

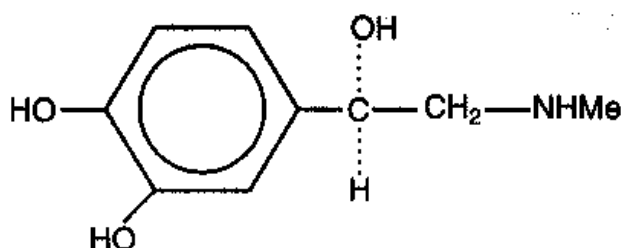
EPIPEN[®] JR**Adrenaline 150 µg/0.3 mL****EpiPen[®] Jr. 150 µg Adrenaline Auto-Injector**

Auto-Injector for Intramuscular Injection of Adrenaline for the Emergency Treatment of Anaphylactic Reactions. Delivers a single 150 microgram (µg) intramuscular dose of adrenaline from Adrenaline Injection 1:2,000 USP (0.3 mL).

Active ingredient: Adrenaline.

Chemical name: (*R*)-1-(3,4-dihydroxyphenyl)-2-methylaminoethanol

Structural formula:



Molecular formula: C₉H₁₃NO₃

Molecular weight: 183.2

CAS Registry No: 51-43-4

Description

Adrenaline is a white odourless crystalline powder, soluble in solutions of mineral acids and alkalis.

The EpiPen[®] Jr. device provides adrenaline for intramuscular auto-injection in a sterile solution prepared from adrenaline with the aid of hydrochloric acid in Pyrogen Free Water. The EpiPen[®] Jr. Auto-Injector contains 2 mL Adrenaline Injection 1:2,000 USP and is designed to deliver a single 0.3 mL dose of 150 µg. Each 0.3 mL dose contains: *Active*; 150 µg adrenaline, *Inactive*; 1.8 mg sodium chloride, 500 µg sodium metabisulfite and hydrochloric acid to adjust pH. The pH range is 2.2 – 5.0.

Adrenaline solution deteriorates rapidly on exposure to air or light, turning pink from oxidation to adrenochrome and brown from the formation of melanin. Replace the EpiPen[®] Jr. Auto-Injector if the adrenaline solution appears discoloured.

Pharmacology

Adrenaline is a sympathomimetic drug, acting on both alpha and beta receptors. Through its action on alpha adrenergic receptors, adrenaline lessens the vasodilatation and increased vascular permeability that occurs during anaphylaxis, which can lead to a loss of intravascular fluid volume and hypotension. