

EPINEPHRINE- epinephrine injection

REMEDYREPACK INC.

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use EPINEPHRINE INJECTION, 0.3 mg and EPINEPHRINE INJECTION, 0.15 mg safely and effectively. See full prescribing information for EPINEPHRINE INJECTION, 0.3 mg and EPINEPHRINE INJECTION, 0.15 mg.

EPINEPHRINE injection, 0.3 mg, for intramuscular or subcutaneous use

EPINEPHRINE injection, 0.15 mg, for intramuscular or subcutaneous use

Initial U.S. Approval: 1939

INDICATIONS AND USAGE

Epinephrine Injection, 0.3 mg and Epinephrine Injection, 0.15 mg contain epinephrine, are non-selective alpha and beta-adrenergic receptor agonist indicated in the emergency treatment of allergic reactions (Type I) including anaphylaxis. (1)

DOSAGE AND ADMINISTRATION

- Patients greater than or equal to 30 kg (66 lbs): Epinephrine injection, 0.3 mg (2)
- Patients 15 to 30 kg (33 lbs to 66 lbs): Epinephrine injection, 0.15 mg (2)

Inject intramuscularly or subcutaneously into the anterolateral aspect of the thigh, through clothing if necessary. Each device is a single-dose injection. (2)

DOSAGE FORMS AND STRENGTHS

- Injection: 0.15 mg (0.15 mg/0.3 mL) single-dose pre-filled auto-injector (3)

CONTRAINDICATIONS

None (4)

WARNINGS AND PRECAUTIONS

- In conjunction with use, seek immediate medical or hospital care. (5.1)
- Do not inject intravenously, into buttock, or into digits, hands, or feet. (5.2)
- To minimize the risk of injection related injury, hold the child's leg firmly in place and limit movement prior to and during injection when administering to young children. (5.2)
- Rare cases of serious skin and soft tissue infections have been reported following epinephrine injection. Advise patients to seek medical care if they develop signs or symptoms of infection. (5.3)
- The presence of a sulfite in this product should not deter use. (5.4)
- Administer with caution in patients with heart disease; may aggravate angina pectoris or produce ventricular arrhythmias. (5.5)

ADVERSE REACTIONS

Adverse reactions to epinephrine include anxiety, apprehensiveness, restlessness, tremor, weakness, dizziness, sweating, palpitations, pallor, nausea and vomiting, headache, and/or respiratory difficulties. (6)

To report SUSPECTED ADVERSE REACTIONS, contact Teva Pharmaceuticals USA, Inc. at 1-888-838-2872 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Cardiac glycosides or diuretics: observe for development of cardiac arrhythmias. (7)
- Tricyclic antidepressants, monoamine oxidase inhibitors, levothyroxine sodium, and certain antihistamines potentiate effects of epinephrine. (7)
- Beta-adrenergic blocking drugs antagonize cardiostimulating and bronchodilating effects of epinephrine. (7)
- Alpha-adrenergic blocking drugs antagonize vasoconstricting and hypertensive effects of epinephrine. (7)
- Ergot alkaloids may reverse the pressor effects of epinephrine. (7)

USE IN SPECIFIC POPULATIONS

- Elderly patients may be at greater risk of developing adverse reactions. (5.5, 8.5)

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

FULL PRESCRIBING INFORMATION: CONTENTS*

1 INDICATIONS AND USAGE

2 DOSAGE AND ADMINISTRATION

- 2.1 Recommended Dosage According to Patient Body Weight
- 2.2 Administration Instructions

3 DOSAGE FORMS AND STRENGTHS

4 CONTRAINDICATIONS

5 WARNINGS AND PRECAUTIONS

- 5.1 Emergency Treatment
- 5.2 Injection-Related Complications
- 5.3 Serious Infections at the Injection Site
- 5.4 Allergic Reactions Associated with Sulfite
- 5.5 Disease Interactions

6 ADVERSE REACTIONS

7 DRUG INTERACTIONS

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

13 NONCLINICAL TOXICOLOGY

- 13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

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

Epinephrine Injection, 0.3 mg and Epinephrine Injection, 0.15 mg are indicated in the emergency treatment of allergic reactions (Type I) including anaphylaxis to stinging insects (e.g., order Hymenoptera, which include bees, wasps, hornets, yellow jackets and fire ants) and biting insects (e.g., triatoma, mosquitoes), allergen immunotherapy, foods, drugs, diagnostic testing substances (e.g., radiocontrast media) and other

allergens, as well as idiopathic anaphylaxis or exercise-induced anaphylaxis.

Epinephrine Injection, 0.3 mg and Epinephrine Injection, 0.15 mg are intended for immediate administration in patients who are determined to be at increased risk for anaphylaxis, including individuals with a history of anaphylactic reactions.

Anaphylactic reactions may occur within minutes after exposure and consist of flushing, apprehension, syncope, tachycardia, thready or unobtainable pulse associated with a fall in blood pressure, convulsions, vomiting, diarrhea and abdominal cramps, involuntary voiding, wheezing, dyspnea due to laryngeal spasm, pruritus, rashes, urticaria or angioedema.

Epinephrine Injection, 0.3 mg and Epinephrine Injection, 0.15 mg are intended for immediate administration as emergency supportive therapy only and are not a substitute for immediate medical care.

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosage According to Patient Body Weight

- Patients greater than or equal to 30 kg (approximately 66 pounds or more):
Epinephrine injection, 0.3 mg
- Patients 15 kg to 30 kg (33 pounds to 66 pounds): Epinephrine injection, 0.15 mg

2.2 Administration Instructions

- Inject the single-dose epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg intramuscularly or subcutaneously into the anterolateral aspect of the thigh, through clothing if necessary. Do not inject intravenously, and do not inject into buttocks, into digits, hands or feet [*see Warnings and Precautions (5.2)*].
- Instruct caregivers of young children who are prescribed an epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg and who may be uncooperative and kick or move during an injection to hold the leg firmly in place and limit movement prior to and during an injection [*see Warnings and Precautions (5.2)*].
- Each epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg is a single-dose epinephrine injection for single use. Since the doses of epinephrine delivered from epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg are fixed, consider using other forms of injectable epinephrine if doses lower than 0.15 mg are deemed necessary.
- With severe persistent anaphylaxis, repeat injections with an additional epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg may be necessary. More than two sequential doses of epinephrine should only be administered under direct medical supervision [*see Warnings and Precautions (5.1)*].
- The epinephrine solution in the clear window of the Epinephrine Injection 0.3 mg (Auto-Injector) or Epinephrine Injection 0.15 mg (Auto-Injector) should be inspected visually for particulate matter and discoloration.

Discarding After Use:

The epinephrine injection 0.3 mg and epinephrine injection 0.15 mg each contain 1 mL epinephrine solution. Approximately 0.7 mL remains in the auto-injector after activation, but is not available for future use, and should be discarded.

