul style="list-style-type: none"> Permanently discontinue therapy with ELREXFIO. Provide supportive therapy, which may include intensive care. |
| Grade 4 | <ul style="list-style-type: none"> Temperature $\geq 100.4^{\circ}\text{F}$ (38°C) with either: Hypotension requiring multiple vasopressors (excluding vasopressin), and/or Oxygen requirement of positive pressure (e.g., continuous positive airway pressure [CPAP], bilevel positive airway pressure [BiPAP], intubation, and mechanical ventilation) | <ul style="list-style-type: none"> Permanently discontinue therapy with ELREXFIO. Provide supportive therapy, which may include intensive care. |

^a. Based on American Society for Transplantation and Cellular Therapy (ASTCT) 2019 grading for CRS.

^b. Attributed to CRS. Fever may not always be present concurrently with hypotension or hypoxia as it may be masked by interventions such as antipyretics or anti-cytokine therapy.

^c. See Table 2 in USPI for recommendations on restarting ELREXFIO after dose delays.

^d. Low-flow nasal cannula is ≤ 6 L/min, and high-flow nasal cannula is >6 L/min.

Management of Neurologic Toxicity and Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS)

Manage neurologic toxicity and ICANS according to the recommendations in Tables 2 and 3 below and consider further management per current practice guidelines.

At the first sign of neurologic toxicity, including ICANS, withhold ELREXFIO and consider neurology evaluation. Rule out other causes of neurologic symptoms. Provide supportive therapy, which may include intensive care, for severe or life-threatening neurologic toxicities including ICANS.

Table 2: Recommendations for Management of Neurologic Toxicity, excluding ICANS

| Adverse Reaction | Severity | Actions |
|---------------------------------------|---------------------------------------|---|
| Neurologic Toxicity (excluding ICANS) | Grade 1 | <ul style="list-style-type: none"> Withhold ELREXFIO until neurologic toxicity symptoms resolve or stabilize. |
| | Grade 2 Grade 3 (First occurrence) | <ul style="list-style-type: none"> Withhold ELREXFIO until neurologic toxicity symptoms improve to Grade 1 or less. Provide supportive therapy. |
| | Grade 3 (Recurrent) Grade 4 | <ul style="list-style-type: none"> Permanently discontinue ELREXFIO. Provide supportive therapy, which may include intensive care. |

Table 3: Recommendations for Management of ICANS

| Grade ^a | Presenting Symptoms ^b | Actions |
|--------------------|---|--|
| Grade 1 | ICE score 7-9 ^c Or depressed level of consciousness ^d : awakens spontaneously. | <ul style="list-style-type: none"> Withhold ELREXFIO until ICANS resolves.^e Monitor neurologic symptoms and consider consultation with a neurologist and other specialists for further evaluation and management. Consider non-sedating, anti-seizure medications (e.g., levetiracetam) for seizure prophylaxis. |
| Grade 2 | ICE score 3-6 ^c or depressed level of consciousness ^d : awakens to voice. | <ul style="list-style-type: none"> Withhold ELREXFIO until ICANS resolves. Administer dexamethasone^f 10 mg intravenously every 6 hours. Continue dexamethasone use until resolution to Grade 1 or less, then taper. Monitor neurologic symptoms and consider consultation with a neurologist and other specialists for further evaluation and management. Consider non-sedating, anti-seizure medications (e.g., levetiracetam) for seizure prophylaxis. Monitor patients daily for 48 hours following the next dose of ELREXFIO. Instruct patients to remain within proximity of a healthcare facility, and consider hospitalization. |

| Grade ^a | Presenting Symptoms | Actions |
|-------------------------------|---|---|
| Grade 3 (First occurrence) | <p>ICE score 0-2^c</p> <p>or depressed level of consciousness^d: awakens only to tactile stimulus,</p> <p>or seizures^d, either:</p> <ul style="list-style-type: none"> • any clinical seizure, focal or generalized, that resolves rapidly, or • non-convulsive seizures on electroencephalogram (EEG) that resolve with intervention, <p>or raised intracranial pressure: focal/local edema on neuroimaging^d</p> | <ul style="list-style-type: none"> • Withhold ELREXFIO until ICANS resolves.^e • Administer dexamethasone^f 10 mg intravenously every 6 hours. Continue dexamethasone use until resolution to Grade 1 or less, then taper. • Monitor neurologic symptoms and consider consultation with a neurologist and other specialists for further evaluation and management. • Consider non-sedating, anti-seizure medications (e.g., levetiracetam) for seizure prophylaxis. • Provide supportive therapy, which may include intensive care. • Patients should be hospitalized for 48 hours following the next dose of ELREXFIO. |
| Grade 3 (recurrent) | <p>ICE score 0-2^c</p> <p>or depressed level of consciousness^d: awakens only to tactile stimulus,</p> <p>or seizures^d, either:</p> <ul style="list-style-type: none"> • any clinical seizure, focal or generalized, that resolves rapidly, or • non-convulsive seizures on electroencephalogram (EEG) that resolve with intervention, <p>or raised intracranial pressure: focal/local edema on neuroimaging^d</p> | <ul style="list-style-type: none"> • Permanently discontinue ELREXFIO. • Administer dexamethasone^f 10 mg intravenously every 6 hours. Continue dexamethasone use until resolution to Grade 1 or less, then taper. • Monitor neurologic symptoms and consider neurology consultation and other specialists for further evaluation and management. • Consider non-sedating, anti-seizure medications (e.g., levetiracetam) for seizure prophylaxis. • Provide supportive therapy, which may include intensive care. |

