

**EPINEPHRINE- epinephrine injection, solution
HF Acquisition Co LLC, DBA HealthFirst**

EPINEPHRINE INJECTION, USP 1mg/10mL (0.1mg/mL) 10mL SYR

HIGHLIGHTS OF PRESCRIBING INFORMATION

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

EPINEPHRINE INJECTION, for intravenous use
Initial U.S. Approval: 1939

INDICATIONS AND USAGE

Epinephrine is a non-selective alpha- and beta-adrenergic agonist indicated to increase mean arterial blood pressure in adult patients with hypotension associated with septic shock. (1.1)

DOSAGE AND ADMINISTRATION

Hypotension associated with septic shock:

Dilute epinephrine in dextrose solution prior to infusion. (2.2)

Infuse epinephrine into a large vein. (2.2)

Intravenous infusion rate of 0.05 mcg/kg/min to 2 mcg/kg/min, titrated to achieve desired mean arterial pressure. (2.2)

Wean gradually. (2.2)

DOSAGE FORMS AND STRENGTHS

Injection: 1 mg/10 mL (0.1 mg/mL) in ABBOJECT® Syringe. (3)

CONTRAINDICATIONS

None. (4)

WARNINGS AND PRECAUTIONS

Monitor patient for acute severe hypertension. (5.1)

Potential for pulmonary edema, which may be fatal. (5.2)

May induce potentially serious cardiac arrhythmias and myocardial ischemia, particularly in patients with underlying heart disease. (5.3)

Avoid extravasation into tissues, which can cause local necrosis. (5.4)

Potential for oliguria or renal impairment. (5.5)

Presence of sulfite in this product should not deter use for hypotension associated with septic shock (5.6)

ADVERSE REACTIONS

Most common adverse reactions to systemically administered epinephrine are headache; anxiety; apprehensiveness; restlessness; tremor; weakness; dizziness; diaphoresis;

nausea/vomiting; and/or respiratory difficulties. Arrhythmias, including fatal ventricular fibrillation, rapid rises in blood pressure producing cerebral hemorrhage, and angina have occurred. (6)

To report SUSPECTED ADVERSE REACTIONS, contact Hospira, Inc. at 1-800-441-4100, or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

Drugs that counter the pressor effects of epinephrine include alpha blockers, vasodilators such as nitrates, diuretics, antihypertensives, and ergot alkaloids. (7.1)

Drugs that potentiate the effects of epinephrine include sympathomimetics, beta blockers, tricyclic antidepressants, MAO inhibitors, COMT inhibitors, clonidine, doxapram, oxytocin, levothyroxine sodium, and certain antihistamines. (7.2)

Drugs that increase the arrhythmogenic potential of epinephrine include beta blockers, cyclopropane and halogenated hydrocarbon anesthetics, quinidine, antihistamines, exogenous thyroid hormones, diuretics, and cardiac glycosides. Observe for development of cardiac arrhythmias. (7.3)

Potassium-depleting drugs, including corticosteroids, diuretics, and theophylline, potentiate the hypokalemic effects of epinephrine. (7.4)

USE IN SPECIFIC POPULATIONS

Pregnancy: Epinephrine may cause fetal harm. (8.1)

Elderly patients and pregnant women may be at greater risk of developing adverse reactions when epinephrine is administered parenterally. (8.1, 8.5)

Revised: 11/2019

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1. INDICATIONS & USAGE

1.1 Hypotension associated with Septic Shock

Epinephrine Injection USP, 1 mg/10 mL (0.1 mg/mL) is indicated to increase mean arterial blood pressure in adult patients with hypotension associated with septic shock.

2. DOSAGE & ADMINISTRATION

2.1 General Considerations

Inspect visually for particulate matter and discoloration prior to administration, whenever solution and container permit. Do not use if the solution is colored or cloudy, or if it contains particulate matter. Discard all unused drug.

2.2 Hypotension associated with Septic Shock

Dilute 10 mL (1 mg) of epinephrine from the syringe in 1,000 mL of 5 percent dextrose solution or 5 percent dextrose and sodium chloride solution to produce a 1 mcg per mL dilution. The diluted solutions can be stored for up to 4 hours at room temperature or 24 hours under refrigerated conditions.

Administration in saline solution alone is not recommended. If indicated, administer whole blood or plasma separately.

Whenever possible, give infusions of epinephrine into a large vein. Avoid using a catheter tie-in technique, because the obstruction to blood flow around the tubing may cause stasis and increased local concentration of the drug. Avoid the veins of the leg in elderly patients or in those suffering from occlusive vascular diseases.

To provide hemodynamic support in septic shock associated hypotension in adult

patients, the suggested dosing infusion rate of intravenously administered epinephrine is 0.05 mcg/kg/min to 2 mcg/kg/min, and is titrated to achieve a desired mean arterial pressure (MAP). The dosage may be adjusted periodically, such as every 10 – 15 minutes, in increments of 0.05 mcg/kg/min to 0.2 mcg/kg/min, to achieve the desired blood pressure goal.

After hemodynamic stabilization, wean incrementally over time, such as by decreasing doses of epinephrine every 30 minutes over a 12- to 24-hour period.

3. DOSAGE FORMS AND STRENGTHS

Injection solution: 10 mL clear glass vial containing 1 mg/10 mL (0.1 mg/mL) epinephrine in a sterile, nonpyrogenic, clear and colorless solution. Each vial is co-packaged with an injector, which together make a single-dose ABBOJECT® Syringe.

4. CONTRAINDICATIONS

None.

5. WARNINGS AND PRECAUTIONS

5.1 Hypertension

Because individual response to epinephrine may vary significantly, monitor blood pressure frequently and titrate to avoid excessive increases in blood pressure.

Patients receiving monoamine oxidase inhibitors (MAOI) or antidepressants of the triptyline or imipramine types may experience severe, prolonged hypertension when given epinephrine.

5.2 Pulmonary Edema

Epinephrine increases cardiac output and causes peripheral vasoconstriction, which may result in pulmonary edema.

5.3 Cardiac Arrhythmias and Ischemia

Epinephrine may induce cardiac arrhythmias and myocardial ischemia in patients, especially patients with coronary artery disease, or cardiomyopathy [see ADVERSE REACTIONS and DRUG INTERACTIONS (7.3)].

5.4 Extravasation and Tissue Necrosis with Intravenous Infusion

Avoid extravasation of epinephrine into the tissues, to prevent local necrosis. When Epinephrine Injection is administered intravenously, check the infusion site frequently for free flow. Blanching along the course of the infused vein, sometimes without obvious extravasation, may be attributed to vasa vasorum constriction with increased permeability of the vein wall, permitting some leakage. This also may progress on rare occasions to a superficial slough. Hence, if blanching occurs, consider changing the infusion site at intervals to allow the effects of local vasoconstriction to subside.

There is potential for gangrene in a lower extremity when infusions of catecholamine are given in an ankle vein.

Antidote for Extravasation Ischemia: To prevent sloughing and necrosis in areas in which extravasation has taken place, infiltrate the area with 10 mL to 15 mL of saline solution containing from 5 mg to 10 mg of phentolamine, an adrenergic blocking agent. Use a syringe with a fine hypodermic needle, with the solution being infiltrated liberally throughout the area, which is easily identified by its cold, hard, and pallid appearance. Sympathetic blockade with phentolamine causes immediate and conspicuous local hyperemic changes if the area is infiltrated within 12 hours.

5.5 Renal Impairment

Epinephrine constricts renal blood vessels, which may result in oliguria or renal impairment.

5.6 Allergic Reactions associated with Sulfite

Epinephrine Injection contains sodium metabisulfite which may cause mild to severe allergic reactions including anaphylaxis or asthmatic episodes, particularly in patients with a history of allergies. The presence of sodium metabisulfite in this product should not preclude its use for the treatment of hypotension associated with septic shock even if the patient is sulfite-sensitive, as the alternatives to using epinephrine in a life-threatening situation may not be satisfactory. In susceptible patients, consider using a formulation of epinephrine or another vasoconstrictor that does not contain sodium metabisulfite.

6. ADVERSE REACTIONS

The following adverse reactions are discussed elsewhere in labeling:

Hypertension [see WARNINGS AND PRECAUTIONS (5.1)]

Pulmonary Edema [see WARNINGS AND PRECAUTIONS (5.2)]

Cardiac Arrhythmias and Ischemia [see WARNINGS AND PRECAUTIONS (5.3)]

Extravasation and Tissue Necrosis with Intravenous Infusion [see WARNINGS AND PRECAUTIONS (5.4)]

Renal Impairment [see WARNINGS AND PRECAUTIONS (5.5)]

Allergic Reactions associated with Sulfite [see WARNINGS AND PRECAUTIONS (5.6)]

The following adverse reactions have been associated with use of epinephrine. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to estimate their frequency reliably or to establish a causal relationship to drug exposure.