3 DOSAGE FORMS AND STRENGTHS

- Injection: 0.15 mg (0.15 mg/0.3 mL), clear and colorless solution in single-dose pre-filled auto-injector

4 CONTRAINDICATIONS

None

5 WARNINGS AND PRECAUTIONS

5.1 Emergency Treatment

Epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg are intended for immediate administration as emergency supportive therapy and are not intended as a substitute for immediate medical care. **In conjunction with the administration of epinephrine, the patient should seek immediate medical or hospital care.** More than two sequential doses of epinephrine should only be administered under direct medical supervision [*see Indications and Usage (1), Dosage and Administration (2) and Patient Counseling Information (17)*].

5.2 Injection-Related Complications

Epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg should **only** be injected into the anterolateral aspect of the thigh [*see Dosage and Administration (2) and Patient Counseling Information (17)*].

Do not inject intravenously

Large doses or accidental intravenous injection of epinephrine may result in cerebral hemorrhage due to sharp rise in blood pressure. Rapidly acting vasodilators can counteract the marked pressor effects of epinephrine if there is such inadvertent administration.

Do not inject into buttock

Injection into the buttock may not provide effective treatment of anaphylaxis. Advise the patient to go immediately to the nearest emergency room for further treatment of anaphylaxis. Additionally, injection into the buttock has been associated with Clostridial infections (gas gangrene). Cleansing with alcohol does not kill bacterial spores, and therefore, does not lower this risk.

Do not inject into digits, hands or feet

Since epinephrine is a strong vasoconstrictor, accidental injection into the digits, hands or feet may result in loss of blood flow to the affected area. Advise the patient to go immediately to the nearest emergency room and to inform the healthcare provider in the emergency room of the location of the accidental injection. Treatment of such inadvertent administration should consist of vasodilation, in addition to further appropriate treatment of anaphylaxis [*see Adverse Reactions (6)*].

Hold leg firmly during injection

Lacerations, bent needles, and embedded needles have been reported when epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg have been injected into the thigh of young children who are uncooperative and kick or move during an injection. To minimize the risk of injection related injury when administering, hold the child's leg firmly in place and limit movement prior to and during injection.

5.3 Serious Infections at the Injection Site

Rare cases of serious skin and soft tissue infections, including necrotizing fasciitis and myonecrosis caused by *Clostridia* (gas gangrene), have been reported at the injection site following epinephrine injection for anaphylaxis. *Clostridium* spores can be present on the skin and introduced into the deep tissue with subcutaneous or intramuscular injection. While cleansing with alcohol may reduce presence of bacteria on the skin, alcohol cleansing does not kill *Clostridium* spores. To decrease the risk of *Clostridium* infection, do not inject epinephrine injection into the buttock [see *Warnings and Precautions (5.2)*]. Advise patients to seek medical care if they develop signs or symptoms of infection, such as persistent redness, warmth, swelling, or tenderness, at the epinephrine injection site.

5.4 Allergic Reactions Associated with Sulfite

The presence of a sulfite in this product should not deter administration of the drug for treatment of serious allergic or other emergency situations even if the patient is sulfite-sensitive.

Epinephrine is the preferred treatment for serious allergic reactions or other emergency situations even though this product contains sodium metabisulfite, a sulfite that may, in other products, cause allergic-type reactions including anaphylactic symptoms or life-threatening or less severe asthmatic episodes in certain susceptible persons.

The alternatives to using epinephrine in a life-threatening situation may not be satisfactory.

5.5 Disease Interactions

Some patients may be at greater risk for developing adverse reactions after epinephrine administration. Despite these concerns, it should be recognized that the presence of these conditions is not a contraindication to epinephrine administration in an acute, life-threatening situation. Therefore, patients with these conditions, and/or any other person who might be in a position to administer epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg to a patient experiencing anaphylaxis should be carefully instructed in regard to the circumstances under which epinephrine should be used.

Patients with Heart Disease

Epinephrine should be administered with caution to patients who have heart disease, including patients with cardiac arrhythmias, coronary artery or organic heart disease, or hypertension . In such patients, or in patients who are on drugs that may sensitize the heart to arrhythmias , epinephrine may precipitate or aggravate angina pectoris as well as produce ventricular arrhythmias [see *Drug Interactions (7)* and *Adverse Reactions (6)*].

Other Patients and Diseases

Epinephrine should be administered with caution to patients with hyperthyroidism, diabetes, elderly individuals, and pregnant women. Patients with Parkinson's disease may notice a temporary worsening of symptoms.

6 ADVERSE REACTIONS

Due to the lack of randomized, controlled clinical trials of epinephrine for the treatment of anaphylaxis, the true incidence of adverse reactions associated with the systemic use of epinephrine is difficult to determine. Adverse reactions reported in observational trials, case reports, and studies are listed below.

Common adverse reactions to systemically administered epinephrine include anxiety; apprehensiveness; restlessness; tremor; weakness; dizziness; sweating; palpitations; pallor; nausea and vomiting; headache; and/or respiratory difficulties. These symptoms occur in some persons receiving therapeutic doses of epinephrine, but are more likely to occur in patients with hypertension or hyperthyroidism [*see Warnings and Precautions (5.5)*].

Cardiovascular Reactions

- Arrhythmias, including fatal ventricular fibrillation, have been reported, particularly in patients with underlying cardiac disease or those receiving certain drugs [*see Warnings and Precautions (5.5)* and *Drug Interactions (7)*].
- Rapid rises in blood pressure have produced cerebral hemorrhage, particularly in elderly patients with cardiovascular disease [*see Warnings and Precautions (5.5)*].
- Angina may occur in patients with coronary artery disease [*see Warnings and Precautions (5.5)*].
- Rare cases of stress cardiomyopathy have been reported in patients treated with epinephrine.

Reactions from Accidental Injection and/or Improper Technique

- Accidental injection into the digits, hands or feet may result in loss of blood flow to the affected area [*see Warnings and Precautions (5.2)*].
- Adverse reactions experienced as a result of accidental injections may include increased heart rate, local reactions including injection site pallor, coldness and hypoaesthesia or injury at the injection site resulting in bruising, bleeding, discoloration, erythema or skeletal injury.
- Lacerations, bent needles, and embedded needles have been reported when epinephrine injection has been injected into the thigh of young children who are uncooperative and kick or move during the injection [*see Warnings and Precautions (5.2)*].
- Injection into the buttock has resulted in cases of gas gangrene [*see Warnings and Precautions (5.2)*].

Skin and Soft Tissue Infections

- Rare cases of serious skin and soft tissue infections, including necrotizing fasciitis and myonecrosis caused by Clostridia (gas gangrene), have been reported following epinephrine injection, including epinephrine injection 0.3 mg, in the thigh [*see Warnings and Precautions (5.3)*].

