

EBGLYSS- lebrikizumab-lbkz injection, solution

Eli Lilly and Company

HIGHLIGHTS OF PRESCRIBING INFORMATION

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

EBGLYSS (lebrikizumab-lbkz), injection, for subcutaneous use
Initial U.S. Approval: 2024

RECENT MAJOR CHANGES

Dosage and Administration, single-dose prefilled pen warming instructions (2.4) removed	5/2025
Dosage and Administration, single-dose prefilled syringe warming instructions (2.4) removed	10/2025

INDICATIONS AND USAGE

EBGLYSS[®] is an interleukin-13 antagonist indicated for the treatment of adults and pediatric patients 12 years of age and older who weigh at least 40 kg with moderate-to-severe atopic dermatitis whose disease is not adequately controlled with topical prescription therapies or when those therapies are not advisable. EBGLYSS can be used with or without topical corticosteroids. (1)

DOSAGE AND ADMINISTRATION

- Prior to EBGLYSS treatment, complete all age-appropriate vaccinations according to current immunization guidelines. (2.1)
- The recommended dosage of EBGLYSS is 500 mg (two 250 mg injections) at Week 0 and Week 2, followed by 250 mg (one injection) every 2 weeks until Week 16 or later, when adequate clinical response is achieved. The maintenance dose is EBGLYSS 250 mg every 4 weeks. (2.2)
- Administer by subcutaneous injection. (2.4)

DOSAGE FORMS AND STRENGTHS

Injection:

- 250 mg/2 mL in a single-dose prefilled pen (3)
- 250 mg/2 mL (125 mg/mL) in a single-dose prefilled syringe with needle shield (3)

CONTRAINDICATIONS

Prior serious hypersensitivity to lebrikizumab-lbkz or any excipients in EBGLYSS. (4)

WARNINGS AND PRECAUTIONS

- **Hypersensitivity:** Hypersensitivity reactions including angioedema and urticaria, have occurred after administration of EBGLYSS. Discontinue EBGLYSS in the event of a serious hypersensitivity reaction. (5.1)
- **Conjunctivitis and Keratitis:** Report new onset or worsening eye symptoms to a healthcare provider. (5.2)
- **Parasitic (Helminth) Infections:** Treat patients with pre-existing helminth infections before initiating EBGLYSS. If patients become infected while receiving EBGLYSS and do not respond to anti-helminth treatment, discontinue treatment with EBGLYSS until the infection resolves. (5.3)
- **Vaccinations:** Avoid use of live vaccines during treatment with EBGLYSS. (5.4)

ADVERSE REACTIONS

Most common ($\geq 1\%$) adverse reactions are conjunctivitis, injection site reactions, and herpes zoster. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Eli Lilly and Company at 1-800-LillyRx (1-800-545-5979) or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling.

Revised: 10/2025

FULL PRESCRIBING INFORMATION: CONTENTS*

1 INDICATIONS AND USAGE

2 DOSAGE AND ADMINISTRATION

- 2.1 Vaccination Prior to Administration of EBGLYSS
- 2.2 Recommended Dosage
- 2.3 Concomitant Topical Therapies
- 2.4 Important Administration Instructions
- 2.5 Missed Dose

3 DOSAGE FORMS AND STRENGTHS

4 CONTRAINDICATIONS

5 WARNINGS AND PRECAUTIONS

- 5.1 Hypersensitivity
- 5.2 Conjunctivitis and Keratitis
- 5.3 Parasitic (Helminth) Infections
- 5.4 Vaccinations

6 ADVERSE REACTIONS

- 6.1 Clinical Trials Experience

8 USE IN SPECIFIC POPULATIONS

- 8.1 Pregnancy
- 8.2 Lactation
- 8.4 Pediatric Use
- 8.5 Geriatric Use

10 OVERDOSAGE

11 DESCRIPTION

12 CLINICAL PHARMACOLOGY

- 12.1 Mechanism of Action
- 12.2 Pharmacodynamics
- 12.3 Pharmacokinetics
- 12.6 Immunogenicity

13 NONCLINICAL TOXICOLOGY

- 13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

14 CLINICAL STUDIES

- 14.1 Atopic Dermatitis

16 HOW SUPPLIED/STORAGE AND HANDLING

17 PATIENT COUNSELING INFORMATION

* Sections or subsections omitted from the full prescribing information are not listed.

FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

EBGLYSS is indicated for the treatment of adults and pediatric patients 12 years of age and older who weigh at least 40 kg with moderate-to-severe atopic dermatitis whose disease is not adequately controlled with topical prescription therapies or when those therapies are not advisable. EBGLYSS can be used with or without topical

corticosteroids.

2 DOSAGE AND ADMINISTRATION

2.1 Vaccination Prior to Administration of EBGLYSS

Complete all age-appropriate vaccinations according to current immunization guidelines [see *Warnings and Precautions (5.4)*].

2.2 Recommended Dosage

The recommended dosage of EBGLYSS is an initial dose of 500 mg (two 250 mg injections) at Week 0 and Week 2, followed by 250 mg every two weeks until Week 16 or later, when adequate clinical response is achieved. The maintenance dosage is 250 mg every four weeks [see *Clinical Studies (14.1)*].

2.3 Concomitant Topical Therapies

EBGLYSS can be used with or without topical corticosteroids (TCS). Topical calcineurin inhibitors (TCI) may be used, but reserved for sensitive areas only, such as the face, neck, intertriginous and genital areas.

2.4 Important Administration Instructions

- EBGLYSS is for subcutaneous administration.
- EBGLYSS is intended for use under the guidance of a healthcare professional. Provide proper training to patients and/or caregivers on the subcutaneous injection technique of EBGLYSS. Adult patients may self-inject, or caregivers may give EBGLYSS after training in subcutaneous injection technique. For pediatric patients, caregivers may give injections after training in subcutaneous injection technique.
- Sites for injection include the abdomen, thigh, and back of the upper arm. Administration of EBGLYSS in the back of the upper arm may be performed by a caregiver or healthcare provider.
- Alternate the injection site with each injection. Do not inject EBGLYSS within 2 inches (5 cm) of the navel or into areas where the skin is tender, bruised, red, hard, or in an area of skin that is affected by atopic dermatitis or skin lesions.
- It is not necessary to allow EBGLYSS prefilled pen or EBGLYSS prefilled syringe to warm up to room temperature before use.
- Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. EBGLYSS is a clear to opalescent, colorless to slightly yellow to slightly brown solution. Do not use if the liquid contains visible particles, is discolored or cloudy [see *Dosage Forms and Strengths (3)*, *How Supplied/Storage and Handling (16)*].
- Refer to the Instructions for Use for complete administration instructions with illustrations [see *Instructions for Use*].

2.5 Missed Dose

If a dose is missed, administer the dose as soon as possible. Thereafter, resume dosing at the regular scheduled time.

3 DOSAGE FORMS AND STRENGTHS

EBGLYSS is a clear to opalescent, colorless to slightly yellow to slightly brown solution available as follows:

- Injection: 250 mg/2 mL in a single-dose prefilled pen
- Injection: 250 mg/2 mL (125 mg/mL) in a single-dose prefilled syringe with needle shield

4 CONTRAINDICATIONS

EBGLYSS is contraindicated in patients with prior serious hypersensitivity to lebrikizumab-lbkz or any excipients of EBGLYSS [see *Warnings and Precautions (5.1)*].

5 WARNINGS AND PRECAUTIONS

5.1 Hypersensitivity

Hypersensitivity reactions, including angioedema and urticaria, have been reported with use of EBGLYSS. If a serious hypersensitivity reaction occurs, discontinue EBGLYSS and institute appropriate therapy.

5.2 Conjunctivitis and Keratitis

Conjunctivitis and keratitis adverse reactions have been reported in clinical trials.

Conjunctivitis and keratitis occurred more frequently in atopic dermatitis subjects who received EBGLYSS compared to those who received placebo. Conjunctivitis was the most frequently reported eye disorder. Most subjects with conjunctivitis or keratitis recovered during the treatment period [see *Adverse Reactions (6.1)*].

Advise patients to report new onset or worsening eye symptoms to their healthcare provider.

5.3 Parasitic (Helminth) Infections

Patients with known helminth infections were excluded from participation in clinical studies. It is unknown if EBGLYSS will influence the immune response against helminth infections by inhibiting IL-13 signaling.

Treat patients with pre-existing helminth infections before initiating treatment with EBGLYSS. If patients become infected while receiving EBGLYSS and do not respond to antihelminth treatment, discontinue treatment with EBGLYSS until the infection resolves.

5.4 Vaccinations

EBGLYSS may alter a patient's immunity and increase the risk of infection following administration of live vaccines. Prior to therapy with EBGLYSS, complete all age-appropriate vaccinations according to current immunization guidelines. Avoid use of live vaccines immediately prior to or during treatment with EBGLYSS. No data are available on the response to live vaccines.

6 ADVERSE REACTIONS

The following adverse reactions are described elsewhere in the labeling:

- Hypersensitivity [see *Warnings and Precautions (5.1)*]
- Conjunctivitis and Keratitis [see *Warnings and Precautions (5.2)*]

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying and controlled 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.

Atopic Dermatitis

The safety of EBGLYSS was evaluated across 4 randomized, double-blind, placebo-controlled, multicenter trials in subjects with moderate-to-severe atopic dermatitis including 3 phase 3 trials (ADvocate 1, ADvocate 2, ADhere) and 1 phase 2 dose ranging trial (KGAF). In these 4 trials, mean age was 37 years; 50% of subjects were male; 62% were White, 13% were Black, and 20% were Asian. In terms of co-morbid conditions, in the phase 3 trials, 30% of the subjects had asthma, 50% had allergic rhinitis, 31% had food allergy, and 14% had allergic conjunctivitis at baseline.

A total of 891 subjects were treated with EBGLYSS for at least 1 year in the atopic dermatitis development program.

ADvocate 1, ADvocate 2, and KGAF compared the safety of EBGLYSS monotherapy to placebo. ADhere compared the safety of EBGLYSS + TCS to placebo + TCS through 16 weeks. All subjects from the phase 3 trials were allowed to enroll in the long-term extension study.

Weeks 0 to 16

Table 1 summarizes the adverse reactions that occurred at a rate of at least 1% in the EBGLYSS 250 mg every 2 weeks monotherapy group, or in the EBGLYSS 250 mg every 2 weeks + TCS group, all at a higher rate than placebo during the first 16 weeks of treatment.

Table 1: Adverse Reactions Occurring in $\geq 1\%$ of the EBGLYSS Monotherapy Group or the EBGLYSS + TCS Group in the Atopic Dermatitis Trials through Week 16

Adverse Reactions	EBGLYSS Monotherapy ^a		EBGLYSS + TCS ^b	
	EBGLYSS 250 mg Q2W ^c N = 638 n (%)	Placebo N = 338 n (%)	EBGLYSS 250 mg Q2W ^c + TCS N = 145 n (%)	Placebo + TCS N = 66 n (%)
Conjunctivitis ^d	61 (10)	10 (3)	7 (5)	0
Injection Site Reactions ^e	16 (3)	4 (1)	4 (3)	1 (2)
Herpes Zoster	3 (<1)	0	2 (1)	0

^a Integrated analysis of ADvocate 1, ADvocate 2, and the phase 2 dose finding trial (KGAF)

^b Analysis of TCS concomitant therapy trial ADhere

^c EBGLYSS 500 mg at Week 0 and Week 2, followed by 250 mg every two weeks

^d Conjunctivitis cluster includes conjunctivitis, conjunctivitis allergic, and conjunctivitis bacterial

^e Injection Site Reactions cluster includes injection site-related: pain, erythema, reaction, discomfort, dermatitis, pruritus, swelling, and rash

In the monotherapy trials (ADvocate 1, ADvocate 2, and KGAF) through Week 16, the proportion of subjects who discontinued treatment due to adverse events was 2.4% in the EBGLYSS 250 mg every 2 weeks group and 1.8% in the placebo group. In the TCS trial (ADhere) through Week 16, the proportion of subjects who discontinued treatment due to adverse events was 2.1% in the EBGLYSS 250 mg every 2 weeks + TCS group and 0% in the placebo + TCS group. The most common adverse reactions leading to discontinuation of EBGLYSS compared to the placebo group were conjunctivitis and keratitis (0.6% vs. 0.3%), and injection site reactions (0.2% vs. 0) in the monotherapy trials; and conjunctivitis (0.7% vs. 0), and injection site reactions (0.7% vs. 0) in the TCS trial.

Eosinophilia

Increased post-baseline blood eosinophils were observed at a higher frequency in EBGLYSS-treated subjects compared to placebo. During the first 16 weeks, eosinophilia (>5000 cells/mcL) was observed in 0.4% in the EBGLYSS-treated subjects and 0% in subjects receiving placebo. Blood eosinophil elevations were generally transient and did not result in discontinuation.

Safety Weeks 16 to 52

Among those EBGLYSS-treated subjects who responded at Week 16 and who were re-randomized in the maintenance period of the monotherapy trials ADvocate 1 and ADvocate 2, a total of 113 and 118 subjects received EBGLYSS 250 mg every 2 weeks or every 4 weeks, respectively. The safety profile of EBGLYSS 250 mg every 4 weeks was generally consistent with EBGLYSS every 2 weeks during Weeks 16 to 52. The safety profile of EBGLYSS during maintenance treatment was generally consistent with the safety profile observed through Week 16.

Specific Adverse Drug Reactions

Conjunctivitis and Keratitis

Conjunctivitis was the most frequently reported eye disorder. Most cases of conjunctivitis and keratitis were mild or moderate in severity and recovered or resolved without treatment interruption or discontinuation.

During the initial 16-week treatment period of the monotherapy trials, conjunctivitis, including allergic conjunctivitis, was reported by 61 subjects (10%) in the EBGLYSS 250 mg every 2 weeks group and 10 subjects (3%) in the placebo group. In the TCS concomitant therapy trial, conjunctivitis was reported by 7 subjects (5%) in the EBGLYSS 250 mg every 2 weeks + TCS group compared to 0% in the placebo + TCS group. During the 16-week placebo-controlled induction period, 68 subjects reported 73 events of conjunctivitis. All events were nonserious and mild or moderate in severity. Conjunctivitis led to treatment discontinuation in 3 subjects. The exposure adjusted incidence rate of conjunctivitis for subjects treated with EBGLYSS 250 mg every 2 weeks was 30.6 events per 100 patient years through Week 16 (KGAF, ADvocate 1, ADvocate 2, ADhere).

During the maintenance treatment period of the monotherapy trials (ADvocate 1 and ADvocate 2) from 16 to 52 weeks, conjunctivitis, including allergic conjunctivitis, was reported by 2 subjects (1.8%) in the EBGLYSS 250 mg every 2 weeks group and 12 subjects (10.1%) in the EBGLYSS 250 mg every 4 weeks group, compared to 5 subjects (8.3%) in the placebo group. During the maintenance treatment period, 14 subjects treated with EBGLYSS reported 18 events of conjunctivitis. All events were mild or moderate in severity. Conjunctivitis led to treatment discontinuation in 2 subjects in the EBGLYSS 250 mg every 4 weeks group. The exposure adjusted incidence rate of conjunctivitis for subjects treated with EBGLYSS 250 mg every 2 weeks was 18.3 events per 100 patient years and for those treated with EBGLYSS 250 mg every 4 weeks was 20.6 events per 100 patient years through Week 52 (ADvocate 1, ADvocate 2, ADhere + the long-term extension study).

During the initial 16-week treatment period of the monotherapy trials, keratitis, including atopic and vernal keratoconjunctivitis, was reported by 4 subjects (0.6%) in the EBGLYSS 250 mg every 2 weeks group and 1 subject (0.3%) in the placebo group. In the TCS concomitant therapy trial, vernal keratoconjunctivitis was reported by 1 subject (0.7%) in the EBGLYSS 250 mg every 2 weeks + TCS group, compared to 0% in the placebo + TCS group. All events were nonserious and mild or moderate in severity. Keratitis led to treatment discontinuation in 2 subjects. The exposure adjusted incidence rate of keratitis for subjects treated with EBGLYSS 250 mg every 2 weeks was 2.2 events per 100 patient years through Week 16 (KGAF, ADvocate 1, ADvocate 2, ADhere).

During the maintenance treatment period of the monotherapy trials (ADvocate 1 and ADvocate 2) from 16 to 52 weeks, atopic keratoconjunctivitis was reported by 1 subject (0.8%) in the EBGLYSS 250 mg every 4 weeks group, and vernal keratoconjunctivitis was reported by 1 subject (0.9%) in the EBGLYSS 250 mg every 2 weeks group, compared to 0% in the placebo group. One (0.9%) event of severe vernal keratoconjunctivitis in an EBGLYSS 250 mg every 2 weeks subject led to treatment discontinuation. The exposure adjusted incidence rate of keratitis for subjects treated with EBGLYSS 250 mg every 2 weeks was 1.0 event per 100 patient years and for those treated with EBGLYSS 250 mg every 4 weeks was 0.7 events per 100 patient years through Week 52 (ADvocate 1, ADvocate 2, ADhere + the long-term extension study).

Injection Site Reactions

Injection site reactions were reported by 3% of the EBGLYSS group and 1% of the placebo group in the first 16 weeks of the monotherapy trials. Incidence of injection site reactions declined with continued treatment. Most events were mild or moderate and recovered without treatment discontinuation.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Available data on lebrikizumab-lbkz use in pregnant women are insufficient to evaluate for a drug-associated risk of major birth defects, miscarriage, or other adverse maternal or fetal outcomes. Monoclonal antibodies are actively transported across the placenta (*see Clinical Considerations*). In animal reproduction studies, no effects on embryo-fetal

development were observed after subcutaneous administration of lebrikizumab-lbkz to cynomolgus monkeys during organogenesis at doses up to 18 times the human exposure at the maximum recommended human dose (MRHD) (*see Data*).

All pregnancies have a background risk of birth defect, loss, or other adverse outcomes. The background risk of major birth defects and miscarriage for the indicated population is unknown. 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.

Report pregnancies to Eli Lilly and Company at 1-800-LillyRx (1-800-545-5979).

Clinical Considerations

Fetal/Neonatal Adverse Reactions

Transport of endogenous IgG antibodies across the placenta increases as pregnancy progresses and peaks during the third trimester. Therefore, EBGLYSS may be present in infants exposed in utero. The potential clinical impact of EBGLYSS exposure in infants exposed in utero should be considered.

Data

Animal Data

In an embryofetal development study, no malformations or embryofetal toxicity were observed in fetuses from pregnant cynomolgus monkeys administered lebrikizumab-lbkz during organogenesis at doses up to 150 mg/kg initial dose followed by 50 mg/kg per week by subcutaneous injection, which was associated with plasma exposure ($C_{avg,ss}$) approximately 18 times the human exposure at the MRHD. Lebrikizumab-lbkz crossed the placenta in monkeys.

In a prenatal and postnatal development study, pregnant cynomolgus monkeys were administered lebrikizumab-lbkz during organogenesis to parturition at doses up to 150 mg/kg initial dose followed by 50 mg/kg per week by subcutaneous injection, which was associated with plasma exposure ($C_{avg,ss}$) approximately 18 times the human exposure at the MRHD. No embryofetal toxicity or malformations, or effects on morphological, functional, or immunological development were observed in the infants from birth through 6 months of age.

8.2 Lactation

Risk Summary

There are no data on the presence of lebrikizumab-lbkz in human milk, the effects on the breastfed infant, or the effects on milk production. Endogenous IgG and monoclonal antibodies are transferred in human milk. The effects of local gastrointestinal exposure and limited systemic exposure in the breastfed infant to lebrikizumab-lbkz are unknown. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for EBGLYSS and any potential adverse effects on the breastfed infant from EBGLYSS or from the underlying maternal condition.

8.4 Pediatric Use

The safety and effectiveness of EBGLYSS have been established in pediatric patients 12 years of age and older who weigh at least 40 kg with moderate-to-severe atopic

dermatitis whose disease is not adequately controlled with topical prescription therapies or when those therapies are not advisable. A total of 372 pediatric subjects were exposed to EBGLYSS with 270 subjects exposed to EBGLYSS for at least one year. The safety and effectiveness were generally consistent between pediatric and adult subjects [see *Adverse Reactions (6.1)*, *Clinical Studies (14.1)*].

The safety and effectiveness of EBGLYSS have not been established in pediatric patients younger than 12 years of age and pediatric patients 12 years and older who weigh less than 40 kg.

8.5 Geriatric Use

Of the 1348 adult subjects with moderate-to-severe atopic dermatitis exposed to EBGLYSS, a total of 123 were 65 years or older, and 29 subjects were 75 years or older. Clinical studies of EBGLYSS did not include sufficient numbers of subjects 65 years of age and older to determine whether they respond differently from younger adult subjects [see *Clinical Pharmacology (12.3)*].

10 OVERDOSAGE

In the event of overdose, contact Poison Control (1-800-222-1222) for the latest recommendations and monitor the patient for any signs or symptoms of adverse reactions and institute appropriate symptomatic treatment immediately.

11 DESCRIPTION

Lebrikizumab-lbkz, an interleukin-13 antagonist, is an immunoglobulin G4 (IgG4) monoclonal antibody that binds to interleukin (IL)-13 and inhibits IL-13 signaling. Lebrikizumab-lbkz is produced in Chinese Hamster Ovary (CHO) cells by recombinant DNA technology. Lebrikizumab-lbkz has an approximate molecular weight of 145 kDa.

EBGLYSS (lebrikizumab-lbkz) injection is a sterile, preservative free, clear to opalescent, colorless to slightly yellow to slightly brown solution for subcutaneous use. EBGLYSS is available as either a 250 mg/2 mL single-dose prefilled pen or a single-dose prefilled syringe with needle shield. The EBGLYSS prefilled pen and prefilled syringe with needle shield are not made with natural rubber latex.

Each prefilled pen or prefilled syringe delivers 250 mg lebrikizumab-lbkz in 2 mL solution which also contains glacial acetic acid (1.8 mg), histidine (6.2 mg), polysorbate 20 (0.6 mg), sucrose (119.6 mg) and Water for Injection. The pH is 5.4 – 6.0.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Lebrikizumab-lbkz is an IgG4 monoclonal antibody that binds with high affinity and slow off-rate to interleukin (IL)-13 and allows IL-13 to bind to IL-13R α 1 but inhibits human IL-13 signaling through the IL-4R α /IL-13R α 1 receptor complex. IL-13 is a naturally occurring cytokine that is involved in Type 2 inflammation, which is an important component in the pathogenesis of atopic dermatitis. Lebrikizumab-lbkz inhibits IL-13-induced responses including the release of proinflammatory cytokines, chemokines and

IgE. Lebrikizumab-lbkz-bound IL-13 can still bind IL-13R α 2 allowing subsequent internalization and natural clearance of IL-13.

12.2 Pharmacodynamics

In clinical studies, lebrikizumab-lbkz reduced the levels of serum periostin, total immunoglobulin E (IgE), CC chemokine ligand (CCL)17 [thymus and activation-regulated chemokine (TARC)], CCL18 [pulmonary and activation-regulated chemokine (PARC)], and CCL13 [monocyte chemotactic protein-4 (MCP-4)]. The clinical relevance of these biomarkers is not completely understood.

12.3 Pharmacokinetics

Lebrikizumab-lbkz steady-state exposure following either a subcutaneous dose of 250 mg every 2 weeks or every 4 weeks in patients with atopic dermatitis are presented in Table 2. Lebrikizumab-lbkz exposure increases dose-proportionally over a subcutaneous dose range of 37.5 to 500 mg. Lebrikizumab-lbkz steady state is achieved at Week 4 following the approved recommended loading doses.

Table 2: Lebrikizumab-lbkz Steady-State Exposure Following Subcutaneous Administration in Patients with Atopic Dermatitis

Lebrikizumab-lbkz Dosage^a	C_{max}	C_{avg}	C_{trough}
250 mg every 2 weeks	108 mcg/mL	100 mcg/mL	87 mcg/mL
250 mg every 4 weeks	63 mcg/mL	51 mcg/mL	36 mcg/mL

C_{max} = Maximum concentration, C_{avg} = Average concentration, C_{trough} = Trough concentration
^a Following approved recommended loading doses

Absorption

Following a single subcutaneous 250 mg dose of lebrikizumab-lbkz, peak serum concentrations were achieved approximately 7 to 8 days post dose. The absolute bioavailability for a subcutaneous dose was approximately 86%.

Injection site locations did not influence the absorption of lebrikizumab-lbkz.

Distribution

The lebrikizumab-lbkz steady-state volume of distribution is 5.14 L.

Metabolism/Elimination

Lebrikizumab-lbkz is expected to be degraded into small peptides and amino acids via catabolic pathways in the same manner as endogenous IgG.

The lebrikizumab-lbkz half-life is 24.5 days and clearance is 0.154 L/day. Lebrikizumab-lbkz exhibits linear elimination that is independent of dose.

Specific Populations

Age, Sex, Race

Age, sex, or race did not have a significant effect on the pharmacokinetics of lebrikizumab-lbkz.

Weight

Lebrikizumab-lbkz trough concentrations were lower in subjects with higher body weight.

Patients with Renal or Hepatic Impairment

Specific clinical pharmacology studies to evaluate the effects of renal impairment and hepatic impairment on the pharmacokinetics of lebrikizumab-lbkz have not been conducted. Lebrikizumab-lbkz, as a monoclonal antibody, is not expected to undergo significant hepatic or renal elimination. No clinically significant differences in the pharmacokinetics of lebrikizumab-lbkz were observed in patients with mild or moderate renal impairment.

Drug Interaction Studies

The effect of lebrikizumab-lbkz on the PK of co-administered medications has not been studied.

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 EBGLYSS or of other drug products.

Antibodies to lebrikizumab-lbkz developed in 4/145 (2.8%) of subjects treated with EBGLYSS 250 mg every 2 weeks followed by 250 mg every four weeks during the 12-month treatment period in EBGLYSS studies. Most of these antibodies were neutralizing and of low titer. Similar results were observed in pediatric subjects who received EBGLYSS up to 12 months. The presence of anti-drug antibodies was not associated with changes to pharmacokinetics, efficacy, or safety of lebrikizumab-lbkz. The clinical relevance of these findings is unknown because of the low occurrence of ADA.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Animal studies have not been conducted to evaluate the carcinogenic or mutagenic potential of lebrikizumab-lbkz.

No effects on fertility parameters such as reproductive organs, reproductive hormones or menstrual cycle length were observed in sexually mature female cynomolgus monkeys that were administered intravenous doses of lebrikizumab-lbkz up to 25 mg/kg/week for 37 weeks, which was associated with plasma exposure ($C_{avg,ss}$) approximately 15 times the human exposure at the MRHD. No effects on reproductive organs or sperm analysis were observed in sexually mature male cynomolgus monkeys that were administered subcutaneous doses of lebrikizumab-lbkz up to 25 mg/kg/week for 13 weeks, which was associated with plasma exposure ($C_{avg,ss}$) approximately 11 times the human exposure at the MRHD.

14 CLINICAL STUDIES

14.1 Atopic Dermatitis

Three multicenter, randomized, double-blind, placebo-controlled trials, ADvocate 1, ADvocate 2 and ADhere (NCT04146363, NCT04178967, NCT04250337) enrolled a total of 1062 subjects 12 years of age and older with moderate-to-severe atopic dermatitis not adequately controlled by topical medication(s) and who were candidates for systemic therapy. A total of 148 subjects (14%) were 12 to <18 years who weighed at least 40 kg and 914 (86%) were adult subjects. Disease severity was defined by an Investigator's Global Assessment (IGA) score ≥ 3 in the overall assessment of AD lesions on a severity scale of 0 to 4, an Eczema Area and Severity Index (EASI) score ≥ 16 on a scale of 0 to 72, and a minimum body surface area involvement of $\geq 10\%$.

At baseline, 50% of subjects were male, 63% were White, 11% were Black or African American, and 21% were Asian; 12.8% identified as Hispanic or Latino, 63% of subjects had a baseline IGA score of 3 (moderate AD) and 37% of subjects had a baseline IGA of 4 (severe AD). The baseline mean EASI was 29, and the baseline Pruritus Numeric Rating Scale (NRS) was 7 on a scale of 0-10. Of all subjects, 99% had received prior treatment for AD.

In all three trials, subjects in the EBGLYSS group received subcutaneous injections of EBGLYSS 500 mg at Week 0 and at Week 2, followed by 250 mg every other week (Q2W) through Week 16.

To evaluate the maintenance and durability of response in the monotherapy trials (ADvocate 1 and ADvocate 2), subjects originally randomized to EBGLYSS who achieved an IGA score of 0 or 1, or at least a 75% reduction in EASI from baseline [EASI-75] at Week 16 and did not require rescue therapy were re-randomized to an additional 36 weeks of either a maintenance dose of EBGLYSS 250 mg Q2W (every 2 weeks), EBGLYSS 250 mg Q4W (every 4 weeks), or placebo.

Subjects who did not achieve IGA 0 or 1 or EASI-75 at Week 16 or subjects who required rescue therapy during the first 16 weeks were treated with open-label EBGLYSS 250 mg Q2W.

In the concomitant therapy trial (ADhere), subjects received EBGLYSS + TCS or placebo + TCS. Topical calcineurin inhibitors (TCI) were permitted for sensitive areas only, such as the face, neck, intertriginous and genital areas.

All three trials assessed the primary endpoint, the proportion of subjects who achieved an IGA score of 0 (clear) or 1 (almost clear) and at least a 2-point improvement from baseline at Week 16. Other evaluated outcomes at Week 16 included the proportion of subjects with EASI-75 and EASI-90, and improvement in itch severity as defined by a reduction of at least 4 points on an 11-point Pruritus NRS. ADvocate 1 and ADvocate 2 also evaluated the maintenance and durability of response through Week 52.

The results of the EBGLYSS monotherapy trials (ADvocate 1 and ADvocate 2) are presented in Table 3.

Table 3: Efficacy Results of EBGLYSS Monotherapy at Week 16^a in ADvocate 1 and ADvocate 2 in Subjects with Moderate-to-Severe Atopic Dermatitis

	ADvocate 1			ADvocate 2		
	EBGLYSS	Placebo	Difference	EBGLYSS	Placebo	Difference

	250 mg Q2W ^b		from Placebo (95% CI)	250 mg Q2W ^b		from Placebo (95% CI)
Number of subjects	283	141	--	281	146	--
IGA 0 or 1 ^c	43%	13%	30% (22%, 38%)	33%	11%	22% (14%, 30%)
EASI-75	59%	16%	42% (33%, 51%)	52%	18%	33% (24%, 42%)
EASI-90	38%	9%	29% (21%, 36%)	31%	10%	21% (13%, 28%)
Number of subjects with baseline Pruritus NRS score ≥ 4	263	130	--	253	134	--
Pruritus NRS ≥ 4 point improvement	46%	13%	33% (25%, 41%)	40%	12%	28% (20%, 37%)

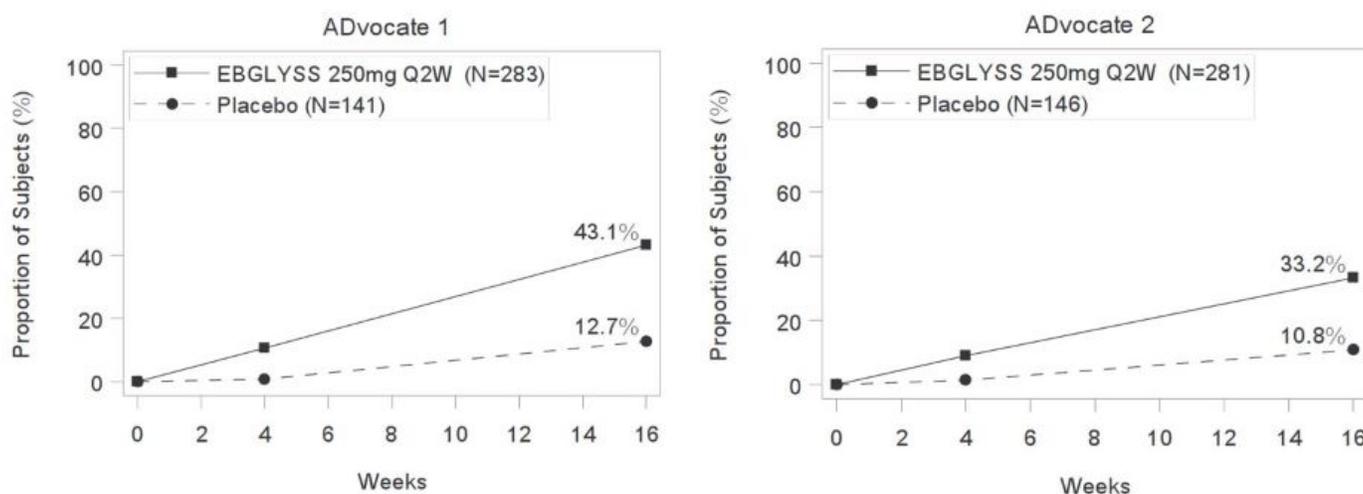
^a Subjects who received rescue therapy or discontinued treatment due to lack of efficacy were analyzed as non-responders. Data after treatment discontinuation due to any other reason were considered missing. Any missing data was imputed using MCMC-MI.

^b Subjects received 500 mg of EBGLYSS at Week 0 and Week 2, and 250 mg Q2W up to Week 16

^c Primary endpoint. Responder was defined as a subject with an IGA 0 or 1 ("clear" or "almost clear") and a reduction of ≥ 2 points on a 0-4 IGA scale

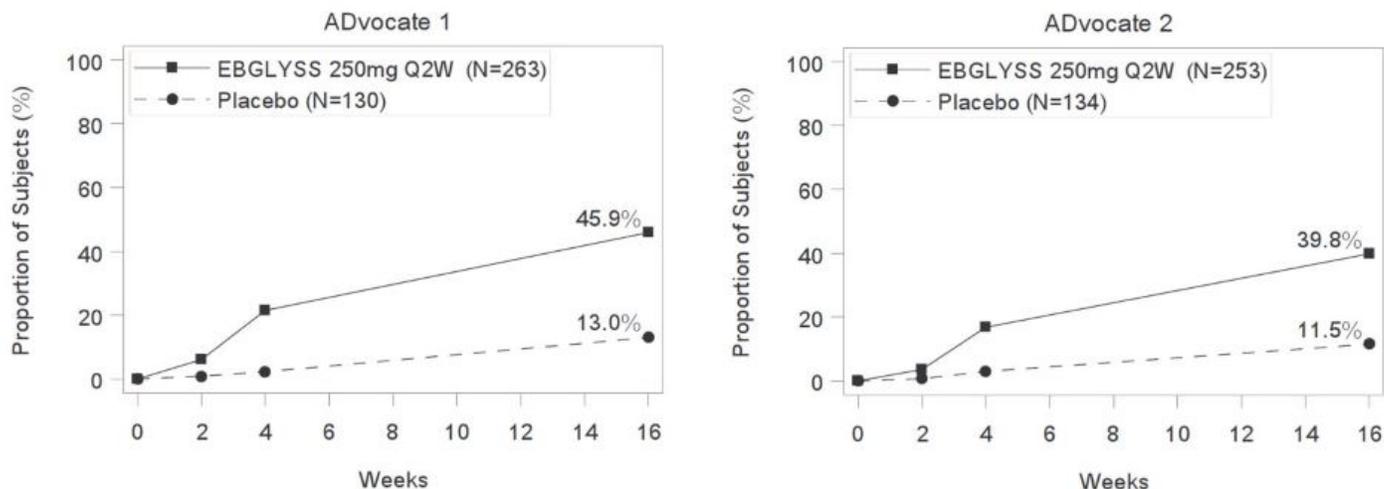
The proportion of EBGLYSS-treated subjects who achieved IGA 0 or 1 (with ≥ 2 -point improvement from baseline) by visit in ADvocate 1 and ADvocate 2 are presented in Figure 1.

Figure 1: Proportion of Subjects with Moderate-to-Severe Atopic Dermatitis achieving IGA 0 or 1, with a ≥ 2 -point improvement from baseline through Week 16 in ADvocate 1 and ADvocate 2



The proportion of EBGLYSS-treated subjects who achieved at least a 4-point improvement from baseline in Pruritus NRS by visit in ADvocate 1 and ADvocate 2 are presented in Figure 2.

Figure 2: Proportion of Subjects with Moderate-to-Severe Atopic Dermatitis with ≥ 4 -point improvement in Pruritus NRS through Week 16 in ADvocate 1 or ADvocate 2



Examination of age, sex, and White, Asian, Black or African American race subgroups did not identify differences in response to EBGLYSS among these subgroups. The database was not large enough to adequately assess differences in effects in other races.

The results in the concomitant therapy trial (ADhere) at Week 16, where subjects received EBGLYSS + TCS or placebo + TCS were consistent with the results in the monotherapy trials (ADvocate 1 and ADvocate 2).

Maintenance and Durability of Response (Week 16 to Week 52)

EBGLYSS-treated subjects achieving IGA 0 or 1 or EASI-75, and who did not receive rescue therapy at Week 16 were re-randomized to 36 weeks of maintenance treatment with EBGLYSS 250 mg Q2W, EBGLYSS 250 mg Q4W, or placebo in ADvocate 1 and ADvocate 2. The results are presented in Table 4.

Table 4: Efficacy Results of EBGLYSS at Week 52 in ADvocate 1 and ADvocate 2^a in Subjects with Moderate-to-Severe Atopic Dermatitis

	ADvocate 1			ADvocate 2		
	EBGLYSS 250 mg Q2W	EBGLYSS 250 mg Q4W	Placebo	EBGLYSS 250 mg Q2W	EBGLYSS 250 mg Q4W	Placebo
Number of subjects who were IGA of 0 or 1 Responders at Week 16^b	45	45	22	32	32	16
IGA of 0 or 1 ^b at	76.0%	71.0%	17.0%	65.0%	61.0%	50.0%

Week 52	10/0	14/0	41/0	63/0	61/0	30/0
Number of subjects who were EASI-75 Responders at Week 16	61	62	30	51	53	27
EASI-75 at Week 52	79%	79%	61%	77%	85%	72%

^a Subjects who received systemic rescue therapy, discontinued treatment due to lack of efficacy were analyzed as non-responders. Data after topical rescue medication or treatment discontinuation due to any other reason were considered missing. Any missing data were imputed using MCMC-MI.

^b Responder was defined as a subject with an IGA 0 or 1 (“clear” or “almost clear”) and a reduction of ≥ 2 points on a 0-4 IGA scale

16 HOW SUPPLIED/STORAGE AND HANDLING

How Supplied

EBGLYSS (lebrikizumab-lbkz) injection is a sterile, preservative free, clear to opalescent, colorless to slightly yellow to slightly brown solution, available in a single-dose prefilled pen or a single-dose prefilled syringe with needle shield. Each prefilled pen and prefilled syringe with needle shield is designed to deliver 250 mg of EBGLYSS in 2 mL.