Cardiovascular disorders: tachycardia, supraventricular tachycardia, ventricular arrhythmias (including fatal ventricular fibrillation), myocardial ischemia, myocardial infarction, limb ischemia, pulmonary edema, hypertension

Gastrointestinal disorders: nausea, vomiting

General disorders and administrative site conditions: extravasation

Metabolic: insulin resistance, hypokalemia, lactic acidosis

Nervous system disorders: headache, paresthesia, tremor, stroke, central nervous system bleeding, weakness, dizziness, disorientation, impaired memory, panic, psychomotor agitation, somnolence

Psychiatric disorders: excitability, anxiety, apprehensiveness, nervousness, restlessness

Renal disorders: renal insufficiency

Respiratory: rales

Skin and subcutaneous tissue disorders: diaphoresis, pallor, piloerection, skin blanching, skin necrosis with extravasation.

7. DRUG INTERACTIONS

7.1 Drugs Antagonizing Pressor Effects of Epinephrine

α-blockers, such as phentolamine

Vasodilators, such as nitrates

Diuretics

Antihypertensives

Ergot alkaloids

Phenothiazine antipsychotics

7.2 Drugs Potentiating Pressor Effects of Epinephrine

Sympathomimetics

β-blockers, such as propranolol

Tricyclic anti-depressants

Monoamine oxidase (MAO) inhibitors

Catechol-O-methyl transferase (COMT) inhibitors, such as entacapone

Clonidine

Doxapram

Oxytocin

7.3 Drugs Potentiating Arrhythmogenic Effects of Epinephrine

Patients who are concomitantly receiving any of the following drugs should be observed carefully for the development of cardiac arrhythmias [see WARNINGS AND PRECAUTIONS (5.5) and ADVERSE REACTIONS (6)].

β-blockers, such as propranolol

Cyclopropane or halogenated hydrocarbon anesthetics, such as halothane

Antihistamines

Thyroid hormones

Diuretics

Cardiac glycosides, such as digitalis glycosides

Quinidine

7.4 Drugs Potentiating Hypokalemic Effects of Epinephrine

Potassium depleting diuretics

Corticosteroids

Theophylline

8. USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Limited published data on epinephrine use in pregnant women are not sufficient to determine a drug-associated risk for major birth defects or miscarriage. However, there are risks to the mother and fetus associated with epinephrine use during labor or delivery, and risks due to untreated hypotension associated with septic shock (see CLINICAL CONSIDERATIONS below). In animal reproduction studies, epinephrine demonstrated adverse developmental effects when administered to pregnant rabbits (gastroschisis), mice (teratogenic effects, embryonic lethality, and delayed skeletal ossification), and hamsters (embryonic lethality and delayed skeletal ossification) during organogenesis at doses approximately 15 times, 3 times and 2 times, respectively, the maximum recommended daily intramuscular or subcutaneous dose (see DATA below).

All pregnancies have a background risk of birth defects, loss, or other adverse outcomes. The estimated 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.

Clinical Considerations

Disease-associated maternal and/or embryo/fetal risk

Hypotension associated with septic shock is a medical emergency in pregnancy which can be fatal if left untreated. Delaying treatment in pregnant women with hypotension associated with septic shock may increase the risk of maternal and fetal morbidity and mortality. Do not withhold life-sustaining therapy for a pregnant woman.

Labor or Delivery

Epinephrine usually inhibits spontaneous or oxytocin induced contractions of the pregnant human uterus and may delay the second stage of labor. Avoid epinephrine during the second stage of labor. In dosage sufficient to reduce uterine contractions, the drug may cause a prolonged period of uterine atony with hemorrhage. Avoid epinephrine in obstetrics when maternal blood pressure exceeds 130/80 mmHg.

Although epinephrine improves maternal hypotension associated with anaphylaxis, it may result in uterine vasoconstriction, decreased uterine blood flow, and fetal anoxia.

Data

Animal Data

In a study in pregnant rabbits administered 1.2 mg/kg/day epinephrine (approximately 15 times the maximum recommended intramuscular or subcutaneous dose on a mg/m² basis) subcutaneously during organogenesis (on days 3 to 5, 6 to 7 or 7 to 9 of gestation), epinephrine caused teratogenic effects (including gastroschisis). Animals treated on days 6 to 7 had decreased number of implantations.

In a teratology study, pregnant mice were subcutaneously administered epinephrine (0.1 to 10 mg/kg/day) on Gestation Days 6 to 15. Teratogenic effects, embryonic lethality, and delays in skeletal ossification were observed at approximately 3 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 2 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).