7 DRUG INTERACTIONS

Cardiac Glycosides, Diuretics, and Anti-arrhythmics

Patients who receive epinephrine while concomitantly taking cardiac glycosides, diuretics, or anti-arrhythmics should be observed carefully for the development of cardiac arrhythmias [*see Warnings and Precautions (5.5)*].

Antidepressants, Monoamine Oxidase Inhibitors, Levothyroxine, and Antihistamines

The effects of epinephrine may be potentiated by tricyclic antidepressants, monoamine oxidase inhibitors, levothyroxine sodium, and certain antihistamines, notably chlorpheniramine, tripeleminamine, and diphenhydramine.

Beta-Adrenergic Blockers

The cardiostimulating and bronchodilating effects of epinephrine are antagonized by beta-adrenergic blocking drugs, such as propranolol.

Alpha-Adrenergic Blockers

The vasoconstricting and hypertensive effects of epinephrine are antagonized by alpha-adrenergic blocking drugs, such as phentolamine.

Ergot Alkaloids

Ergot alkaloids may also reverse the pressor effects of epinephrine.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

There are no adequate and well controlled studies of the acute effect of epinephrine in pregnant women. In animal reproductive studies, epinephrine administered by the subcutaneous route to rabbits, mice, and hamsters during the period of organogenesis was teratogenic at doses 7 times and higher than the maximum recommended human intramuscular and subcutaneous dose on a mg/m² basis. Epinephrine is the first-line medication of choice for the treatment of anaphylaxis during pregnancy in humans. Epinephrine should be used for treatment of anaphylaxis during pregnancy in the same manner as it is used in non-pregnant patients.

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.

Clinical Considerations

Disease-associated maternal and embryo/fetal risk:

During pregnancy, anaphylaxis can be catastrophic and can lead to hypoxic-ischemic encephalopathy and permanent central nervous system damage or death in the mother and, more commonly, in the fetus or neonate. The prevalence of anaphylaxis occurring during pregnancy is reported to be approximately 3 cases per 100,000 deliveries.

Management of anaphylaxis during pregnancy is similar to management in the general population. Epinephrine is the first line-medication of choice for treatment of anaphylaxis; it should be used in the same manner in pregnant and non-pregnant patients. In conjunction with the administration of epinephrine, the patient should seek immediate medical or hospital care.

Data

Animal Data:

In an embryofetal development study with rabbits dosed during the period of organogenesis, epinephrine was shown to be teratogenic (including gastroschisis and embryonic lethality) at doses approximately 40 times the maximum recommended intramuscular or subcutaneous dose (on a mg/m² basis at a maternal subcutaneous dose of 1.2 mg/kg/day for two to three days).

In an embryofetal development study with mice dosed during the period of organogenesis, epinephrine was shown to be teratogenic (including embryonic lethality) at doses approximately 8 times the maximum recommended intramuscular or subcutaneous dose (on a mg/m² basis at maternal subcutaneous dose of 1 mg/kg/day for 10 days). These effects were not seen in mice at approximately 4 times the maximum recommended daily intramuscular or subcutaneous dose (on a mg/m² basis at a subcutaneous maternal dose of 0.5 mg/kg/day for 10 days).

In an embryofetal development study with hamsters dosed during the period of organogenesis from gestation days 7 to 10, epinephrine was shown to be teratogenic at doses approximately 7 times the maximum recommended intramuscular or subcutaneous dose (on a mg/m² basis at a maternal subcutaneous dose of 0.5 mg/kg/day).

8.2 Lactation

Risk Summary

There is no information on the presence of epinephrine in human milk, the effects on breastfed infants, or the effects on milk production. Epinephrine is the first line-medication of choice for treatment of anaphylaxis; it should be used in the same manner in breastfeeding and non-breastfeeding patients.

8.4 Pediatric Use

Epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg may be administered to pediatric patients at a dosage appropriate to body weight [*see Dosage and Administration (2.1)*]. Clinical experience with the use of epinephrine suggests that the adverse reactions seen in children are similar in nature and extent to those both expected and reported in adults. Since the doses of epinephrine delivered from epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg are fixed, consider using other forms of injectable epinephrine if doses lower than 0.15 mg are deemed necessary.

8.5 Geriatric Use

Clinical studies for the treatment of anaphylaxis have not been performed in subjects aged 65 and over to determine whether they respond differently from younger subjects.

However, other reported clinical experience with use of epinephrine for the treatment of anaphylaxis has identified that geriatric patients may be particularly sensitive to the effects of epinephrine. Therefore, epinephrine injection, 0.3 mg should be administered with caution in elderly individuals, who may be at greater risk for developing adverse reactions after epinephrine administration [see *Warnings and Precautions* (5.5), *Overdosage* (10)].

10 OVERDOSAGE

Overdosage of epinephrine may produce extremely elevated arterial pressure, which may result in cerebrovascular hemorrhage, particularly in elderly patients. Overdosage may also result in pulmonary edema because of peripheral vascular constriction together with cardiac stimulation. Treatment consists of rapidly acting vasodilators or alpha-adrenergic blocking drugs and/or respiratory support.

Epinephrine overdosage can also cause transient bradycardia followed by tachycardia, and these may be accompanied by potentially fatal cardiac arrhythmias. Premature ventricular contractions may appear within one minute after injection and may be followed by multifocal ventricular tachycardia (prefibrillation rhythm). Subsidence of the ventricular effects may be followed by atrial tachycardia and occasionally by atrioventricular block. Treatment of arrhythmias consists of administration of a beta-adrenergic blocking drug such as propranolol.

Overdosage sometimes results in extreme pallor and coldness of the skin, metabolic acidosis, and kidney failure. Suitable corrective measures must be taken in such situations.

11 DESCRIPTION

Epinephrine Injection USP, 0.3 mg and Epinephrine Injection USP, 0.15 mg are single-dose auto-injectors and combination products containing drug and device components.

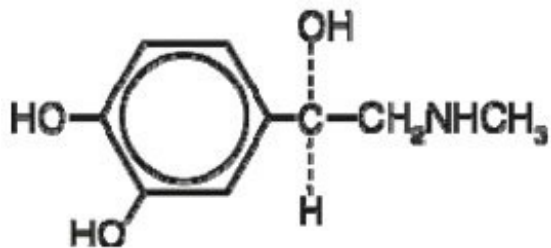
Each Epinephrine Injection USP, 0.3 mg (Auto-Injector) delivers a single dose of 0.3 mg epinephrine, USP from epinephrine injection USP, 0.3 mg/0.3 mL in a sterile solution.

Each Epinephrine Injection USP, 0.15 mg (Auto-Injector) delivers a single dose of 0.15 mg epinephrine, USP from epinephrine injection USP, 0.15 mg/0.3 mL in a sterile solution.

Each 0.3 mL in the Epinephrine Injection USP, 0.3 mg (Auto-Injector) contains 0.3 mg epinephrine USP, 1.8 mg sodium chloride, 0.4 mg sodium metabisulfite, 0.4 mg sodium tartrate (dihydrate), hydrochloric acid to adjust pH, and water for injection. The pH range is 2.2 to 5.0.