EBGLYSS is supplied as:

	Pack Size	NDC
Prefilled Pen		
250 mg/2 mL single-dose	Carton of 1	0002-7772-11
Prefilled syringe with needle shield		
250 mg/2 mL (125 mg/mL) single-dose	Carton of 1	0002-7797-11

Storage and Handling

Store refrigerated at 2°C to 8°C (36°F to 46°F).

If necessary, EBGLYSS can be stored at room temperature up to 30°C (86°F) for up to 7 days in the original carton. Dispose of EBGLYSS that has been left at room temperature for longer than 7 days.

Store in the original carton to protect from light until use.

Do not freeze. Do not use EBGLYSS if it has been frozen.

Do not shake.

Do not microwave, run hot water over it, or leave it in direct sunlight.

Not made with natural rubber latex.

Discard the EBGLYSS single-dose prefilled pen or prefilled syringe with needle shield after use in a puncture-resistant container.

17 PATIENT COUNSELING INFORMATION

Advise the patient and/or caregiver to read the FDA-approved patient labeling (Patient Information and Instructions for Use).

Administration Instructions: Provide guidance to patients and caregivers on proper subcutaneous injection technique, including aseptic technique, and how to use the prefilled pen and prefilled syringe correctly. Advise patients to follow sharps disposal recommendations [see *Dosage and Administration (2.4), Instructions for Use*].

Hypersensitivity: Advise patients to discontinue EBGLYSS and to seek immediate medical attention if they experience any symptoms of systemic hypersensitivity reactions [see *Warnings and Precautions (5.1)*].

Conjunctivitis and Keratitis: Advise patients to consult their healthcare provider if new onset or worsening eye symptoms develop [see *Warnings and Precautions (5.2)*].

Parasitic (Helminth) Infections: Advise patients to notify their healthcare provider if they present with clinical features consistent with helminthic infection [see *Warnings and Precautions (5.3)*].

Vaccinations: Advise patients that EBGLYSS may increase the risk of infection following administration of live vaccines and that vaccination with live vaccines is not recommended during EBGLYSS treatment. Instruct patients to inform the healthcare provider that they are taking EBGLYSS prior to a potential vaccination [see *Warnings and Precautions (5.4)*].

Pregnancy: Inform patients to report their pregnancy to Eli Lilly and Company at 1-800-LillyRx (1-800-545-5979) [see *Use in Specific Populations (8.1)*].

Eli Lilly and Company, Indianapolis, IN 46285, USA

US License No. 1891

Copyright © 2024, 2025, Eli Lilly and Company. All rights reserved.

EBG-0005-USPI-20251028

<p style="text-align: center;">PATIENT INFORMATION EBGLYSS™ (EHB-glihs) (lebrikizumab-lbkz) injection, for subcutaneous use</p>

What is EBGLYSS?

EBGLYSS is a prescription medicine used to treat adults and children 12 years of age and older who weigh at least 88 pounds (40 kg) with moderate-to-severe eczema (atopic dermatitis) that is not well controlled with prescription therapies used on the skin (topical), or who cannot use topical therapies. EBGLYSS can be used with or without topical corticosteroids.

It is not known if EBGLYSS is safe and effective in children less than 12 years of age or in children 12 years to less than 18 years of age who weigh less than 88 pounds (40 kg).

Do not use EBGLYSS if you are allergic to lebrikizumab-lbkz or to any of the ingredients in EBGLYSS. See the end of this leaflet for a complete list of ingredients in EBGLYSS.

Before you use EBGLYSS, tell your healthcare provider about all your medical conditions, including if you:

- have a parasitic (helminth) infection
- are scheduled to receive any vaccinations. You should not receive a “live vaccine” if you are treated with EBGLYSS.
- are pregnant or plan to become pregnant. It is not known if EBGLYSS will harm your unborn baby. If you become pregnant during treatment with EBGLYSS, you or your healthcare provider can call Eli Lilly and Company at 1-800-LillyRx (1-800-545-5979) to report the pregnancy.
- are breastfeeding or plan to breastfeed. It is not known if EBGLYSS passes into your breast milk.

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

How should I take EBGLYSS?

- **See the detailed “Instructions for Use” that comes with EBGLYSS for information about how to prepare and inject EBGLYSS and how to properly store and throw away (dispose of) used EBGLYSS prefilled pens and prefilled syringes.**
- Use EBGLYSS exactly as prescribed by your healthcare provider.
- Your healthcare provider will tell you how much EBGLYSS to inject and how often to inject it.
- EBGLYSS comes as a single-dose prefilled pen or prefilled syringe with needle shield.
- EBGLYSS is given as an injection under the skin (subcutaneous injection).
- If your healthcare provider decides that you or a caregiver can give the injections of EBGLYSS, you or a caregiver should receive training on the right way to prepare and inject EBGLYSS. Do not try to inject EBGLYSS until you have been shown the right way by your healthcare provider. In children 12 years of age and older, EBGLYSS should be given by a caregiver.
- If you miss a dose of EBGLYSS, inject the missed dose as soon as possible, then inject your next dose at your regular scheduled time.
- If you inject too much EBGLYSS (overdose), get medical help or contact a Poison Center expert right away at 1-800-222-1222.
- Your healthcare provider may prescribe other medicines to use with EBGLYSS. Use the other prescribed medicines exactly as your healthcare provider tells you to.

What are the possible side effects of EBGLYSS?

EBGLYSS can cause serious side effects, including:

- **Allergic reactions. EBGLYSS can cause allergic reactions that may sometimes be severe.** Stop using EBGLYSS and tell your healthcare provider or get emergency help right away if you get any of the following signs or symptoms:
 - breathing problems or wheezing
 - swelling of the face, lips, mouth, tongue, or throat
 - hives
 - itching
 - fainting, dizziness, feeling lightheaded
 - skin rash
 - cramps in your stomach area (abdomen)
- **Eye problems.** Tell your healthcare provider if you have any new or worsening eye problems, include eye pain or changes in vision, such as blurred vision.

The most common side effects of EBGLYSS include:

- eye and eyelid inflammation, including redness, swelling, and itching
- injection site reactions
- shingles (herpes zoster)

These are not all of the possible side effects of EBGLYSS. Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

How should I store EBGLYSS?

- Store EBGLYSS in the refrigerator between 36°F to 46°F (2°C to 8°C).
- EBGLYSS can be stored at room temperature up to 86°F (30°C) for up to 7 days in the original carton. Throw away (dispose of) EBGLYSS that has been left at room temperature for longer than 7 days.
- Store EBGLYSS in the original carton to protect from light until use.
- **Do not** freeze. **Do not** shake.
- **Do not** microwave EBGLYSS, or run hot water over it, or leave it in direct sunlight.

Keep EBGLYSS and all medicines out of the reach of children.**General information about the safe and effective use of EBGLYSS.**

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. Do not use EBGLYSS for a condition for which it is not prescribed. Do not give EBGLYSS to other people, even if they have the same symptoms you have. It may harm them. You can ask your pharmacist or healthcare provider for information about EBGLYSS that is written for health professionals.

What are the ingredients in EBGLYSS?

Active ingredient: lebrikizumab-lbkz

Inactive ingredient: glacial acetic acid, histidine, polysorbate 20, sucrose, and Water for Injection.

EBGLYSS prefilled pen and prefilled syringe with needle shield are not made with natural rubber latex.

EBGLYSS is a trademark of Eli Lilly and Company.

Eli Lilly and Company, Indianapolis, IN 46285

US License Number 1891

Copyright © 2024, Eli Lilly and Company. All rights reserved.

For more information about EBGLYSS, call 1-800-545-5979 (1-800-Lilly-Rx) or go to www.ebglyss.com

This Patient Information has been approved by the U.S. Food and Drug Administration
EBG-0001-PPI-202409

Revised: 09/2024

PREFILLED PEN INSTRUCTIONS FOR USE**INSTRUCTIONS FOR USE**

EBGLYSS™ [EHB-glihs]

(lebrikizumab-lbkz)

injection, for subcutaneous use

Single-Dose Prefilled Pen

This Instructions for Use contains information on how to inject EBGLYSS.

Before you use the EBGLYSS Prefilled Pen (Pen), read and carefully follow all the step-by-step instructions.



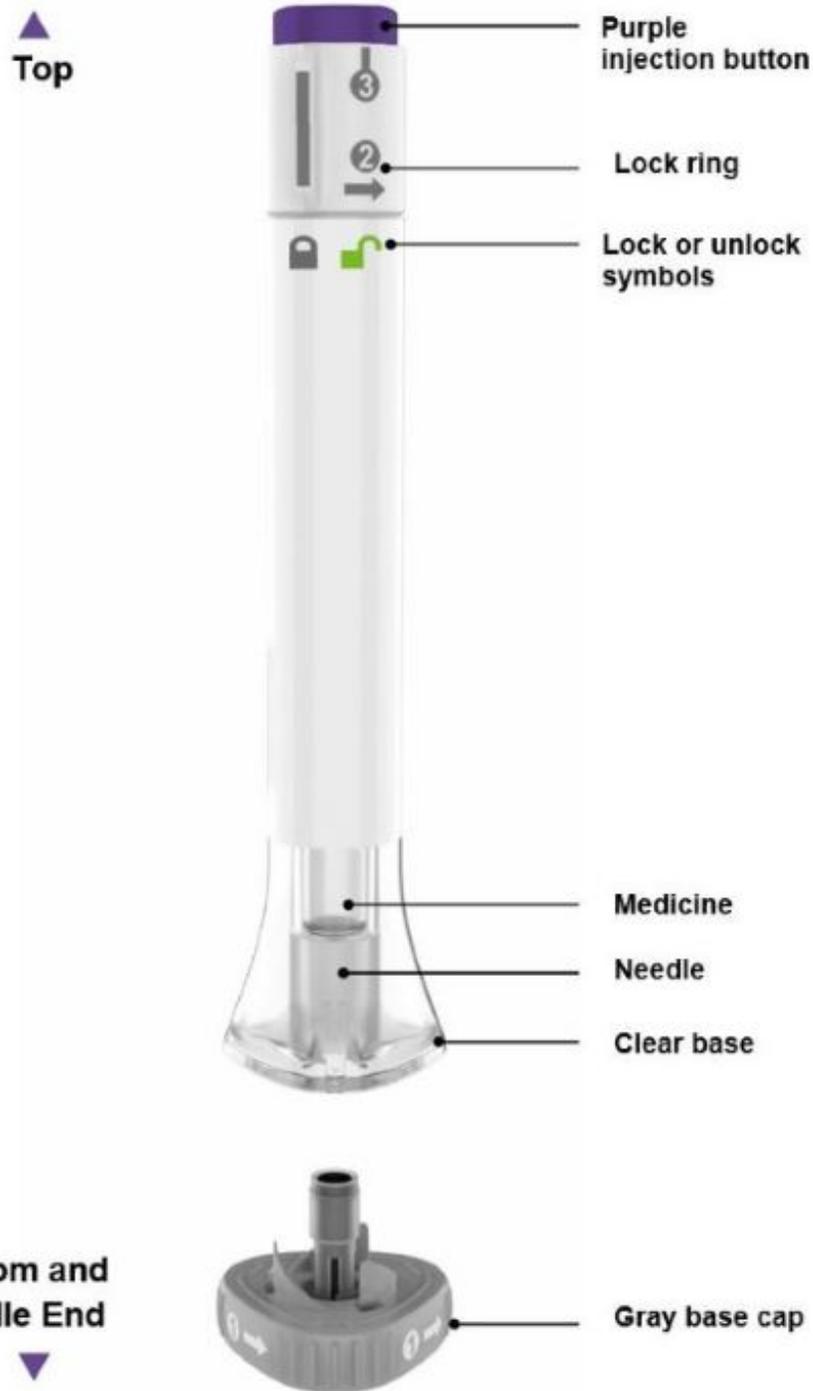
Important information you need to know before injecting EBGLYSS