Subcutaneous administration of epinephrine to pregnant hamsters at a dose of 0.5 mg/kg/day (approximately 2 times the maximum recommended intramuscular or subcutaneous dose on a mg/m² basis) on Gestation Days 7 to 10 resulted in delayed skeletal ossification and a reduction in litter size.

8.2 Lactation

Risk Summary

There is no information regarding the presence of epinephrine in human milk, or the effects of epinephrine on the breastfed infant or on milk production. However, due to its poor oral bioavailability and short half-life, epinephrine exposure is expected to be very low in the breastfed infant. The lack of clinical data during lactation precludes a clear determination of the risk of epinephrine to a breastfed infant.

8.4 Pediatric Use

Safety and effectiveness in pediatric patients have not been established.

8.5 Geriatric Use

Clinical studies of epinephrine for the treatment of hypotension associated with septic shock did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects. Other reported clinical experience has not identified differences in responses between the elderly and younger patients. In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy.

CLOSE

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. Epinephrine overdosage may 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. Myocardial ischemia and infarction, cardiomyopathy, extreme pallor and coldness of the skin, metabolic acidosis due to elevated blood lactic acid levels, and renal insufficiency and failure have also been reported.

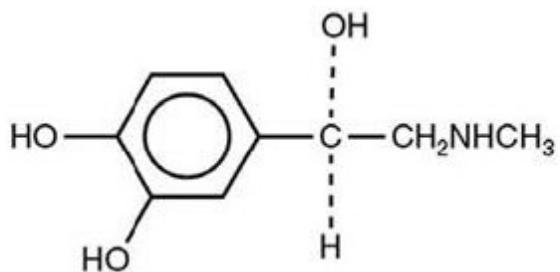
Epinephrine is rapidly inactivated in the body and treatment following overdose is primarily supportive. Treatment of pulmonary edema consists of a rapidly acting alpha-adrenergic blocking drug (such as phentolamine mesylate) and respiratory support. Treatment of arrhythmias consists of administration of a beta-adrenergic blocking drug (such as propranolol). If necessary, pressor effects may be counteracted by rapidly

acting vasodilators (such as nitrites) or alpha-adrenergic blocking drugs. If prolonged hypotension follows such measures, it may be necessary to administer another pressor drug.

11. DESCRIPTION

Epinephrine Injection USP, 1 mg/10 mL (0.1 mg/mL) is supplied as a sterile aqueous solution with a pH range of 2.3-3.5, that is clear, colorless, and nonpyrogenic. Each milliliter contains 0.1 mg epinephrine, 8.16 mg sodium chloride, 0.46 mg sodium metabisulfite, and 2.13 mg citric acid, anhydrous and 0.41 mg sodium citrate, dihydrate added as buffers. Additional citric acid and/or sodium citrate may be added for pH adjustment. The solution contains no preservatives.

This sterile solution is to be administered after dilution by the intravenous route. Epinephrine is a sympathomimetic catecholamine (adrenergic agent) designated chemically as 4-[1-hydroxy-2-(methylamino) ethyl]-1,2 benzenediol, a white, crystalline powder. It has the following structural formula:



The molecular weight of epinephrine is 183.2.

Epinephrine solution deteriorates rapidly on exposure to air or light, turning pink from oxidation to adrenochrome and brown from the formation of melanin.

12. CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Epinephrine acts on both alpha (α)- and beta (β)-adrenergic receptors. The mechanism of the rise in blood pressure is 3-fold: a direct myocardial stimulation that increases the strength of ventricular contraction (positive inotropic action), an increased heart rate (positive chronotropic action), and peripheral vasoconstriction.

12.2 Pharmacodynamics

Following intravenous administration of epinephrine, increases in systolic blood pressure and heart rate are observed. Decreases in systemic vascular resistance and diastolic blood pressure are observed at low doses of epinephrine because of β_2 -mediated vasodilation, but are overtaken by α_1 -mediated peripheral vasoconstriction at higher doses leading to increase in diastolic blood pressure. The onset of blood pressure increase following an intravenous dose of epinephrine is < 5 minutes and the time to offset blood pressure response occurs within 20 min. Most vascular beds are constricted including renal, splanchnic, mucosal and skin.

Epinephrine increases glycogenolysis, reduces glucose uptake by tissues, and inhibits insulin release in the pancreas, resulting in hyperglycemia and increased blood lactic acid.

12.3 Pharmacokinetics

When administered parenterally epinephrine has a rapid onset and short duration of action.

Following intravenous injection, epinephrine is rapidly cleared from the plasma with an effective half-life of < 5 min. A pharmacokinetic steady state following continuous intravenous infusion is achieved within 10–15 min. In patients with septic shock, epinephrine displays dose-proportional pharmacokinetics in the infusion dose range of 0.03 to 1.7 mcg/kg/min.