| Grade ^a | Presenting Symptoms | Actions |
|--------------------|--|---|
| Grade 4 | <p>ICE score 0^c</p> <p>or, depressed level of consciousness^d either:</p> <ul style="list-style-type: none"> • patient is unarousable or requires vigorous or repetitive tactile stimuli to arouse, or • stupor or coma, <p>or seizures^d, either:</p> <ul style="list-style-type: none"> • life-threatening prolonged seizure (>5 minutes), or • repetitive clinical or electrical seizures without return to baseline in between, <p>or motor findings^d:</p> <ul style="list-style-type: none"> • deep focal motor weakness such as hemiparesis or paraparesis, <p>or raised intracranial pressure / cerebral edema^d, with signs/symptoms such as:</p> <ul style="list-style-type: none"> • diffuse cerebral edema on neuroimaging, or • decerebrate or decorticate posturing, or • cranial nerve VI palsy, or • papilledema, or • Cushing's triad | <ul style="list-style-type: none"> • Permanently discontinue ELREXFIO. • Administer dexamethasone^f 10 mg intravenously every 6 hours. Continue dexamethasone use until resolution to Grade 1 or less, then taper. • Alternatively, consider administration of methylprednisolone 1,000 mg per day intravenously for 3 days. • Monitor neurologic symptoms and consider consultation with a neurologist and other specialists for further evaluation and management. • Consider non-sedating, anti-seizure medications (e.g., levetiracetam) for seizure prophylaxis. • Provide supportive therapy, which may include intensive care. |

^a Based on American Society for Transplantation and Cellular Therapy (ASTCT) 2019 grading for ICANS.
^b Management is determined by the most severe event, not attributable to any other cause.
^c If patient is arousable and able to perform Immune Effector Cell-Associated Encephalopathy (ICE) Assessment, assess: Orientation (oriented to year, month, city, hospital = 4 points); Naming (name 3 objects, e.g., point to clock, pen, button = 3 points); Following Commands (e.g., “show me 2 fingers” or “close your eyes and stick out your tongue” = 1 point); Writing (ability to write a standard sentence = 1 point; and Attention (count backwards from 100 by ten = 1 point). If patient is unarousable and unable to perform ICE Assessment (Grade 4 ICANS) = 0 points.
^d Not attributable to any other cause.
^e See Table 2 in USPI for recommendations on restarting ELREXFIO after dose delays.
^f All references to dexamethasone administration are dexamethasone or equivalent medications.

Please visit www.ELXREXFIOREMS.com for further information and resources.



ELREXFIO™ REMS Knowledge Assessment



PRESCRIBER INFORMATION (All Fields Required)

| | | | |
|--|--|-----------------------|--------------------|
| Prescriber First Name: | | Prescriber Last Name: | |
| NPI #: | | State of Practice: | |
| | | State License #: | |
| Address: | | | |
| City: | | State: | Zip Code: |
| Credentials: <input type="checkbox"/> MD <input type="checkbox"/> DO <input type="checkbox"/> NP <input type="checkbox"/> PA <input type="checkbox"/> Other (please specify) _____ | | | |
| Phone: | | Fax: | E-mail: |
| Signature: | | | Date (MM/DD/YYYY): |

As a condition of certification, the prescriber must complete the **Knowledge Assessment**. All 10 questions must be answered correctly.

The **Knowledge Assessment** can be completed online at www.ELREXFIOREMS.com or a completed hard copy can be submitted via fax to 1-800-349-5131 or e-mail at ELREXFIOREMS@ubc.com. You must also complete the **Prescriber Enrollment Form** at www.ELREXFIOREMS.com for your certification to be complete.

The ELREXFIO REMS will verify that both the **Knowledge Assessment** and **Prescriber Enrollment Form** are complete, and will provide confirmation of certification via e-mail after processing.



ELREXFIO™ REMS Knowledge Assessment



- 1- **ELREXFIO is a B-cell maturation antigen (BCMA)-directed and CD3-directed bispecific antibody indicated for the treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy, including a proteasome inhibitor, an immunomodulatory agent, and an anti-CD38 monoclonal antibody:**
 - A- True
 - B- False

- 2- **What is not typically a symptom of CRS:**
 - A- Fever
 - B- Hypotension
 - C- Tinnitus
 - D- Hypoxia
 - E- Tachycardia

- 3- **Which one of the following is true regarding the median time to onset of CRS for ELREXFIO? It typically occurs:**
 - A- 2 (range: 1 to 9) days after the most recent dose
 - B- 6 (range: 3-21) days after the most recent dose
 - C- 10 (range 6-15) days after the most recent dose
 - D- Rarely starts during the first week following ELREXFIO dosing

- 4- **Which of the following regarding Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) are correct:**
 - A- The onset of ICANS can be concurrent with CRS, following resolution of CRS, or in the absence of CRS.
 - B- At the first sign of neurologic toxicity including ICANS immediately evaluate patient and institute treatment based on severity
 - C- ELREXFIO should be withheld from patients experiencing Grade 1 or Grade 2 and Grade 3 (first occurrence) ICANS until resolution of the event
 - D- The most common signs or symptoms of ICANS included depressed level of consciousness and Grade 1 or 2 changes in the Immune Effector Cell-Associated Encephalopathy (ICE) score
 - E- All of the above