- Your healthcare provider should show you how to prepare and inject EBGLYSS using the Pen. **Do not** inject yourself or someone else until you have been shown how to inject EBGLYSS.
- Keep this Instructions for Use and read it as needed.
- Each EBGLYSS Pen contains 1 dose of EBGLYSS. **The Pen is for one-time use only.**
- The EBGLYSS Pen contains glass parts. Handle it carefully. If you drop it on a hard surface, **do not** use it. Use a new EBGLYSS Pen for your injection.
- Your healthcare provider may help you decide where on your body to inject your dose. You can also read the **Choose and clean your injection site** section of these instructions to help you choose which area can work best for you.
- If you have vision or hearing problems, **do not** use EBGLYSS Pen without help from a caregiver.
- See **Storing EBGLYSS** for important storage information.

INSTRUCTIONS FOR USE

Before you use the EBGLYSS Pen, read and carefully follow all the step-by-step instructions.

Parts of the EBGLYSS Pen



Preparing to inject EBGLYSS

Gather supplies and EBGLYSS Pen:

- EBGLYSS Pen from the refrigerator
- 1 alcohol wipe
- 1 cotton ball or piece of gauze
- 1 sharps disposal container
(See **Disposing of EBGLYSS**)

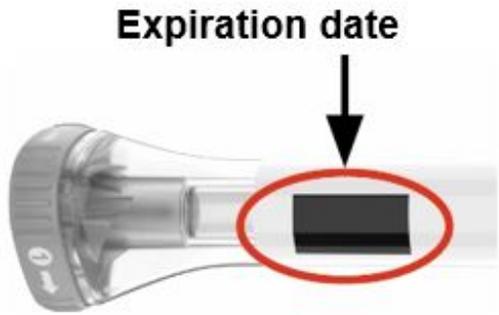
Note: You do not need to allow your Pen to warm up to room temperature before use.

Inspect the Pen and the medicine

Leave the gray base cap on until you are ready to inject. Make sure you have the right medicine. The medicine inside should be clear. It may be colorless to slightly yellow to slightly brown.

Do not use the Pen (see **Disposing of EBGLYSS**) if the:

- Pen looks damaged
- medicine is frozen
- medicine is cloudy, is discolored, or has particles
- expiration date printed on the label has passed

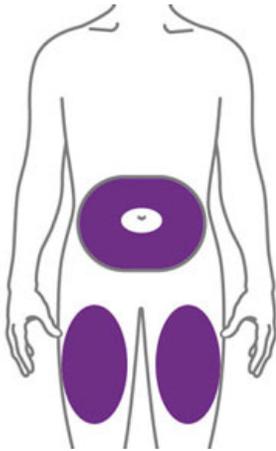


Wash your hands with soap and water

Choose and clean your injection site

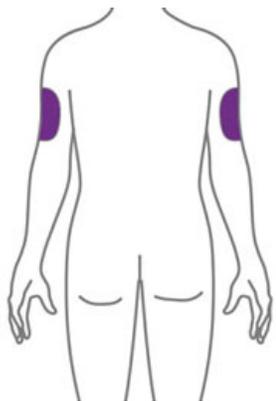
Your healthcare provider can help you choose the injection site that is best for you.

Clean the injection site with an alcohol wipe and let dry.



You or another person may inject into these areas.

- **Stomach area (abdomen) —**
At least 2 inches away from the belly button (navel).
- **Front of thigh —**
At least 2 inches above the knee and 2 inches below the groin.



Another person should inject into this area.

- **Back of upper arm —**
Another person should inject into the back of your upper arm.

Do not inject in the exact same spot every time.

Do not inject into areas where the skin is tender, bruised, red, hard, or in an area of skin that is affected by eczema (atopic dermatitis) or other skin lesions.

Injecting EBGLYSS

1 Uncap the Pen



Make sure the Pen is **locked**.



When you are ready to inject, twist

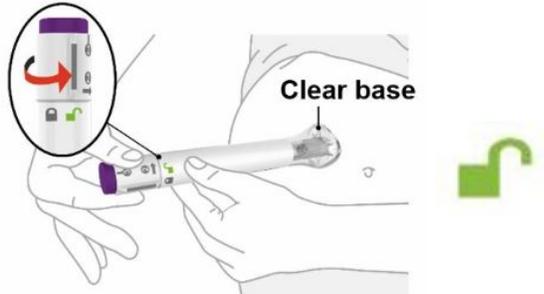


off the gray base cap and throw it away in your household trash.

Do not put the gray base cap back on; this could damage the needle.

Do not touch the needle inside the clear base.

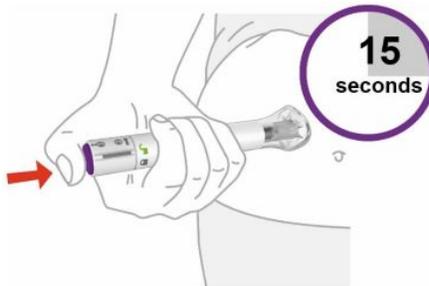
2 Place and unlock



Place and hold the clear base flat and firmly against the skin.

Keep the clear base on the skin, then turn the lock ring to the **unlock** position.

3 Press and hold for 15 seconds



Press and hold the purple injection button and **listen** for 2 loud clicks:

- First click: injection started
- Second click: injection completed

The injection may take up to 15 seconds.

You will know the injection is complete when the gray plunger is visible.



Disposing of EBGLYSS

Dispose of (throw away) the used Pen



Put the used EBGLYSS Pen in an FDA-cleared sharps disposal container right away after use.

Do not throw away (dispose of) the EBGLYSS Pen in your household trash. If you do not have an FDA-cleared sharps disposal container, you may use a household container that is:

- made of a heavy-duty plastic,
- can be closed with a tight-fitting, puncture-resistant lid, without sharps being able to come out,
- upright and stable during use,
- leak-resistant, and
- properly labeled to warn of hazardous waste inside the container.

When your sharps disposal container is almost full, you will need to follow your community guidelines for the right way to dispose of your sharps disposal container.

There may be state or local laws about how you should throw away needles and syringes.

For more information about safe sharps disposal, and for specific information about sharps disposal in the state you live in, go to the FDA's website at: <http://www.fda.gov/safesharpsdisposal>.

Do not recycle your used sharps disposal container.

Commonly asked questions

Q. What if I see air bubbles in the Pen?

A. Air bubbles are normal. They will not harm you or affect your dose.

Q. What if my Pen is not at room temperature?

A. You do not need to allow your Pen to warm up to room temperature before injecting.

Q. What if there is a drop of liquid on the tip of the needle when I remove the gray base cap?

A. A drop of liquid on the tip of the needle is normal. This will not harm you or affect your dose.

Q. What if I unlock the Pen and press the purple injection button before twisting off the gray base cap?

A. **Do not** remove the gray base cap. Throw away (dispose of) the Pen and use a new one.

Q. Do I need to hold the purple injection button down until the injection is complete?

A. You do not need to hold the purple injection button down, but it may help you keep the Pen steady and firm against your skin.

Q. What if the needle did not retract after my injection?

A. **Do not** touch the needle or replace the gray base cap. Store the Pen in a safe place to avoid an accidental needlestick and contact Lilly at 1-800-Lilly-Rx (1-800-545-5979) for instructions on how to return the Pen.

Q. What if there is a drop of liquid or blood on my skin after my injection?

A. This is normal. Press a cotton ball or gauze over the injection site. **Do not** rub the injection site.

Q. How can I tell if my injection is complete?

A. After you press the purple injection button, you will hear 2 loud clicks. The second loud click tells you that your injection is complete. You will also see the gray plunger at the top of the clear base. The injection may take up to 15 seconds.

Q. What if I remove the Prefilled Pen before the second loud click or before the gray plunger stops moving?

A. You may not have received your full dose. **Do not** give another injection. Call your healthcare provider for help.

Q. What if I heard more than 2 clicks during my injection, 2 loud clicks and 1 soft one. Did I get my complete injection?

A. Some people may hear a soft click right before the second loud click. That is the normal operation of the Prefilled Pen. **Do not** remove the Prefilled Pen from your skin until you hear the second loud click.