Each 0.3 mL in the Epinephrine Injection USP, 0.15 mg (Auto-Injector) contains 0.15 mg epinephrine USP, 1.8 mg sodium chloride, 0.4 mg sodium metabisulfite, 0.2 mg sodium tartrate (dihydrate), hydrochloric acid to adjust pH, and water for injection. The pH range is 2.2 to 5.0.

Epinephrine, USP is a sympathomimetic catecholamine. Chemically, epinephrine, USP is (-)-3,4-Dihydroxy- α -[(methylamino)methyl]benzyl alcohol with the following structure:



Epinephrine solution deteriorates rapidly on exposure to air or light, turning pink from oxidation to adrenochrome and brown from the formation of melanin. Replace Epinephrine Injection USP, 0.3 mg and Epinephrine Injection USP, 0.15 mg if the epinephrine solution appears discolored (pinkish or darker than slightly yellow), cloudy, or if it contains a precipitate.

Thoroughly review the patient instructions and operation of Epinephrine Injection USP, 0.3 mg or Epinephrine Injection USP, 0.15 mg with patients and caregivers prior to use [see *Patient Counseling Information (17)*].

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Epinephrine acts on both alpha- and beta-adrenergic receptors.

12.2 Pharmacodynamics

Through its action on alpha-adrenergic receptors, epinephrine lessens the vasodilation and increased vascular permeability that occurs during anaphylaxis, which can lead to loss of intravascular fluid volume and hypotension.

Through its action on beta-adrenergic receptors, epinephrine causes bronchial smooth muscle relaxation and helps alleviate bronchospasm, wheezing and dyspnea that may occur during anaphylaxis.

Epinephrine also alleviates pruritus, urticaria, and angioedema and may relieve gastrointestinal and genitourinary symptoms associated with anaphylaxis because of its relaxer effects on the smooth muscle of the stomach, intestine, uterus and urinary bladder.

When given subcutaneously or intramuscularly, epinephrine has a rapid onset and short duration of action.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Long-term studies to evaluate the carcinogenic potential of epinephrine have not been conducted.

Epinephrine and other catecholamines have been shown to have mutagenic potential *in vitro*. Epinephrine was positive in the *Salmonella* bacterial reverse mutation assay, positive in the mouse lymphoma assay, and negative in the *in vivo* micronucleus assay.

Epinephrine is an oxidative mutagen based on the *E. coli* WP2 Mutoxitest bacterial reverse mutation assay. This should not prevent the use of epinephrine where indicated [see *Indications and Usage (1)*].

The potential for epinephrine to impair reproductive performance has not been evaluated, but epinephrine has been shown to decrease implantation in female rabbits dosed subcutaneously with 1.2 mg/kg/day (40-fold the highest human intramuscular or subcutaneous daily dose) during gestation days 3 to 9.

16 HOW SUPPLIED/STORAGE AND HANDLING

Epinephrine Injection USP, 0.3 mg 2-Pack and Epinephrine Injection USP, 0.15 mg 2-Pack also includes a W-clip to clip two auto-injectors together.

NDC: 70518-3173-00

PACKAGING: 2 in 1 CARTON, 1 in 1 CONTAINER, 0.3 mL in 1 SYRINGE TYPE 2

Storage and Handling

Protect from light. Epinephrine, USP is light sensitive and the auto-injector is manufactured from transparent UV stabilized polycarbonate to protect it from light.

Store at 20° to 25°C (68° to 77°F); excursions permitted to 15° to 30°C (59° to 86°F) [See USP Controlled Room Temperature].

Do not refrigerate.

Before using, check to make sure the solution in the auto-injector is clear and colorless.

Replace the auto-injector if the solution is discolored (pinkish or darker than slightly yellow), cloudy, or contains a precipitate.

Properly dispose all used, unwanted or expired epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg auto-injectors.

KEEP THIS AND ALL MEDICATIONS OUT OF THE REACH OF CHILDREN.

Repackaged and Distributed By:

Remedy Repack, Inc.

625 Kolter Dr. Suite #4 Indiana, PA 1-724-465-8762

17 PATIENT COUNSELING INFORMATION

See FDA-Approved Patient Labeling (Patient Information and Instructions for Use)

A healthcare provider should review the patient instructions and operation of epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg in detail, with the patient or caregiver.

Epinephrine is essential for the treatment of anaphylaxis. Patients who are at risk of or with a history of severe allergic reactions (anaphylaxis) to insect stings or bites, foods, drugs, and other allergens, as well as idiopathic and exercise-induced anaphylaxis, should be carefully instructed about the circumstances under which epinephrine should be used.

Administration

Instruct patients and/or caregivers in the appropriate use of epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg should be injected into the middle of the outer thigh (through clothing, if necessary). Each device is a single-use injection. Advise patients to seek immediate medical care in conjunction with administration of epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg.

Instruct caregivers to hold the leg of young children firmly in place and limit movement prior to and during injection. Lacerations, bent needles, and embedded needles have been reported when epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg have been injected into the thigh of young children who are uncooperative and kick or move during an injection [*see Warnings and Precautions (5.2)*].

Instruct patients and/or caregivers to throw away the blue safety release immediately after using epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg. This small part may pose a choking hazard for children.

Complete patient information, including dosage, directions for proper administration and precautions can be found inside each epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg carton. A printed label on the surface of epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg shows instructions for use and a diagram depicting the injection process.

Training

Instruct patients and/or caregivers to use and practice with the Trainer to familiarize themselves with the use of epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg in an allergic emergency. The Trainer may be used multiple times. A Trainer device is provided in 2-Pack cartons.

Instruct patients and/or caregivers to immediately place the blue safety release back on the Trainer and reset it after practicing. This small part may pose a choking hazard for children.

Adverse Reactions

Epinephrine may produce symptoms and signs that include an increase in heart rate, the sensation of a more forceful heartbeat, palpitations, sweating, nausea and vomiting, difficulty breathing, pallor, dizziness, weakness or shakiness, headache, apprehension, nervousness, or anxiety. These signs and symptoms usually subside rapidly, especially with rest, quiet and recumbency. Patients with hypertension or hyperthyroidism may develop more severe or persistent effects, and patients with coronary artery disease could experience angina. Patients with diabetes may develop increased blood glucose levels following epinephrine administration. Patients with Parkinson's disease may notice a temporary worsening of symptoms [*see Warnings and Precautions (5.5)*].

Accidental Injection

Advise patients to seek immediate medical care in the case of accidental injection. Since epinephrine is a strong vasoconstrictor when injected into the digits, hands, or feet, treatment should be directed at vasodilatation if there is such an accidental injection to these areas [*see Warnings and Precautions (5.2)*].