- 5- **As a part of patient counseling, which of the following are correct regarding ELREXFIO:**
 - A- Before treatment initiation (first step-up dose), the **Patient Wallet Card** must be provided through the processes and procedures established as a requirement of the REMS Program
 - B- Patients should be hospitalized for 48 hours after administration of the first step-up dose, and for 24 hours after administration of the second step-up dose
 - C- Patients should seek immediate medical attention if they experience symptoms of CRS and/or neurologic toxicity including ICANS
 - D- All of the above

- 6- **Which one of the following is true regarding the median time to onset of Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) for ELREXFIO? It typically occurs:**
 - A- 3 (range: 1 to 4) days after the most recent dose
 - B- 5 (range: 3-15) days after the most recent dose
 - C- 8 (range 6-12) days after the most recent dose
 - D- Rarely starts during the first week following ELREXFIO dosing

ELREXFIO™ REMS Knowledge Assessment



- 7- Administer acetaminophen, dexamethasone, diphenhydramine (or their equivalent) approximately 1 hour before the first 3 doses of the ELREXFIO in the dosing schedule to reduce the risk of CRS:
- A- True
 - B- False
- 8- Potentially life-threatening complications of CRS may include:
- A- Cardiac dysfunction
 - B- Adult respiratory distress
 - C- Neurologic toxicity including ICANS
 - D- Renal and/or hepatic failure
 - E- Disseminated intravascular coagulation (DIC)
 - F- All of the above
- 9- Which neurologic toxicity has been reported at any time during treatment with ELREXFIO:
- A- Headache
 - B- Confusion
 - C- Muscular weakness
 - D- Sensory neuropathy
 - E- Guillain-Barre Syndrome
 - F- All of the above
- 10- If CRS is suspected during treatment with ELREXFIO, which of the following supportive measures should be considered:
- A- Withhold ELREXFIO until Grade 1 or Grade 2 and Grade 3 (first occurrence) CRS resolves
 - B- Administer supportive therapy for CRS, which may include intensive care for severe or life-threatening CRS
 - C- Laboratory testing to monitor for disseminated intravascular coagulation, hematology parameters, as well as pulmonary, cardiac, renal, and hepatic function
 - D- All of the above

Fold

• First Treatment Dose _____

• Step-up Dose 2 _____

• Step-up Dose 1 _____

Dates of ELREXFIO Injections : _____

Healthcare Setting Name: _____

After Hours Phone Number: _____

Office Phone Number: _____

Name of ELREXFIO treating Oncologist: _____

⚠ This Patient has received ELREXFIO.

+ IMPORTANT SAFETY INFORMATION YOU SHOULD KNOW: ELREXFIO therapy can cause cytokine release syndrome (CRS) or neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) which may be fatal or life threatening. CRS may involve multiple organ systems.

FOR HEALTHCARE PROVIDERS

**ELREXFIO™ REMS
Patient Wallet Card**

1-844-923-7845 www.ELREXFIOREMS.com



Carry this card with you at all times.

SHOW THIS CARD to any healthcare provider involved in your care and if you go to the emergency room.



**IMPORTANT SAFETY INFORMATION FOR PATIENTS
RECEIVING TREATMENT WITH ELREXFIO**

FOR THE PATIENT

Call your healthcare provider or get emergency help right away if you have any of these symptoms:

- Fever 100.4°F (38°C) or higher
- Trouble breathing
- Chills
- Dizziness or light-headedness
- Fast heartbeat
- Headache
- Agitation, trouble staying awake, confusion or disorientation, seeing or hearing things that are not real
- Trouble speaking, thinking, remembering things, paying attention, or understanding things
- Problems walking or 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

IMPORTANT TO REMEMBER:



If you have **any** of these symptoms call your doctor or seek emergency medical attention right away! These are not all of the possible symptoms of ELREXFIO. Tell your doctor if you have any symptom that bothers you or does not go away.

You should be hospitalized after administration of each of the first 2 step-up doses within the step-up dosing schedule.

You should always ask your doctor about taking other medications while taking ELREXFIO.

Fold

Fold



Fold

Print out, cut along the dotted lines and fold in half and half again.



PHARMACY AND HEALTHCARE SETTING TRAINING PROGRAM

Risk Evaluation and Mitigation Strategy (REMS)



ELREXFIO Pharmacy and Healthcare Setting Training Module

- This training module provides details related to pharmacy and healthcare setting requirements of the ELREXFIO REMS.
- Please refer to the Prescribing Information for additional information.

1 ELREXFIO Pharmacy and Healthcare Setting Training Module



What is a REMS?

- A Risk Evaluation and Mitigation Strategy (REMS) is a program required by the FDA to manage known or potential serious risks associated with a drug product. The FDA has determined that a REMS is necessary to ensure that the benefits of ELXREXFIO outweigh its risks.
- The goal of the ELXREXFIO REMS is to mitigate the risks of Cytokine Release Syndrome (CRS) and neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) by ensuring prescribers are aware of the importance of monitoring for signs and symptoms of CRS and neurologic toxicity including ICANS in patients exposed to ELXREXFIO.

2 ELXREXFIO Pharmacy and Healthcare Setting Training Module



ELREXFIO Indication

- ELREXFIO is a B-cell maturation antigen (BCMA)-directed and CD3-directed bispecific antibody indicated for the treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy, including a proteasome inhibitor, an immunomodulatory agent, and an anti-CD38 monoclonal antibody.