If you have more questions about how to use the EBGLYSS Prefilled Pen:

- Call your healthcare provider
- Call 1-800-Lilly-Rx (1-800-

Scan this code to

545-5979)

- Visit www.ebglyss.com



launch
www.ebglyss.com

Storing EBGLYSS

- Store your Pen in the refrigerator between 36°F to 46°F (2°C to 8°C).
- EBGLYSS can be stored at room temperature up to 7 days in the original carton. **Do not** store above 86°F (30°C). Throw away (dispose of) EBGLYSS that has been left at room temperature for longer than 7 days.
- Store your Pen in the original carton to protect from light until use.
- **Do not** freeze your Pen. **Do not** shake your Pen.
- **Do not** microwave your Pen, or run hot water over it, or leave it in direct sunlight.
- Throw away (dispose of) your Pen if any of the above conditions are not followed.

Keep your Pen and all medicines out of the reach of children.

Read the Patient Information insert for EBGLYSS inside this box to learn more about your medicine.

Eli Lilly and Company
Indianapolis, IN 46285, USA
US License Number 1891

EBGLYSS is a trademark of Eli Lilly and Company.
Copyright © 2024, Eli Lilly and Company. All rights reserved.

This Instructions for Use has been approved by the U.S. Food and Drug Administration.
Revised: MAY 2025
EBG-0003-PEN-IFU-20250521

PREFILLED SYRINGE INSTRUCTIONS FOR USE

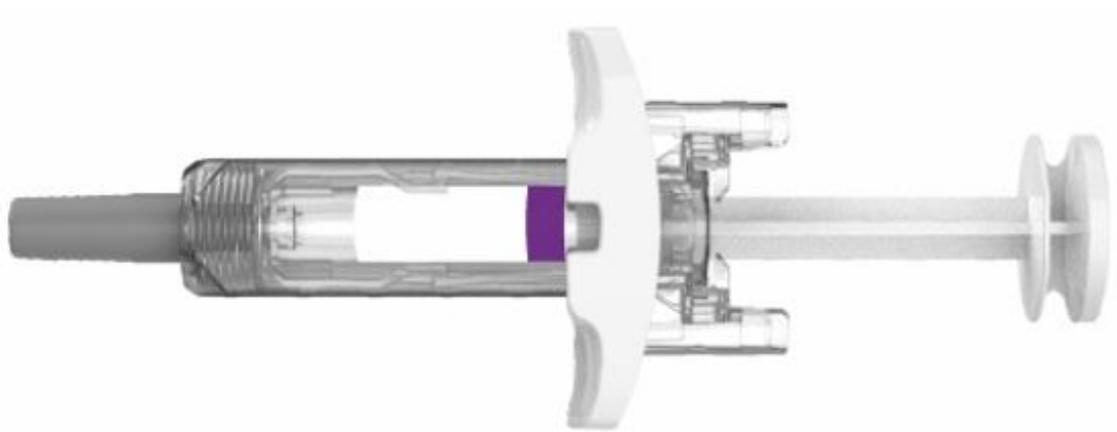
INSTRUCTIONS FOR USE

EBGLYSS™ [EHB-glihs]

(lebrikizumab-lbkz)

injection, for subcutaneous use

Single-Dose Prefilled Syringe with Needle Shield



This Instructions for Use contains information on how to inject EBGLYSS.

Before you use the EBGLYSS Prefilled Syringe with Needle Shield (Prefilled Syringe), read and carefully follow all the step-by-step instructions.

Important information you need to know before injecting EBGLYSS

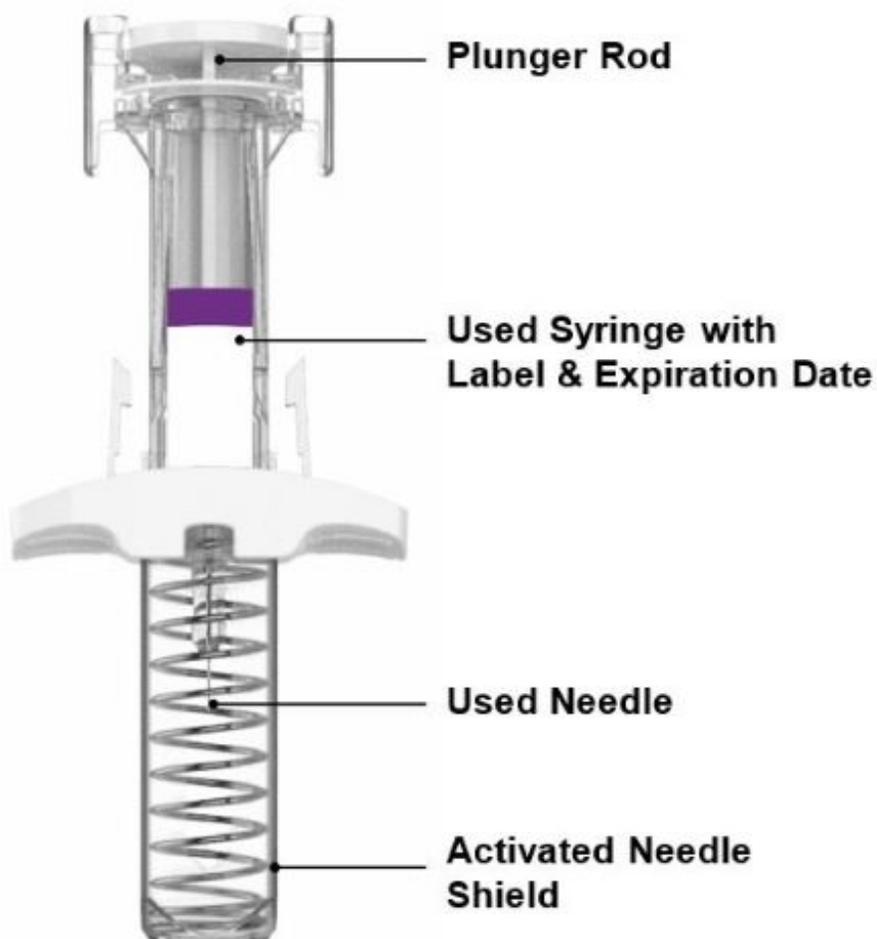
- Your healthcare provider should show you how to prepare and inject EBGLYSS using the Prefilled Syringe. **Do not** inject yourself or someone else until you have been shown how to inject EBGLYSS.
- Keep these Instructions for Use and read it as needed.
- Each EBGLYSS Prefilled Syringe contains 1 dose of EBGLYSS. **The Prefilled Syringe is for one-time use only.**
- The EBGLYSS Prefilled Syringe contains glass parts. Handle it carefully. If you drop it on a hard surface, **do not** use it. Use a new EBGLYSS Prefilled Syringe for your injection.
- Your healthcare provider may help you decide where on your body to inject your dose. You can also read the **Choose and clean your injection site** section of these instructions to help you choose which area can work best for you.
- If you have vision problems, **do not** use the EBGLYSS Prefilled Syringe without help from a caregiver.
- See **Storing EBGLYSS** for important storage information.

INSTRUCTIONS FOR USE

Before you use the EBGLYSS Prefilled Syringe, read and carefully follow all the step-by-step instructions.

Parts of the EBGLYSS Prefilled Syringe with Needle Shield

After Use



Preparing to inject EBGLYSS

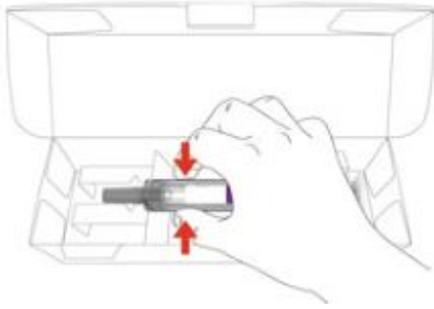
Gather Supplies and EBGLYSS Prefilled Syringe

- EBGLYSS Prefilled Syringe from the refrigerator
- 1 alcohol wipe
- 1 cotton ball or piece of gauze
- 1 sharps disposal container (see **Disposing of EBGLYSS**)

Note: You do not need to allow your Prefilled Syringe to warm up to room temperature before use.

Remove Prefilled Syringe from carton

Remove the EBGLYSS Prefilled Syringe from the carton by holding the middle of the Syringe Body



and then place the EBGLYSS Prefilled Syringe on a clean flat surface.

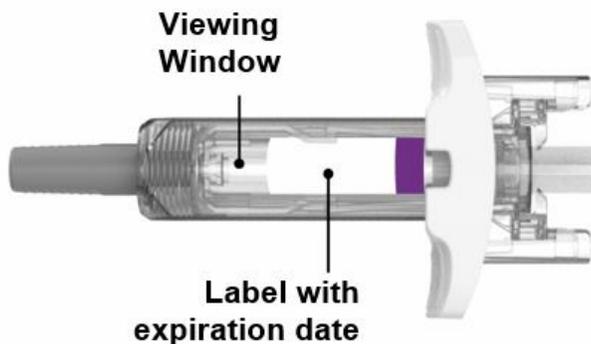
Leave the needle cap on the syringe until you are ready to inject.

Do not pull back on the Plunger Rod at any time.

Inspect the Prefilled Syringe and the medicine

Make sure you have the right medicine. The medicine inside should be clear. It may be colorless to slightly yellow to slightly brown. Some air bubbles are normal.

Note: You may gently rotate the plunger rod to view the syringe label if needed.



Do not use the Prefilled Syringe (see **Disposing of EBGLYSS**) if the:

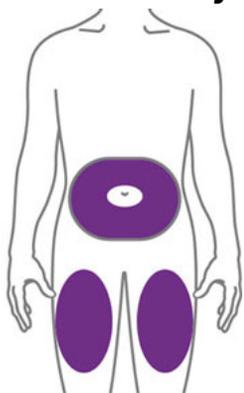
- Needle Cap is missing or not securely attached
- Prefilled Syringe looks damaged
- medicine is frozen
- medicine is cloudy, is discolored, or has particles
- expiration date printed on the label has passed.