Epinephrine is extensively metabolized with only a small amount excreted unchanged. Epinephrine is rapidly degraded to vanillylmandelic acid, an inactive metabolite, by monoamine oxidase and catechol-O-methyltransferase that are abundantly expressed in the liver, kidneys and other extraneuronal tissues. The tissues with the highest contribution to removal of circulating exogenous epinephrine are the liver (32%), kidneys (25%), skeletal muscle (20%), and mesenteric organs (12%).

Specific Populations

Elderly

In a pharmacokinetic study of 45-minute epinephrine infusions given to healthy men aged 20 to 25 years and healthy men aged 60 to 65 years, the mean plasma metabolic clearance rate of epinephrine at steady state was greater among the older men (144.8 versus 78 mL/kg/min for a 14.3 ng/kg/min infusion).

Body Weight

Body weight has been found to influence epinephrine pharmacokinetics. Higher body weight was associated with a higher plasma epinephrine clearance and a lower concentration plateau.

13. NON CLINICAL 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 a mutagen based on the E. coli WP2 Mutoxitest bacterial reverse mutation assay.

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 (15-fold the highest human intramuscular or subcutaneous daily dose) during gestation days 3 to 9.

13.2 Animal Toxicology and/or Pharmacology

Epinephrine was associated with metabolic effects, decreased mesentery, coronary and renal conductance in a sheep model of septic shock. Data from hemolysis study have shown that epinephrine at 1:1000 dilution is non-hemolytic. Epinephrine infusion significantly increased the MAP (69 vs. 86 mmHg) and cardiac output (6.4 vs. 7.1 L/min) and decreased renal blood flow (330 vs. 247 mL/min).

14. CLINICAL STUDIES

14.1 Hypotension associated with Septic Shock

Fourteen clinical studies from the literature documented that epinephrine increases the mean arterial pressure (MAP) in patients with hypotension associated with septic shock.

16. HOW SUPPLIED/STORAGE AND HANDLING

Epinephrine Injection 1 mg/10 mL (0.1 mg/mL) is a clear and colorless solution available in glass vials. Each vial is co-packaged with an injector, which together make a single-dose ABBOJECT® Syringe. It is supplied in the following presentation.

Unit of Sale	Concentration
NDC 0409-4933-01	1 mg/10 mL
Bundle of 10 Single-dose ABBOJECT® Glass Syringes (10 mL each)	(0.1 mg/mL)

Epinephrine is light sensitive. Protect from light until ready to use.

Do not refrigerate. Protect from freezing.

Store at room temperature, between 20°C to 25°C (68°F to 77°F). (See USP Controlled Room Temperature.)

Protect from alkalis and oxidizing agents.

Inspect visually for particulate matter and discoloration prior to administration. Do not use the solution if it is colored or cloudy, or if it contains particulate matter.