3 ELREXFIO Pharmacy and Healthcare Setting Training Module



/01

Pharmacy and Healthcare Setting Certification Requirements

 **ELREXFIO**TM
(elranatamab-bcmm)
INJECTION FOR SUBCUTANEOUS USE | 44 mg/11 mL
76 mg/19 mL

Pharmacy and Healthcare Setting Certification Requirements

Requirements to become certified to dispense ELREXFIO are as follows:

- Designate an Authorized Representative to carry out the certification process and oversee implementation and compliance with the REMS requirements on behalf of the Pharmacy and/or Healthcare Setting
- Have the Authorized Representative review this **Pharmacy and Healthcare Setting Training Program**
- Have the Authorized Representative enroll in the REMS by completing and submitting the **Pharmacy and Healthcare Setting Enrollment Form** to the REMS
- Train all relevant staff involved in dispensing ELREXFIO on the REMS requirements using this **Pharmacy and Healthcare Setting Training Program**
- Before dispensing, obtain authorization to dispense each prescription by contacting the REMS to verify the prescriber is certified

5 ELREXFIO Pharmacy and Healthcare Setting Training Module

 **ELREXFIO™**
(elranatamab-bcmm)
INJECTION FOR SUBCUTANEOUS USE | 44 mg/1 mL
76 mg/1.9 mL

Pharmacy and Healthcare Setting Responsibilities

At all times:

- Report serious adverse events of CRS and neurologic toxicity including ICANS to the REMS
- Maintain records of staff training
- Maintain records that processes and procedures are in place and are being followed
- Maintain records of all ELREXFIO dispenses and provide data to the REMS and Wholesalers-Distributors, as requested.
- Comply with audits carried out by Pfizer Inc. or a third party acting on behalf of Pfizer Inc., to ensure that all training, processes, and procedures are in place and being followed
- Not distribute, transfer, loan, or sell ELREXFIO except to certified pharmacies and healthcare settings

To maintain certification to dispense:

- Have a new Authorized Representative enroll in the REMS by completing and submitting the **Pharmacy and Healthcare Setting Enrollment Form** to the REMS if the Authorized Representative changes

6 ELREXFIO Pharmacy and Healthcare Setting Training Module

 **ELREXFIO™**
(elranatamab-bcmm)
INJECTION FOR SUBCUTANEOUS USE | 44 mg/1 mL | 76 mg/1.9 mL

Who Can Be an Authorized Representative?

An Authorized Representative at the Pharmacy or Healthcare Setting can be a:

- Pharmacist
- Pharmacy Technician
- Registered Nurse
- Any responsible individual assigned by the Pharmacy or Healthcare Setting

Certified ELREXFIO Prescribers cannot be designated as an Authorized Representative for a certified Pharmacy or Healthcare Setting.

One Authorized Representative must enroll for each Pharmacy or Healthcare Setting and uphold the REMS requirements as stated on the Pharmacy and Healthcare Setting Enrollment Form

7 ELREXFIO Pharmacy and Healthcare Setting Training Module

 **ELREXFIO™**
(elranatamab-bcmm)
INJECTION FOR SUBCUTANEOUS USE | 44 mg/1 mL
76 mg/1.9 mL

Managing Pharmacy and Healthcare Setting Staff

- The Authorized Representative (AR) may grant Pharmacy and Healthcare Setting Staff access via the *Staff Management* tab in the ELREXFIO REMS Portal
- Once the AR has added the Staff member through this page, the Staff member will receive an automated e-mail
- The Staff member will then have the ability to access the Portal to confirm Prescriber Certification prior to dispensing ELREXFIO

8 ELREXFIO Pharmacy and Healthcare Setting Training Module



How to Confirm Prescriber Certification

- When an order for ELREXFIO is received, the Authorized Representative or designated staff will log into the ELREXFIO REMS portal at www.ELREXFIOREMS.com
 - Select **REMS Dispense Authorization (RDA)**
 - After entering either the ordering Prescriber's NPI# or name, one of the following certification statuses will appear:
 - CERTIFIED: OK TO DISPENSE
 - NOT CERTIFIED: DO NOT DISPENSE
 - The RDA code will display for documentation purposes
- ELREXFIO may only be dispensed upon generation of an RDA
- The Authorized Representative or designated staff may also contact the ELREXFIO REMS Coordinating Center via phone at 1-844-923-7845 to obtain an RDA

Adverse Event Reporting

- Reporting of suspected adverse events following administration of therapy is vital for the continued monitoring of the risk/benefit balance of therapy.
- Pharmacy and healthcare setting staff must report serious adverse events of CRS and neurologic toxicity including ICANS to the REMS Coordinating Center at 1-844-923-7845. Report all other suspected adverse events associated with ELREXFIO to Pfizer Inc. at 1-800-438-1985 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

This label may not be the latest approved by FDA.
For current labeling information, please visit <https://www.fda.gov/drugsatfda>



 **ELREXFIO**[™]
(elranatamab-bcmm)
INJECTION FOR SUBCUTANEOUS USE | 44 mg/1.1 mL
76 mg/1.9 mL



**For further information, please visit
www.ELREXFIOREMS.com or
call 1-844-923-7845**





FDA - REQUIRED REMS SAFETY INFORMATION

- Risk of Cytokine Release Syndrome (CRS) and neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) with ELREXFIO
- Required REMS Certification to prescribe ELREXFIO

Dear Healthcare Provider:

The purpose of this letter is to inform you of important safety information for ELREXFIO based on current prescribing information. The FDA has required this safety notice as part of the ELREXFIO REMS (**Risk Evaluation and Mitigation Strategy**) to inform you about the serious risks of CRS and neurologic toxicity including ICANS.