Serious Infections at the Injection Site

Rare cases of serious skin and soft tissue infections, including necrotizing fasciitis and myonecrosis caused by Clostridia (gas gangrene), have been reported at the injection site following epinephrine injection for anaphylaxis. Advise patients to seek medical care if they develop signs or symptoms of infection, such as persistent redness, warmth, swelling, or tenderness, at the epinephrine injection site [*see Warnings and Precautions (5.3)*].

Storage and Handling

Instruct patients to inspect the epinephrine solution visually through the clear window of the auto-injector periodically. Replace epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg if the epinephrine solution appears discolored (pinkish or darker than slightly yellow), cloudy or if it contains a precipitate. Epinephrine is light sensitive and should be protected from light. The auto-injector is not waterproof. Instruct patients that epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg must be used or properly disposed once the blue safety release is removed or after use [*see Storage and Handling (16.2)*].

Advise patients and caregivers to give used epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg auto-injectors to their healthcare provider for inspection and proper disposal.

Advise patients and caregivers to promptly dispose of medicines that are no longer needed. Dispose of expired, unwanted, or unused epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg auto-injectors in an FDA-cleared sharps container. Instruct patients not to dispose epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg in their household trash. Instruct patients that if they do not have a FDA-cleared sharps disposal container, they may use a household container that is made of a heavy-duty plastic, can be closed with a tight-fitting and 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. Inform patients that they can visit the FDA website for additional information on disposal of unused medicines.

Complete patient information, including dosage, directions for proper administration and precautions can be found inside each epinephrine injection, 0.3 mg (Auto-Injector) and epinephrine injection, 0.15 mg (Auto-Injector) carton.

Repackaged By / Distributed By: RemedyRepack Inc.

625 Kolter Drive, Indiana, PA 15701

(724) 465-8762

PATIENT INFORMATION and INSTRUCTIONS FOR USE

Epinephrine Injection USP, 0.3 mg (Auto-Injector)

Epinephrine Injection USP, 0.3 mg = one dose of 0.3 mg epinephrine USP, 0.3 mg/0.3 mL

Epinephrine Injection USP, 0.15 mg (Auto-Injector)

Epinephrine Injection USP, 0.15 mg = one dose of 0.15 mg epinephrine USP, 0.15 mg/0.3 mL

For allergic emergencies (anaphylaxis)

Patient Information

Read this Patient Information Leaflet carefully before using the epinephrine injection, 0.3 mg (auto-injector) or epinephrine injection, 0.15 mg (auto-injector) and each time you get a refill. There may be new information. You, your parent, caregiver, or others who may be in a position to administer epinephrine injection, 0.3 mg (auto-injector) or epinephrine injection, 0.15 mg (auto-injector), should know how to use it before you have an allergic emergency.

This information does not take the place of talking with your healthcare provider about your medical condition or your treatment.

What is the most important information I should know about epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg?

1. Epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg are single-dose automatic injection devices (auto-injectors) that contain epinephrine, a medicine used to treat allergic emergencies (anaphylaxis). Anaphylaxis can be life threatening, can happen within minutes, and can be caused by stinging and biting insects, allergy injections, foods, medicines, exercise, or unknown causes.

Symptoms of anaphylaxis may include:

trouble breathing

wheezing

hoarseness (changes in the way your voice sounds)

hives (raised reddened rash that may itch)

severe itching

swelling of your face, lips, mouth, or tongue

skin rash, redness, or swelling

fast heartbeat

weak pulse

feeling very anxious

confusion

stomach pain

losing control of urine or bowel movements (incontinence)

diarrhea or stomach cramps

dizziness, fainting, or "passing out" (unconsciousness)

2. Always carry your epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg with you because you may not know when anaphylaxis may happen.

Talk to your healthcare provider if you need additional units to keep at work, school, or other locations. Tell your family members, caregivers, and others where you keep your epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg and how to use it before you need it. You may be unable to speak in an allergic emergency.

3. When you have an allergic emergency (anaphylaxis)

Use epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg right away. Get emergency medical help right away. You may need further medical attention. You may need to use a second epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg if symptoms continue or recur. Only a healthcare provider should give additional doses of epinephrine if you need more than 2 injections for a single anaphylaxis episode.

What are epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg?

Epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg are disposable, pre-filled auto-injectors used to treat life-threatening, allergic emergencies including anaphylaxis in people who are at risk for or have a history of serious allergic emergencies. Each device contains a single dose of epinephrine.

Epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg are for immediate self (or caregiver) administration and do not take the place of emergency medical care. You should get emergency help right away after using epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg.

Epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg are for people who have been prescribed this medicine by their healthcare provider.

The Epinephrine Injection, 0.3 mg (Auto-Injector) is for patients who weigh 66 pounds or more (30 kilograms or more).

The Epinephrine Injection, 0.15 mg (Auto-Injector) is for patients who weigh about 33 to 66 pounds (15 to 30 kilograms).

It is not known if epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg are safe and effective in children who weigh less than 33 pounds (15 kilograms).

What should I tell my healthcare provider before using the epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg?

Before you use epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg, tell your healthcare provider about all your medical conditions, but especially if you:

have heart problems or high blood pressure.

have diabetes.

have thyroid problems.

have asthma.

have a history of depression.

have Parkinson's disease.

are pregnant or plan to become pregnant. It is not known if epinephrine will harm your unborn baby.

are breastfeeding or plan to breastfeed. It is not known if epinephrine 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. Tell your healthcare provider about all of your known allergies.

Especially tell your healthcare provider if you take certain asthma medicines.

Epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg and other medicines may affect each other, causing side effects. Epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg may affect the way other medicines work, and other medicines may

affect how epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg work.

Know the medicines you take. Keep a list of them to show your healthcare provider and pharmacist when you get a new medicine.

Use your epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg for treatment of anaphylaxis as prescribed by your healthcare provider, regardless of your medical conditions or the medicines you take.

How should I use epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg?

Each epinephrine injection, 0.3 mg (auto-injector) or epinephrine injection, 0.15 mg (auto-injector) contains only 1 dose of medicine (single-dose).

Epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg Auto-Injectors deliver a fixed dose of epinephrine. The auto-injectors cannot be reused. Do not try to reuse epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg after the device has been activated. It is normal for most of the medicine to remain in the auto-injector after the dose has been injected. The dose has been injected if the orange tip is extended and the window is blocked.

Epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg should be injected into the middle of your outer thigh (upper leg). It can be injected through your clothing if needed. Do not inject into a vein or into the buttocks, fingers, toes, hands or feet.

Read the Instructions for Use at the end of this Patient Information Leaflet about the right way to use epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg.

Your healthcare provider will show you how to safely use the epinephrine injection, 0.3 mg (auto-injector) or epinephrine injection, 0.15 mg (auto-injector).