Wash your hands with soap and water

Choose and clean your injection site

Your healthcare provider can help you choose the injection site that is best for you.

Clean the injection site with an alcohol wipe and let dry.



You or another person may inject into these areas.

- **Stomach area (abdomen) -**

- At least 2 inches away from the belly button (navel).

- **Front of thigh -**

- At least 2 inches above the knee and 2 inches below the groin.

- **Back of upper arm -**

- Another person should inject into the back of your upper arm.

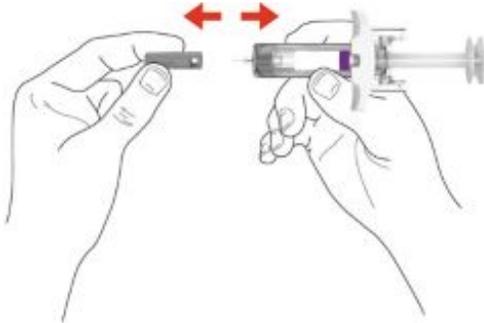
Another person should inject into this area.

Do not inject in the exact same spot every time.

Do not inject into areas where the skin is tender, bruised, red, hard, or in an area of the skin that is affected by eczema (atopic dermatitis) or other skin lesions.

Injecting EBGLYSS

1 Remove Needle Cap

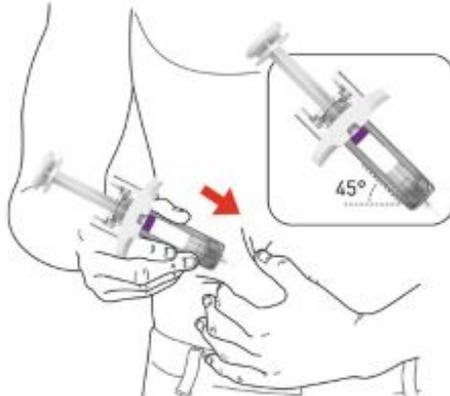


Hold the EBGLYSS Prefilled Syringe in the middle of the Syringe Body and pull the Needle Cap straight off.

Do not put the Needle Cap back on; this could damage the needle.

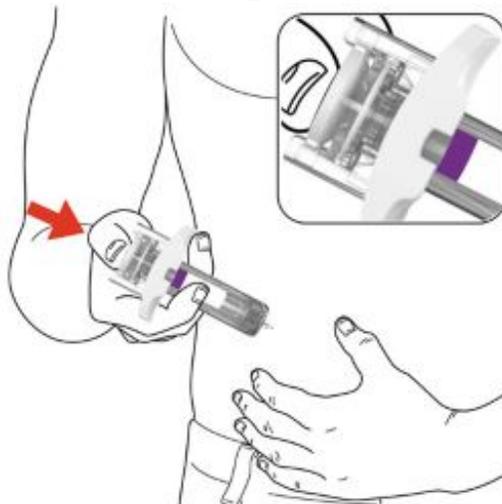
Do not touch the Needle.

2 Pinch the injection site and insert the Needle



Gently pinch a fold of skin at the injection site with one hand. Insert the Needle completely into the fold of skin at about a 45° angle with your other hand.

3 Push in the Plunger Rod



Gently relax the pinch while keeping the needle in place. Push the Plunger Rod all the way down as far as it will go to inject all of the medicine.

Note: It is normal to feel some resistance.

You have to give the full dose to activate the Needle Shield.

4 Release the Plunger Rod



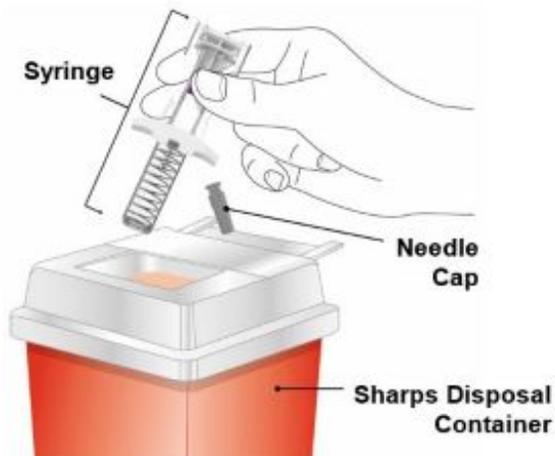
Lift your thumb to release the Plunger Rod until the Needle is covered by the Needle Shield.

Lightly press a cotton ball or gauze on the injection site if you see any blood.

Do not rub your skin after the injection.

Disposing of EBGLYSS

Dispose of (throw away) the used Prefilled Syringe



Put the used EBGLYSS Prefilled Syringe and Needle Cap in an FDA-cleared sharps disposal container right away after use.

Do not throw away (dispose of) the EBGLYSS Prefilled Syringe in your household trash.

If you do not have an FDA-cleared sharps disposal container, you may use a household container that is:

- made of a heavy-duty plastic,
- able to be closed with a tight-fitting, puncture-resistant lid, without sharps being able to come out,
- upright and stable during use,
- leak-resistant, and
- properly labeled to warn of hazardous waste inside the container.

When your sharps disposal container is almost full, you will need to follow your community guidelines for the right way to dispose of your sharps disposal container.

There may be local laws about how you should throw away needles and syringes.

For more information about safe sharps disposal, and for specific information about sharps disposal in the state you live in, go to the FDA website at:

<http://www.fda.gov/safesharpsdisposal>

Do not recycle your used sharps disposal container.

Storing EBGLYSS

- Store your Prefilled Syringe in a refrigerator between 36°F to 46°F (2°C to 8°C).
- EBGLYSS can be stored at room temperature up to 7 days in the original carton. **Do not** store above 86°F (30°C). Throw away (dispose of) EBGLYSS that has been left at room temperature for longer than 7 days.
- Store your Prefilled Syringe in the original carton to protect from light until use.
- **Do not** freeze your Prefilled Syringe. **Do not** shake your Prefilled Syringe.
- **Do not** microwave your Prefilled Syringe, or run hot water over it, or leave it in direct sunlight.
- Throw away (dispose of) your Prefilled Syringe if any of the above conditions are not followed.

Keep your Prefilled Syringes and all medicines out of the reach of children.

If you have more questions about how to use the EBGLYSS Prefilled Syringe:

- Call your healthcare provider
- Call 1-800-Lilly-Rx (1-800-545-5979)
- Visit www.ebglyss.com



**Scan this code to
launch
www.ebglyss.com**

Read the Patient Information insert for EBGLYSS inside this box to learn more about your medicine.

Eli Lilly and Company

Indianapolis, IN 46285, USA

US License Number 1891

EBGLYSS is a trademark of Eli Lilly and Company.

Copyright © 2024, 2025, Eli Lilly and Company. All rights reserved.

This Instructions for Use has been approved by the U.S. Food and Drug Administration.

Revised: October 2025

EBG-0002-PFS-IFU-20251028

PACKAGE LABEL - Ebglyss 250 mg Prefilled Pen

Ebglyss®

(lebrikizumab-lbkz)
injection

250 mg / 2 mL

1 Single-Dose Prefilled Pen

NDC 0002-7772-11

1 x 250 mg/2 mL
Single-Dose Prefilled Pen
For Subcutaneous Use Only
Rx only
Lilly



PACKAGE LABEL - Ebglyss 250 mg Prefilled Syringe

Ebglyss[®]
(lebrikizumab-lbkz)
injection

250 mg / 2 mL

1 Single-Dose Prefilled Syringe with Needle Shield

NDC 0002-7797-11

1 x 250 mg/2 mL

(125 mg/mL)
 Single-Dose Prefilled Syringe with Needle Shield
 For Subcutaneous Use Only
 Rx only
 Lilly



EBGLYSS			
lebrikizumab-lbkz injection, solution			
Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0002-7772
Route of Administration	SUBCUTANEOUS		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	Lebrikizumab (UNII: U9JLP7V031) (Lebrikizumab - UNII:U9JLP7V031)	Lebrikizumab	250 mg in 2 mL

Inactive Ingredients

Ingredient Name	Strength
Acetic acid (UNII: Q40Q9N063P)	1.8 mg in 2 mL
Histidine (UNII: 4QD397987E)	6.2 mg in 2 mL
Polysorbate 20 (UNII: 7T1F30V5YH)	0.6 mg in 2 mL
Sucrose (UNII: C151H8M554)	119.6 mg in 2 mL
Water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0002-7772-11	1 in 1 CARTON	09/13/2024	
1	NDC:0002-7772-01	2 mL in 1 SYRINGE; Type 3: Prefilled Biologic Delivery Device/System (syringe, patch, etc.)		
2	NDC:0002-7772-62	2 in 1 CARTON	09/13/2024	
2		2 mL in 1 SYRINGE; Type 3: Prefilled Biologic Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761306	09/13/2024	

EBGLYSS

lebrikizumab-lbkz injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0002-7797
Route of Administration	SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Lebrikizumab (UNII: U9JLP7V031) (Lebrikizumab - UNII:U9JLP7V031)	Lebrikizumab	250 mg in 2 mL

Inactive Ingredients

Ingredient Name	Strength
Acetic acid (UNII: Q40Q9N063P)	1.8 mg in 2 mL
Histidine (UNII: 4QD397987E)	6.2 mg in 2 mL
Polysorbate 20 (UNII: 7T1F30V5YH)	0.6 mg in 2 mL

Sucrose (UNII: C151H8M554)

119.6 mg in 2 mL

Water (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0002-7797-11	1 in 1 CARTON	09/13/2024	
1	NDC:0002-7797-01	2 mL in 1 SYRINGE; Type 3: Prefilled Biologic Delivery Device/System (syringe, patch, etc.)		
2	NDC:0002-7797-61	1 in 1 CARTON	09/13/2024	
2		2 mL in 1 SYRINGE; Type 3: Prefilled Biologic Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761306	09/13/2024	

Labeler - Eli Lilly and Company (006421325)

Registrant - Eli Lilly and Company (006421325)

Revised: 11/2025

Eli Lilly and Company