Serious Risks of ELREXFIO

- **Cytokine release syndrome (CRS)**, including life-threatening or fatal reactions, can occur in patients receiving ELREXFIO.
- Initiate treatment with the ELREXFIO step-up dosing schedule to reduce the risk of CRS.
- **Neurologic toxicity including ICANS** and serious and life-threatening reactions, can occur in patients receiving ELREXFIO.
- Monitor patients for signs or symptoms of neurologic toxicity including ICANS during treatment.
- Withhold ELREXFIO until CRS or neurologic toxicity including ICANS resolves or permanently discontinue ELREXFIO based on severity.

Enclosed for your review and awareness of these serious risks is the ELREXFIO **REMS Fact Sheet**, a non-promotional educational document reviewed by the FDA. The ELREXFIO **REMS Fact Sheet** will provide you with more detailed information about these serious risks and the ELREXFIO REMS requirements.

REMS Requirements

- Those who prescribe and/or dispense ELREXFIO must be aware of how to manage the risks of CRS and neurologic toxicity including ICANS.
- ELREXFIO is ONLY prescribed and/or dispensed by certified prescribers, pharmacies, and healthcare settings.
- Prescribers must counsel patients on signs and symptoms of CRS and neurologic toxicity including ICANS.
- Prescribers must complete and provide patients or their caregivers with the **Patient Wallet Card** before treatment initiation (first step-up dose).



Prescriber Knowledge, Attitude, and Behavior (KAB) Survey

The Prescriber KAB Survey is a part of the REMS commitment made by Pfizer Inc., to the FDA to help assess stakeholder knowledge as it pertains to the risks associated with the use of ELREXFIO and in determining if the ELREXFIO REMS is meeting its goal.

We ask that you consider participating in this voluntary KAB survey upon receipt of the survey invitation. The timetable in completing this Prescriber KAB Survey will be **[TIMETABLE TO BE INCLUDED]**.

Indication

ELREXFIO is a B-cell maturation antigen (BCMA)-directed and CD3-directed bispecific antibody indicated for the treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy, including a proteasome inhibitor, an immunomodulatory agent, and an anti-CD38 monoclonal antibody.

Adverse Event Reporting

Healthcare Providers must report serious adverse events of CRS and neurologic toxicity including ICANS to the REMS Coordinating Center at 1-844-923-7845.

Healthcare providers should report all other suspected adverse events to Pfizer Inc. at 1-800-438-1985 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

For a complete safety profile of ELREXFIO, please see the [Prescribing Information](#) included.

For additional details about the REMS, please visit www.ELREXFIOREMS.com or contact the ELREXFIO REMS Coordinating Center at 1-844-923-7845.

Sincerely,
Pfizer Inc.



FDA - REQUIRED REMS SAFETY INFORMATION

- Risk of Cytokine Release Syndrome (CRS) and neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) with ELREXFIO
- Required REMS Certification to prescribe ELREXFIO

Dear Professional Society,

We request that you share this safety information with your members.

The purpose of this letter is to inform you of important safety information for ELREXFIO based on current prescribing information. The FDA has required this safety notice as part of the ELREXFIO REMS (Risk Evaluation and Mitigation Strategy) to inform you about the serious risks of CRS and neurologic toxicity including ICANS.

Serious Risks of ELREXFIO

- **Cytokine release syndrome (CRS)**, including life-threatening or fatal reactions, can occur in patients receiving ELREXFIO.
- Initiate treatment with the ELREXFIO step-up dosing schedule to reduce the risk of CRS.
- **Neurologic toxicity including ICANS** and serious and life-threatening reactions, can occur in patients receiving ELREXFIO.
- Monitor patients for signs or symptoms of neurologic toxicity including ICANS during treatment.
- Withhold ELREXFIO until CRS or neurologic toxicity including ICANS resolves or permanently discontinue ELREXFIO based on severity.

Enclosed for your review and awareness of these serious risks is the ELREXFIO **REMS Fact Sheet**, a non-promotional educational document reviewed by the FDA. The ELREXFIO **REMS Fact Sheet** will provide you with more detailed information about these serious risks and the ELREXFIO REMS requirements.

REMS Requirements

- Those who prescribe and/or dispense ELREXFIO must be aware of how to manage the risks of CRS and neurologic toxicity including ICANS.
- ELREXFIO is ONLY prescribed and/or dispensed by certified prescribers, pharmacies, or healthcare settings.
- Prescribers must counsel patients on signs and symptoms of CRS and neurologic toxicity including ICANS.
- Prescribers must complete and provide the **Patient Wallet Card** to the patient or caregiver before treatment initiation (first step-up dose).



Indication

ELREXFIO is a B-cell maturation antigen (BCMA)-directed and CD3-directed bispecific antibody indicated for the treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy, including a proteasome inhibitor, an immunomodulatory agent, and an anti-CD38 monoclonal antibody.