Use your single-dose epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg exactly as your healthcare provider tells you to use it. You may need to use a second epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg if symptoms continue or recur. Only a healthcare provider should give additional doses of epinephrine if you need more than 2 injections for a single anaphylaxis episode.

Caution: Never put your thumb, fingers, or hand over the orange tip. Never press or push the orange tip with your thumb, fingers, or hand. The needle comes out of the orange tip. Accidental injection into finger, hands or feet may cause a loss of blood flow to these areas. If an accidental injection happens, go immediately to the nearest emergency room. Tell the healthcare provider where on your body you received the accidental injection.

Your epinephrine injection, 0.3 mg (auto-injector) and epinephrine injection, 0.15 mg (auto-injector) may come packaged with an Epinephrine Injection Trainer and separate Epinephrine Injection Trainer Instructions for Use. The Epinephrine Injection Trainer has a grey color. The grey Epinephrine Injection Trainer contains no medicine and no needle. Keep the Epinephrine Injection Trainer and the real Epinephrine Injection, 0.3 mg and Epinephrine Injection, 0.15 mg Auto-Injectors away from young children. The real Epinephrine Injection, 0.3 mg and Epinephrine Injection, 0.15 mg Auto-Injectors and Epinephrine Injection Trainer are not toys. For young children, use of the Epinephrine Injection Trainer and the real Epinephrine Injection, 0.3 mg and Epinephrine Injection, 0.15 mg Auto-Injectors should be supervised by an adult. Periodically practice with your Epinephrine Injection Trainer before an allergic emergency happens to make sure you are able to safely use the real epinephrine injection, 0.3 mg (auto-injector) and epinephrine injection, 0.15 mg (auto-injector) in an emergency. Always carry your real epinephrine injection, 0.3 mg (auto-injector) or epinephrine injection 0.15 mg (auto-injector) with you in case of an allergic emergency. Additional training resources are

available at www.TevaEpinephrine.com.

Do not drop the auto-injector. If the auto-injector is dropped, check for damage and leakage. Throw away (dispose of) the auto-injector and replace if damage or leakage is noticed or suspected.

What are the possible side effects of epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg?

Epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg may cause serious side effects.

The epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg should only be injected into the middle of your outer thigh (upper leg). Do not inject the epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg into your:

veins
buttocks
fingers, toes, hands, or feet

If you accidentally inject epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg into any other part of your body, go to the nearest emergency room right away. Tell the healthcare provider where on your body you received the accidental injection.

Rarely, patients who have used epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg may develop infections at the injection site within a few days of an injection. Some of these infections can be serious. Call your healthcare provider right away if you have any of the following at an injection site:

redness that does not go away
swelling
tenderness
the area feels warm to the touch

Cuts on the skin, bent needles, and needles that remain in the skin after the injection, have happened in young children who do not cooperate and kick or move during an injection. If you inject a young child with epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg, hold their leg firmly in place before and during the injection to prevent injuries. Ask your healthcare provider to show you how to:

Hold the young child firmly in place (restrain).

With 1 hand, quickly twist the yellow cap off the Epinephrine Injection, 0.3 mg auto-injector or the green cap off the Epinephrine Injection, 0.15 mg auto-injector in the direction of the "twist arrow" to remove it.

Grasp the auto-injector in your fist with the orange tip (needle end) pointing downward. With your other hand, pull off the blue safety release.

If you have certain medical conditions, or take certain medicines, your condition may get worse or you may have longer lasting side effects when you use your epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg. Talk to your healthcare provider about all your medical conditions.

Common side effects of epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg include:

fast, irregular or "pounding" heartbeat

sweating
headache
weakness
shakiness
paleness
feelings of over excitement, nervousness or anxiety
dizziness
nausea or vomiting
breathing problems

These side effects may go away with rest. Tell your healthcare provider if you have any side effect that bothers you or that does not go away.

These are not all the possible side effects of the epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg. For more information, ask your healthcare provider or pharmacist.

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 epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg?

Store epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg at room temperature between 68° to 77°F (20° to 25°C).

Protect from light.

Do not expose to extreme cold or heat. For example, do not store in your vehicle's glove box and do not store in the refrigerator or freezer.

Examine the contents in the clear window of your auto-injector periodically. The solution should be clear. If the solution is discolored (pinkish or darker than slightly yellow) or contains solid particles, replace the unit.

Always protect your epinephrine injection, 0.3 mg (auto-injector) and epinephrine injection, 0.15 mg (auto-injector) from damage and water.

The blue safety release helps to prevent accidental injection. Keep the blue safety release on until you need to use epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg. Your epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg has an expiration date. Replace it before the expiration date. Throw away (dispose of) expired, unwanted, or unused epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg in an FDA-cleared sharps disposal container. Do not throw away the epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg 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 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 used needles and syringes. For more information about safe sharps disposal, and for specific information about sharps disposal in the state that you live in, go to the FDA's website at:

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

Visit the FDA's website (<https://www.fda.gov/drugs/safe-disposal-medicines/disposal-unused-medicines-what-you-should-know>) for more information about how to throw away (dispose of) unused, unwanted or expired medicines.

Keep epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg and all medicines out of the reach of children.

General information about the safe and effective use of epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information Leaflet. Do not use the epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg for a condition for which it was not prescribed. Do not give your epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg to other people.

This Patient Information Leaflet summarizes the most important information about epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg. If you would like more information, talk to your healthcare provider. You can ask your pharmacist or healthcare provider for information about epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg that is written for health professionals.

For more information and video instructions on the use of epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg go to www.TevaEpinephrine.com or call 1-888-838-2872.

What are the ingredients in epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg?

Active Ingredients: Epinephrine

Inactive Ingredients: sodium chloride, sodium metabisulfite, sodium tartrate (dihydrate), hydrochloric acid, and water

Important Information

The Epinephrine Injection, 0.3 mg (Auto-Injector) has a yellow colored label.

The Epinephrine Injection, 0.15 mg (Auto-Injector) has a green colored label.

The Epinephrine Injection Trainer has a grey color and contains no medicine and no needle.

Your auto-injector is designed to work through clothing.

The blue safety release on the Epinephrine Injection, 0.3 mg (Auto-Injector) and Epinephrine Injection, 0.15 mg (Auto-Injector) helps to prevent accidental injection of the device. Do not remove the blue safety release until you are ready to use it.

Choking hazard: The blue safety release is a small part that may become a choking hazard for children. Throw away the blue safety release immediately after using epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg Auto-Injector.

Only inject into the middle of the outer thigh (upper leg). Never inject into any other part of the body.

Never put your thumb, fingers, or your hand over the orange tip. The needle comes out of the orange tip.

If an accidental injection happens, get emergency medical help right away.