SPL UNCLASSIFIED

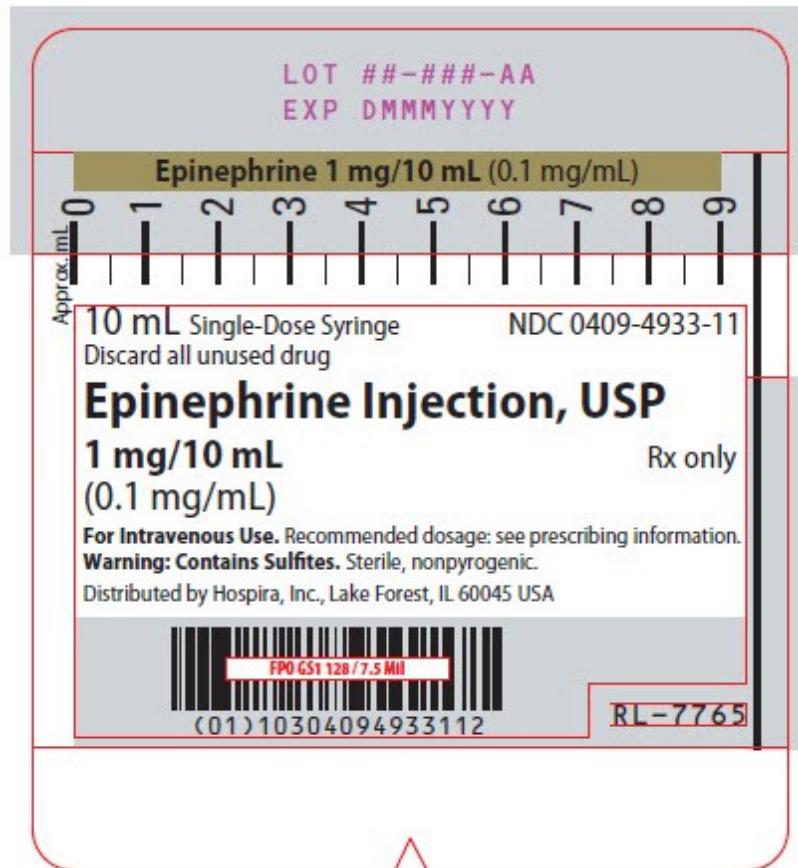


Distributed by Hospira, Inc., Lake Forest, IL 60045 USA

ABBOJECT® is a trademark of Abbott Laboratories.

LAB-0928-2.0

PRINCIPAL DISPLAY PANEL - SYRINGE LABELING



**PRINCIPAL DISPLAY PANEL - 51662-1531-1 CARTON LABELING AND
SERIALIZED LABEL ON CARTON**

CAUTION: Liquid in glass. Handle with care. Inspect vial for damage

USE ASEPTIC TECHNIQUE

prior to assembly.

- Do not assemble until ready to use.**



Insert vial into injector **without exerting excessive force.**
Ensure that vial and injector are properly aligned.
Gently rotate vial clockwise (about 3 turns) until medication enters needle.
If resistance is encountered, remove

ויליאם הנרי פולמן

adapter, push yellow hood in and then twist *counterclockwise*.

To access **needle**, twist and pull yellow hood clockwise to remove hood and green adapter.

4. Apply gentle downward pressure on vial to initiate liquid flow. **DO NOT APPLY EXCESSIVE FORCE TO VIAL.**

Each ml contains epinephrine 0.1 mg; citric acid, anhydrous 2.13 mg; sodium chloride 1.6 mg; sodium citrate, dihydrate 0.41 mg and sodium metabisulfite added 0.06 mg. Contains no preservative and/or flavoring agent. Redesignation. Fluid bath and needle undiluted and nonhypotensive if used as needed cover are sterile and packaged sterile.

Epinephrine
Injection, USP
1 mg/10 mL
(0.1 mg/mL)
Hospira

10 mL NDC 0409-4933-11

NDC 0409-4933-11
Epinephrine
Injection, USP
10 mL
1 mg/10 mL
(0.1 mg/mL)


10 mL
Epinephrine
Injection, USP
1 mg/10 mL
(0.1 mg/mL)
NDC 0409-491

OPEN

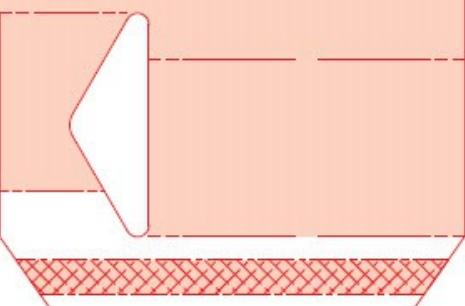
PRESS AND PULL TO OPEN

A standard linear barcode is positioned at the bottom right of the page. It consists of vertical black bars of varying widths on a white background.

Single-Dose Syringe

with male luer lock
adapter and 20-Gauge
protected needle

Single-Dose Syringe. Discard all unused drug. NOTE: Do not use if the solution is colored or cloudy, or if it contains particulate matter. For intravenous use. Recommended dosage: see prescribing information.
Protect from light by retaining product in carton until ready for use.
To prevent needle-stick injuries, needles should not be recapped, purposely bent or broken by hand. Store at 20°C to 25°C (68°F to 77°F). [See USP Controlled Room Temperature.] Do not refrigerate. Protect from freezing.



Epinephrine Injection, USP

1 mg/10 mL
(0.1 mg/mL)

For Intravenous Use

Warning: Contains Sulfites

Protect from light

ABBOJECT®
Single-Dose Syringe

with male luer lock
adapter and 20-Gauge
protected needle

Hospira

LOT 24077DK
EXP 1 MAR 2022

Single-Dose Syringe. Discard all unused drug. NOTE: Do not use if the syringe is colored or cloudy, or if it contains particulate matter. For intravenous Recommended dosage: see prescribing information.

Protect from light by retaining product in carton until ready for use.

To prevent needle-stick injuries, needles should not be recapped, purged or bent or broken by hand. Store at 20°C to 25°C (68°F to 77°F). [See USP Controlled Room Temperature.] Do not refrigerate. Protect from freezing.