For a complete safety profile of ELREXFIO, please see the [Prescribing Information](#) included. For additional details about the REMS, please visit www.ELREXFIOREMS.com or contact the ELREXFIO REMS Coordinating Center at 1-844-923-7845.

Sincerely,
Pfizer Inc.

ELREXFIO™ REMS Fact Sheet



FDA - REQUIRED REMS SAFETY INFORMATION

ELREXFIO REMS Overview

- The ELREXFIO **R**isk **E**valuation and **M**itigation **S**trategy (REMS) is a safety program that manages the risks of Cytokine Release Syndrome (CRS) and Neurologic Toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS). The ELREXFIO REMS is required by the Food and Drug Administration (FDA) to ensure that the benefits of the drug outweigh its risks.
- Healthcare providers, pharmacies, and healthcare settings that prescribe and/or dispense ELREXFIO must be specially certified and trained on how to manage the risks of CRS and neurologic toxicity including ICANS.
- Patients or their caregivers must receive the **Patient Wallet Card** before the first step-up dose of ELREXFIO.
- Wholesalers-distributors must ONLY distribute ELREXFIO to certified pharmacies and healthcare settings.

What Are the Risks?

- CRS, including life-threatening or fatal reactions, can occur in patients receiving ELREXFIO. Initiate treatment with the ELREXFIO step-up dosing schedule to reduce the risk of CRS.
- Neurologic toxicity, including ICANS, and serious and life-threatening reactions, can occur in patients receiving ELREXFIO. Monitor patients for signs or symptoms of neurologic toxicity including ICANS during treatment.

How Can Healthcare Providers Manage the Risks?

- Follow the ELREXFIO step-up dosing schedule as outlined in the **Prescriber Training Program**.
- Administer pretreatment medications approximately 1 hour before each dose in the step-up dosing schedule (step-up dose 1, step-up dose 2, and the first treatment dose) of ELREXFIO to reduce the risk of CRS as outlined in the **Prescriber Training Program**.
- Complete and provide patients or their caregivers with the **Patient Wallet Card** before treatment initiation (first step-up dose).
- Counsel patients about the risks of CRS and neurologic toxicity including ICANS and to seek medical attention should signs or symptoms of CRS and neurologic toxicity occur.
- Advise patients not to drive or operate heavy or potentially dangerous machinery for 48 hours after completing each of the 2 step-up doses and the first treatment dose within the ELREXFIO step-up dosing schedule and in the event of new onset of any neurological toxicity symptoms until symptoms resolve.
- Instruct patients that they should be hospitalized for 48 hours after administration of the first step-up dose, and for 24 hours after administration of the second step-up dose.
- Monitor patients for signs or symptoms of CRS and of neurologic toxicity during treatment.
 - At the first sign of CRS, immediately evaluate the patient for hospitalization. Administer supportive therapy based on severity and consider further management per current practice guidelines.
 - At the first sign of neurologic toxicity including ICANS immediately evaluate the patient and provide supportive therapy based on severity.

ELREXFIO™ REMS Fact Sheet



ELREXFIO REMS Overview (continued)

- Withhold ELREXFIO until CRS or neurologic toxicity including ICANS resolves or permanently discontinue ELREXFIO based on severity, as indicated in the Prescribing Information.
- For further details on recommended actions to be taken and treatment guidance for CRS and neurologic toxicity including ICANS refer to the **Prescriber Training Program** and **Adverse Reaction Management Guide**.

Key Requirements of the ELREXFIO REMS

Healthcare providers who prescribe ELREXFIO



Receive training on the REMS requirements at www.ELREXFIOREMS.com by using the **Prescriber Training Program** and the **Adverse Reaction Management Guide**.



Successfully complete the **Knowledge Assessment** and submit it to the REMS online.

Enroll in the REMS by completing and submitting the **Prescriber Enrollment Form** online to the REMS. Fax and e-mail options are also available.



If ELREXFIO will be dispensed and administered in the same location, an Authorized Representative must complete the Pharmacy and Healthcare Setting certification.



Instruct patients that they should be hospitalized for 48 hours after administration of the first step-up dose, and for 24 hours after administration of the second step-up dose. Monitor patients for signs and symptoms of CRS and neurologic toxicity including ICANS during treatment with ELREXFIO.

Pharmacies and Healthcare Settings that dispense ELREXFIO



Designate an Authorized Representative to carry out the certification process and oversee implementation and compliance with the REMS requirements.



NOTE: Certified ELREXFIO Prescribers cannot be designated as an Authorized Representative for a certified Pharmacy or Healthcare Setting.



Authorized Representative must be trained at www.ELREXFIOREMS.com using the **Pharmacy and Healthcare Setting Training Program**.



Authorized Representative must enroll in the REMS by completing and submitting the **Pharmacy and Healthcare Setting Enrollment Form** online to the REMS. Fax and e-mail options are also available.



Pharmacies and Healthcare Settings must verify prescriber certification in the ELREXFIO REMS before dispensing ELREXFIO.

ELREXFIO™ REMS Fact Sheet



Key Requirements of the ELREXFIO REMS (continued)

Patients



Receive the **Patient Wallet Card** before treatment.



Should be hospitalized for 48 hours after administration of the first step-up dose, and for 24 hours after administration of the second step-up dose.

Adverse Event Reporting

Healthcare Providers must report serious adverse events of CRS and neurologic toxicity including ICANS to the REMS Coordinating Center at 1-844-923-7845.