INSTRUCTIONS FOR USE

Epinephrine Injection USP, 0.3 mg (Auto-Injector)

Epinephrine Injection USP, 0.3 mg = one dose of 0.3 mg epinephrine USP, 0.3 mg/0.3 mL

Epinephrine Injection USP, 0.15 mg (Auto-Injector)

Epinephrine Injection USP, 0.15 mg = one dose of 0.15 mg epinephrine USP, 0.15 mg/0.3 mL

For allergic emergencies (anaphylaxis)

Read this Instructions for Use carefully before you use epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg. Before you need to use your single-dose epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg auto-injector, make sure your healthcare provider shows you the right way to use it. Parents, caregivers, and others who may be in a position to administer epinephrine injection, 0.3 mg (auto-injector) or epinephrine injection, 0.15 mg (auto-injector) should also understand how to use it as well. If you have any questions, ask your healthcare provider.

Your Epinephrine Injection, 0.3 mg (Auto-Injector) and Epinephrine Injection, 0.15 mg (Auto-Injector)

[ppi image]

A dose of epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg requires 3 steps: Prepare, Administer and Get emergency medical help

Step 1. Prepare epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg for injection

[ppi ifu image 1]

Quickly twist the yellow cap off the epinephrine injection, 0.3 mg auto-injector or the green cap off the epinephrine injection, 0.15 mg auto-injector in the direction of the “twist arrow” to remove it.

[ppi ifu image 2]

Grasp the auto-injector in your fist with the orange tip (needle end) pointing downward. With your other hand, pull off the blue safety release.

Important: The blue safety release is a small part that may become a choking hazard for children. Throw away the blue safety release immediately after using epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg.

Note:

The needle comes out of the orange tip.

To avoid an accidental injection, never put your thumb, fingers or hand over the orange tip. If an accidental injection happens, get emergency medical help right away.

Step 2. Administer epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg

If you are administering epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg to a young child, hold the leg firmly in place while administering an injection.

[1]

Place the orange tip against the middle of the outer thigh (upper leg) at a right angle

(perpendicular) to the thigh.

Swing and push the auto-injector firmly until it 'clicks'.

The click signals that the injection has started.

[1]

Hold firmly in place for 3 seconds (count slowly 1,2,3).

The injection is now complete.

[1]

Remove the auto-injector from the thigh. The orange tip will extend to cover the needle. If the needle is still visible, do not attempt to reuse it.

[1]

Massage the injection area for 10 seconds.

Step 3. Get emergency medical help now. You may need further medical attention. You may need to use a second epinephrine injection, 0.3 mg (auto-injector) or epinephrine injection, 0.15 mg (auto-injector) if symptoms continue or recur.

Take your used auto-injector with you when you go to see a healthcare provider.

Tell the healthcare provider that you have received an injection of epinephrine. Show the healthcare provider where you received the injection.

Give your used epinephrine injection, 0.3 mg (auto-injector) or epinephrine injection, 0.15 mg (auto-injector) to the healthcare provider for inspection and proper disposal.

Ask for a refill, if needed.

Note:

Keep the Epinephrine Injection Trainer and the real epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg Auto-Injectors away from young children. The real epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg Auto-Injectors and Epinephrine Injection Trainer are not toys. For young children, use of the Epinephrine Injection Trainer and the real epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg Auto-Injectors should be supervised by an adult.

A carrying tube is not provided as seen with other products.

Epinephrine injection, 0.3 mg and epinephrine injection, 0.15 mg are single-dose injectable devices that deliver a fixed dose of epinephrine. The auto-injector cannot be reused. Do not attempt to reuse epinephrine injection after the device has been activated. It is normal for most of the medicine to remain in the auto-injector after the dose is injected. The correct dose has been administered if the orange needle tip is extended and the window is blocked.

Your epinephrine injection, 0.3 mg (auto-injector) and epinephrine injection, 0.15 mg (auto-injector) may come packaged with an Epinephrine Injection Trainer and separate Epinephrine Injection Trainer Instructions for Use. The Epinephrine Injection Trainer has a grey color. The grey Epinephrine Injection Trainer contains no medicine and no needle. Practice with your Epinephrine Injection Trainer, but always carry your real epinephrine injection, 0.3 mg auto-injector or epinephrine injection, 0.15 mg auto-injector in case of an allergic emergency.

If you are administering epinephrine injection, 0.3 mg or epinephrine injection, 0.15 mg

to a young child, ask your healthcare provider to show you how to (1) hold the young child firmly in place (restrain), (2) with 1 hand, quickly twist the yellow cap off the epinephrine injection, 0.3 mg (auto-injector) or the green cap off the epinephrine injection, 0.15 mg (auto-injector) in the direction of the “twist arrow” to remove it, (3) grasp the auto-injector in your fist with the orange tip (needle end) pointing downward and (4) with your other hand, pull off the blue safety release, and how to properly hold the leg in place while administering a dose.

Do not try to take the epinephrine injection, 0.3 mg (auto-injector) or epinephrine injection, 0.15 mg (auto-injector) apart.

This Patient Information and Instructions for Use have been approved by the U.S. Food and Drug Administration.

KEEP THIS AND ALL MEDICATIONS OUT OF THE REACH OF CHILDREN.

Epinephrine Injection USP, 0.3 mg (Auto-Injector)

Epinephrine Injection USP, 0.3 mg = one dose of 0.3 mg epinephrine USP, 0.3 mg/0.3 mL

Epinephrine Injection USP, 0.15 mg (Auto-Injector)

Epinephrine Injection USP, 0.15 mg = one dose of 0.15 mg epinephrine USP, 0.15 mg/0.3 mL

TevaEpinephrine.com

Register your Epinephrine Injection 0.3 mg (Auto-Injector) or Epinephrine Injection 0.15 mg (Auto-Injector) at www.TevaEpinephrine.com and find out more about:

Free Epinephrine Injection Auto-Injector Refill Reminder Program. It is important to keep your auto-injector up-to-date.

Register up to 6 Epinephrine Injection 0.3 mg (Auto-Injectors) or Epinephrine Injection 0.15 mg (Auto-Injectors) and receive automatic Refill Reminder Alerts.

Receive periodic information related to allergies and allergens.

Instructional Video

For more information about Epinephrine Injection 0.3 mg (Auto-Injector) or Epinephrine Injection 0.15 mg (Auto-Injector) and proper use of the products, call Teva at 1-888-838-2872 or visit www.TevaEpinephrine.com.

Epinephrine Injection Trainer Instructions for Use

In an emergency: Do not use the grey Trainer. Use your real yellow Epinephrine Injection 0.3 mg Auto-Injector or real green Epinephrine Injection 0.15 mg Auto-Injector.

Important Information

The Trainer label has a grey color.

The Trainer contains no medicine and no needle.

Keep the grey Trainer away from young children. The Trainer is not a toy. Young children should only practice with the Trainer under adult supervision.

Periodically practice with the grey Trainer before an allergic emergency (anaphylaxis) happens to make sure you are able to safely use the real yellow epinephrine injection 0.3 mg auto-injector or real green epinephrine injection 0.15 mg auto-injector in case of an

emergency.