RX ONLY



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manufacturer's package insert
distributed by HF Acquisition Co., LLC
Seattle, WA 98275

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◀ PRESS AND PULL TO OPEN

Principal Display Panel - 51662-1531-2

Outer Pouch Labeling

**EPINEPHRINE INJECTION, USP 1mg/10mL
(0.1mg/mL) 10mL SYR**

**NDC : 51662-1531-2
LOT : 12345
EXP : 2030-01-01**

(01) 00351662153123
(10) 12345
(17) 300101
(21) 222127316592

FOR IV USE. PROTECT FROM LIGHT.TO PREVENT NEEDLE-STICK INJURIES,
NEEDLES SHOULD NOT BE RECAPPED, PURPOSELY BENT OR BROKEN BY
HAND.CONTAINS SULFITES.DISCARD ALL UNUSED DRUG.NOTE: DO NOT USE IF
THE SOLUTION IS COLORED OR CLOUDY, OR IF IT CONTAINS PARTICULATE
MATTER.RECOMMENDED DOSAGE: SEE INSERT.STORE AT 20°C TO 25°C (68°F TO
77°F).

See manufacturer's package insert

ORIGINAL MFG NDC: 0409-4933-01

RX ONLY

Manufactured by HF Acquisition Co., LLC
Mukilteo, WA 98275



CAUTION: Liquid in glass. Handle with care. Inspect vial for damage prior to assembly.

USE ASEPTIC TECHNIQUE

Do not assemble until ready to use.

1. Remove caps from vial and injector.



2. Insert vial into injector without exerting excessive force.

Ensure that vial and injector are properly aligned. Gently rotate vial clockwise (about 3 turns) until medication enters needle. If resistance is encountered, remove and repeat procedure.



3. To access **needle**, twist and pull yellow hood clockwise and then twist counterclockwise.
- OR
- To access **needle**, twist and pull yellow hood clockwise to remove hood and green adapter.



4. Apply gentle downward pressure on vial to initiate liquid flow. **DONOT APPLY EXCESSIVE FORCE TO VIAL**

CA-7403

Distributed by Hospira, Inc., Lake Forest, IL 60055 USA
Abbott Laboratories is a trademark of the Abbott group of companies.
MADE IN ITALY

Hospira

0.1 mg/mL
1 mg/10 mL
Hospira

10 mL NDC 0409-4933-11
Rx only
Epinephrine NDC 0409-4933-11

(0.1 mg/mL)
1 mg/10 mL
Hospira
Epinephrine
Injection, USP

10 mL NDC 0409-4933-11
Rx only
Epinephrine
Injection, USP

(0.1 mg/mL)
1 mg/10 mL
Hospira
Epinephrine
Injection, USP

For Intravenous Use
Warning: Contains Sulfites
Protect from light

ABPROJECT®
Single-Dose Syringe
with male luer lock
adapter and 20-Gauge
protected needle

Each mL contains epinephrine 0.1 mg; citric acid, anhydrous 2.13 mg;
sodium chloride 8.16 mg; sodium sulfite 0.46 mg; May contain aditives: 0.1 mg a and sodium
metabisulfite for pH adjustment. Content of this is no better or equivalent to an intermediate if
caps and needle cover are undisturbed and package is intact.

Single-Dose Syringe. Discard all unused drug. NOTE: Do not use if the solution is colored or cloudy, or if it contains particulate matter. For intravenous use. Recommended dosage see prescribing information. Protect from light by retaining product in carton until ready for use. To prevent needle-stick injuries, needles should not be recapped, purposely bent or broken by hand. Store at 20°C to 25°C (68°F to 77°F). [See USP Controlled Room Temperature.] Do not refrigerate. Protect from freezing.