Healthcare providers should report all other suspected adverse events to Pfizer Inc. at 1-800-438-1985 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

For more information, to enroll in the ELREXFIO REMS, and for all REMS materials go to www.ELREXFIOREMS.com.



ELREXFIO™ Risk Evaluation and Mitigation Strategy (REMS)

The goal of the ELREXFIO REMS is to mitigate the risks of Cytokine Release Syndrome (CRS) and neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) by ensuring prescribers are aware of the importance of monitoring for signs and symptoms of CRS and neurologic toxicity including ICANS in patients exposed to ELREXFIO.

Prescribers

Prescribers must be certified in the ELREXFIO REMS to treat patients with ELREXFIO.

[Learn More >](#)

Pharmacies and Healthcare Settings

Pharmacies and Healthcare Settings must be certified in the ELREXFIO REMS to dispense ELREXFIO to patients.

[Learn More >](#)

Patients

All patients treated with ELREXFIO should receive a **Patient Wallet Card**. Patients should carry the **Patient Wallet Card** to remind them of the signs and symptoms of CRS and neurologic toxicity including ICANS and when to seek immediate medical attention. Patients should share this card with any healthcare provider who provides care to them to inform them of receipt of ELREXFIO treatment and when to contact the patient's oncologist.

Resources for Patients

- [Patient Wallet Card - English](#)
- [Patient Wallet Card - Spanish](#)

Resources for Prescribers

- [Prescriber Training Program](#)
- [Adverse Reaction Management Guide](#)
- [Knowledge Assessment](#)
- [Prescriber Enrollment Form](#)
- [Patient Wallet Card - English](#)
- [Patient Wallet Card - Spanish](#)
- [Fact Sheet](#)
- [Healthcare Provider Letter](#)
- [Professional Society Letter](#)

[Download All Prescriber Resources](#)

Resources for Pharmacies and Healthcare Settings

- [Pharmacy and Healthcare Setting Training Program](#)
- [Pharmacy and Healthcare Setting Enrollment Form](#)

[Download All Pharmacy and Healthcare Setting Resources](#)

Have Questions?

Contact the ELREXFIO REMS by calling 1-844-923-7845.

What is the ELREXFIO REMS?

A Risk Evaluation and Mitigation Strategy (REMS) is a safety program to manage known or potential serious risks associated with a drug product and is required by the Food and Drug Administration (FDA).

The FDA has determined that a REMS is necessary to ensure that the benefits of ELREXFIO outweigh the risks of cytokine release syndrome and neurologic toxicity including ICANS.

Indication

ELREXFIO is a B-cell maturation antigen (BCMA)-directed and CD3-directed bispecific antibody indicated for the treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy, including a proteasome inhibitor, an immunomodulatory agent, and an anti-CD38 monoclonal antibody.

Reporting Adverse Events

To report serious adverse events of CRS or neurologic toxicity including ICANS, contact the REMS Coordinating Center at 1-844-923-7845.

To report all other suspected adverse events, contact Pfizer Inc. at 1-800-438-1985 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

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Prescribers

ELREXFIO REMS Requirements

All Healthcare Providers involved in the prescribing of ELREXFIO are trained on the ELREXFIO REMS requirements and must successfully complete the **Knowledge Assessment**.

Prescribers must enroll and become certified in the ELREXFIO REMS to be able to prescribe ELREXFIO .

To become enrolled in the ELREXFIO REMS, Prescribers MUST:

| STEP 1 | STEP 2 | STEP 3 | STEP 4 |
|---|--|---|---|
| Review the following materials: <ul style="list-style-type: none">• Prescribing Information• Prescriber Training Program• Adverse Reaction Management Guide | Successfully complete the Knowledge Assessment and submit it to the REMS | Complete the Prescriber Enrollment Form and submit it to the REMS | Before treatment initiation (first step-up dose), counsel patients and/or their caregivers using the Patient Wallet Card . Complete and provide patients or their caregivers with the Patient Wallet Card . |

Resources for Prescribers

- [Prescriber Training Program](#)
- [Adverse Reaction Management Guide](#)
- [Knowledge Assessment](#)
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Reference ID: 5226170



Pharmacies and Healthcare Settings

ELREXFIO REMS Requirements

All Pharmacies and Healthcare Settings involved in the dispensing of ELREXFIO must become certified in the ELREXFIO REMS.

How does a Pharmacy and Healthcare Setting become certified in the ELREXFIO REMS?



Resources for Pharmacies and Healthcare Settings

[Pharmacy and Healthcare Setting Training Program](#)

[Pharmacy and Healthcare Setting Enrollment Form](#)

[Download All Pharmacy and Healthcare Setting Resources](#)

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ELREXFIO™ REMS Pharmacy and Healthcare Setting Enrollment Form

Instructions

To become certified in the ELREXFIO REMS and dispense ELREXFIO, a pharmacy/healthcare setting must designate an ***Authorized Representative** to:

- Review the **Pharmacy and Healthcare Setting Training Program**
- Complete this **Pharmacy and Healthcare Setting Enrollment Form**:
 - below for immediate enrollment
 - by e-mail at ELREXFIOREMS@ubc.com or
 - via fax to 1-800-349-5131.
- The ELREXFIO REMS will verify that the **Pharmacy and Healthcare Setting Enrollment Form** is complete and provide confirmation of certification via e-mail after processing.
- ELREXFIO cannot be dispensed until the Pharmacy and Healthcare Setting certification is complete.