Always carry your real yellow epinephrine injection 0.3 mg auto-injector or real green epinephrine injection 0.15 mg auto-injector in case of an allergic emergency.

Small parts like the blue safety release may become a choking hazard for children. Put the blue safety release back on the Trainer and reset it immediately after practicing.

The Epinephrine Injection Trainer

Familiarize yourself with this grey Trainer. Practice until you are comfortable using it.

Your grey Trainer: [trainer-ifu image]

- Never put your thumb, other fingers, or hand over the orange tip (below grey safety cap).
- The orange tip is where the needle comes out of your epinephrine injection 0.3 mg auto-injector or epinephrine injection 0.15 mg auto-injector.

Practice Instructions

[1]

1 Prepare the Trainer for Simulated Injection

Grasp the grey Trainer in your fist with the orange tip pointing downward and twist off Grey Cap in the direction of “twist arrow”.

With your other hand, pull off blue safety release.

Removing the blue safety release unlocks the Trainer.

2 Administer the Trainer Simulation

If practicing with a young child, hold the leg firmly in place while using the Epinephrine Injection Trainer.

Ask your healthcare provider to show you how to:

1. Hold the young child firmly in place (restrain).
2. With 1 hand, quickly twist the grey cap off the Epinephrine Injection Trainer in the direction of the “twist arrow” to remove it.
3. Grasp the auto-injector in your fist with the orange tip (needle end) pointing downward.
4. With your other hand, pull off the blue safety release, and how to properly hold the leg to practice so that you will be prepared before an allergic emergency happens.

[1] [1]

Place the orange tip against the middle of the outer thigh (upper leg) at a right angle (perpendicular) to the thigh.

Swing and push the Trainer firmly until it ‘clicks’. The click signals that the injection has started.

Hold firmly in place for 3 seconds (count slowly 1,2,3).

Remove the Trainer from the thigh and massage the injection area for 10 seconds. The orange tip automatically extends out after use.

Note:

In an actual emergency, you would need to seek emergency medical help right away. The actual auto-injector is made to work through clothing. Do not inject into any other part of the body.

[1]

3 To reset the Trainer

Put the blue safety release back on the Trainer.
Replace Grey Cap.

NOTE: With the real yellow Epinephrine Injection 0.3 mg Auto-Injector or real green Epinephrine Injection 0.15 mg Auto-Injector, the orange tip covers the needle after self-injection to help protect you from accidentally sticking yourself or others.

Practice Session Information

In case of an allergic emergency, use the real yellow Epinephrine Injection 0.3 mg Auto-Injector or real green Epinephrine Injection 0.15 mg Auto-Injector and not the grey Trainer.

Follow instructions above. Repeat as often as needed until you are able to self-inject quickly and correctly.

Reread:

The Trainer Instructions for Use

The “Patient Information” that comes with your Epinephrine Injection 0.3 mg Auto-Injector or Epinephrine Injection 0.15 mg Auto-Injector

Train others who could help you in an emergency:

- Your parents, caregivers, and others who may be in a position to administer epinephrine injection 0.3 mg or epinephrine injection 0.15 mg should know how

to help you during an allergic emergency (anaphylaxis). Before an emergency occurs, have them:

Practice activating the Trainer

Read these Trainer Instructions and the “Patient Information”

For more information about the Epinephrine Injection 0.3 mg Auto-Injector and Epinephrine Injection 0.15 mg Auto-Injector and the proper use of the products, go to www.TevaEpinephrine.com.

Caution:

Important differences between the Trainer and your real yellow Epinephrine Injection 0.3 mg (Auto-Injector) or real green Epinephrine Injection 0.15 mg (Auto-Injector)

[1] [0.3 mg injector] [0.15 mg injector]

Trainer

(Grey)

Epinephrine Injection 0.3 mg

(Yellow)

Epinephrine Injection 0.15 mg

(Green)

Contains medicine?

No

Yes

Yes

Has needle?

No

Yes

Yes

Comes in Carrier Tube?

No

No

No

Color of label

Grey

Yellow

Green

Has expiration date?

No

Yes

Yes

Can be reused?

Yes

No (use only one time)

No (use only one time)

Okay to remove and replace cap and/or safety release?

Yes

No (remove just one time
before use)

No (remove just one time
before use)

Pressure needed to hold against thigh?

Moderate

Strong

Strong

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

KEEP THIS AND ALL MEDICATIONS OUT OF THE REACH OF CHILDREN.

Repackaged and Distributed By:

Remedy Repack, Inc.

625 Kolter Dr. Suite #4 Indiana, PA 1-724-465-

DRUG: Epinephrine

GENERIC: Epinephrine

DOSAGE: INJECTION

ADMINSTRATION: INTRAMUSCULAR

NDC: 70518-3173-0

PACKAGING: 0.3 mL in 1 SYRINGE

OUTER PACKAGING: 1 in 1 CONTAINER

OUTER PACKAGING: 2 in 1 CARTON

ACTIVE INGREDIENT(S):

- EPINEPHRINE 0.15mg in 0.3mL

INACTIVE INGREDIENT(S):

- SODIUM CHLORIDE
- SODIUM METABISULFITE
- SODIUM TARTRATE DIHYDRATE
- HYDROCHLORIC ACID
- WATER

EPINEPHrine

For allergic emergencies (anaphylaxis)

0.15mg / 0.3 mL

Single-Dose Auto-Injectors

QTY: 2 - Pack

Two green single-dose auto injectors- One grey trainer; Should be used under adult supervision



RX ONLY

NDC #: 70518-3173-00

Expires:

LOT #:

Source NDC: 00093-5985-27

MFG: Teva, Frazer, PA 19355

Keep this and all medication out of the reach of children



WARNING: The needle comes out of the orange tip. Never put thumb, fingers or hand over orange tip. Do NOT remove the blue safety release until ready to use.

Directions For Use: See Package Insert

Store at 20-25°C (68-77°F); excursions permitted to 15-30°C (59-86°F) [See USP]

Repackaged by: RemedyRepack Inc., Indiana, PA 15701, 724.465.8762

EPINEPHRINE

epinephrine injection

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70518-3173(NDC:0093-5985)
Route of Administration	INTRAMUSCULAR, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
EPINEPHRINE (UNII: YKH834O4BH) (EPINEPHRINE - UNII:YKH834O4BH)	EPINEPHRINE	0.15 mg in 0.3 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
SODIUM TARTRATE DIHYDRATE (UNII: DIA7C37AOW)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
WATER (UNII: 059QF0KOOR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70518-3173-0	2 in 1 CARTON	07/27/2021	
1		1 in 1 CONTAINER		
1		0.3 mL in 1 SYRINGE; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090589	07/27/2021	

Labeler - REMEDYREPACK INC. (829572556)

Revised: 3/2024

REMEDYREPACK INC.