(01)20304094933119

◀ PRESS AND PULL TO OPEN

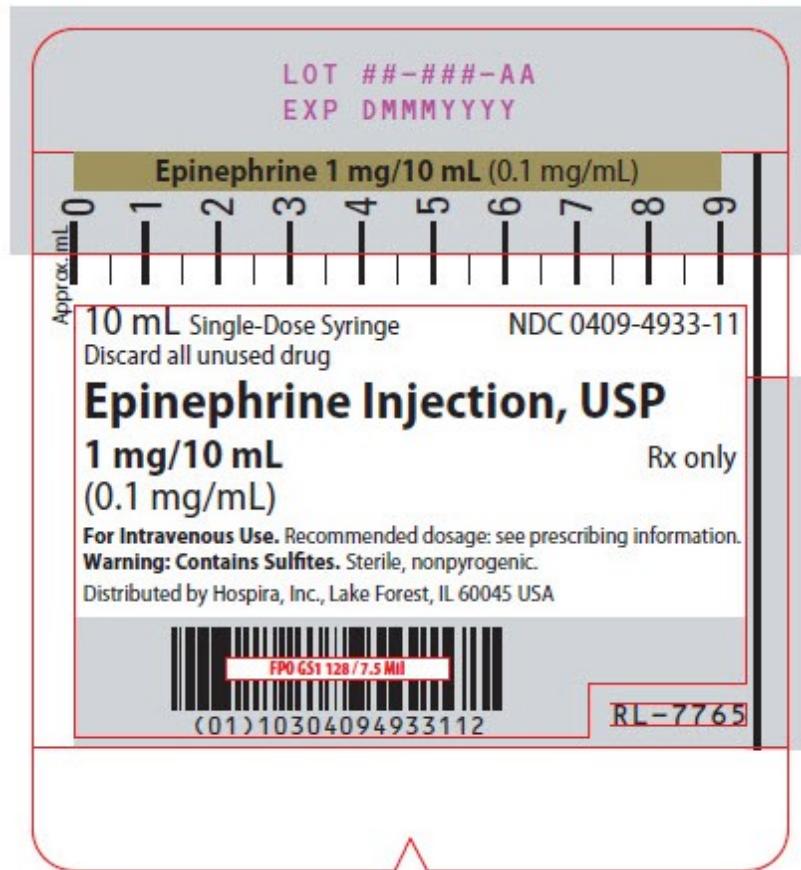
OPEN

10 mL
Epinephrine
Injection, USP
1 mg/10 mL
(0.1 mg/mL)

NDC 0409-4933-11

10 mL
Epinephrine
Injection, USP
1 mg/10 mL
(0.1 mg/mL)

NDC 0409-4933-11



Principal Display Panel - 51662-1531-3 - Carton of 50 Labeling

Case Labeling 51662-1531-3, Case of 50

**EPINEPHRINE INJECTION, USP 1mg/10mL(0.1mg/mL)
10mL SYR (BOX OF 50)**

NDC: 51662-1531-3

LOT: 123456

EXP: 2030-01-01

(01) 00351662153130

(10) 123456

(17) 300101

(21) 61099193435

CASE

FOR IV USE. PROTECT FROM LIGHT. TO PREVENT NEEDLE-STICK INJURIES,
NEEDLES SHOULD NOT BE RECAPPED, PURPOSELY BENT OR BROKEN BY
HAND. CONTAINS SULFITES. DISCARD ALL UNUSED DRUG. NOTE: DO NOT USE
IF THE SOLUTION IS COLORED OR CLOUDY, OR IF IT CONTAINS PARTICULATE
MATTER. RECOMMENDED DOSAGE: SEE INSERT. STORE AT 20°C TO 25°C (68°F
TO 77°F).

See manufacturer's package insert

ORIGINAL MFG NDC: 0409-4933-01

RX ONLY

Manufactured by HF Acquisition Co., LLC
Mukilteo, WA 98275



Serialized Label on Case

(01)00351662153130
(17)230919
(21)351662141212
(10)HF1234TESTING

SEE MANUFACTURE'S INSERT
DISTRIBUTED BY HF ACQUISITIONS CO., LLC
MUKILTEO, WA 98275

RX ONLY



EPINEPHRINE

epinephrine injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:51662-1531(NDC:0409-4933)
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
EPINEPHRINE (UNII: YKH834O4BH) (EPINEPHRINE - UNII:YKH834O4BH)	EPINEPHRINE	0.1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	8.16 mg in 1 mL
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	0.46 mg in 1 mL
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	2.13 mg in 1 mL
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	0.41 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51662-1531-1	1 in 1 CARTON	04/25/2021	
1		10 mL in 1 SYRINGE, GLASS; Type 1: Convenience Kit of Co-Package		
2	NDC:51662-1531-3	50 in 1 CARTON	04/25/2021	
2	NDC:51662-1531-2	1 in 1 POUCH		
2		1 in 1 CARTON		
2		10 mL in 1 SYRINGE, GLASS; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA209359	04/25/2021	

Labeler - HF Acquisition Co LLC, DBA HealthFirst (045657305)

Registrant - HF Acquisition Co LLC, DBA HealthFirst (045657305)

Establishment

Name	Address	ID/FEI	Business Operations
HF Acquisition Co LLC, DBA HealthFirst		045657305	relabel(51662-1531)

Revised: 1/2024

HF Acquisition Co LLC, DBA HealthFirst