***Certified ELREXFIO Prescribers cannot be designated as the Authorized Representatives for a Pharmacy or Healthcare Setting.**

(Fields marked with an * are REQUIRED)

Pharmacy and Healthcare Information

***Pharmacy/Healthcare Setting National Provider Identifier (NPI)#**

CONTINUE

If you do not have an NPI, please reach out to the ELREXFIO REMS by calling 1-844-923-7845

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(Fields marked with an * are REQUIRED)

Pharmacy and Healthcare Information

*Pharmacy/Healthcare Setting National Provider Identifier (NPI)#

1234567890

*Enrollment Type (Please Check One)

- New Certification Change in Authorized Representative

*Pharmacy/Healthcare Setting Name

University Hospital

DEAR (On file with distributor account)

*Pharmacy/Healthcare Setting Type

- Inpatient Hospital Pharmacy Outpatient Hospital Pharmacy Oncology Infusion Center
 Community Oncology Physician Office Specialty Pharmacy Other (please specify)

*Address Line 1

Address Line 2

*City

*State

— Please Select —

*Zip Code

*Phone

*Fax

Ship To Information

*Ship To Contact Name

Shipping Address - Same as above

*Address Line 1

Address Line 2

*City

*State

— Please Select —

*Zip Code

*Phone

*Fax

Authorized Representative Information

*E-Mail

*First Name

*Last Name

*Credentials (select one)

- PharmD R.Ph. RN
 Other (please specify: e.g., MD, DO, Office Administrator, Practice Manager)

*Phone

*Fax

Your pharmacy or healthcare setting's information will be shared with Pfizer Inc.'s wholesaler-distributor partners, to allow your pharmacy or healthcare setting to purchase product.

Authorized Representative Responsibilities

As the Authorized Representative, I must:

- Review the **Pharmacy and Healthcare Setting Training Program**.
- Enroll in the REMS by completing and submitting this **Pharmacy and Healthcare Setting Enrollment Form** to the REMS.
- Train all relevant staff involved in dispensing ELREXFIO on the REMS requirements using the **Pharmacy and Healthcare Setting Training Program**.
- Oversee implementation and compliance with the REMS requirements on behalf of the pharmacy and/or healthcare setting.

Before dispensing, staff must:

- Obtain authorization to dispense each prescription by contacting the REMS to verify the prescriber is certified.

At all times, staff must:

- Report serious adverse events of Cytokine Release Syndrome (CRS) and neurologic toxicity including Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) to the REMS.
- Not distribute, transfer, loan or sell ELREXFIO except to certified pharmacies and healthcare settings.
- Maintain records of staff training.
- Maintain records that processes and procedures are in place and are being followed.
- Maintain records of all ELREXFIO dispenses and provide data to the REMS and Wholesalers-Distributors, as requested.
- Comply with audits carried out by Pfizer Inc. or a third party acting on behalf of Pfizer Inc. to ensure that all training, processes, and procedures are in place and are being followed.
- Have a new Authorized Representative enroll in the REMS by completing and submitting the **Pharmacy and Healthcare Setting Enrollment Form** to the REMS if the Authorized Representative changes.

*Authorized Representative e-signature

*Authorized Representative Attestation

[e-Signature functionality will be available within 30 days of approval of ELREXFIO REMS]

CLEAR

CANCEL

SUBMIT



ELREXFIO™ REMS Pharmacy and Healthcare Setting Enrollment Form

Instructions

To become certified in the ELREXFIO REMS and dispense ELREXFIO, a pharmacy/healthcare setting must designate an

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(Fields marked with an * are REQUIRED)

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*Pharmacy/Healthcare Setting National Provider Identifier (NPI)#

1234567890

*Enrollment Type (Please Check One)

New Certification Change in Authorized Representative

*Pharmacy/Healthcare Setting Name

University Hospital

DEA# (On file with distributor account)

*Pharmacy/Healthcare Setting Type

Inpatient Hospital Pharmacy Outpatient Hospital Pharmacy Oncology Infusion Center
 Community Oncology Physician Office Specialty Pharmacy Other (please specify)

*Other Pharmacy/Healthcare Setting Type

Other

*Address Line 1

Address Line 2

*City

*State

*Zip Code

*Phone

*Fax

Ship To Information

*Ship To Contact Name

Shipping Address - Same as above

Authorized Representative Information

*E-Mail

*First Name

*Last Name

*Credentials (select one)

PharmD R.Ph. RN

Other (please specify: e.g., MD, DO, Office Administrator, Practice Manager)

*Other Credentials

Please Specify

*Phone

*Fax

Your pharmacy or healthcare setting's information will be shared with Pfizer Inc.'s wholesaler-distributor partners, to allow your pharmacy or healthcare setting to purchase product.

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*Authorized Representative e-signature

*Authorized Representative Attestation

[e-Signature functionality will be available within 30 days of approval of ELREXFIO REMS]

CLEAR

CANCEL

SUBMIT



ELREXFIO™ REMS Pharmacy and Healthcare Setting Enrollment Form

Thank you for submitting your information to enroll in the ELREXFIO REMS.

A confirmation of this submission has been sent to the e-mail address provided.

You can expect to receive an e-mail with a link to create your online account.

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Contact Us

Phone

1-844-923-7845

Fax

1-800-349-5131

Hours of Operation

Monday - Friday
08:00 AM – 08:00 PM
Eastern

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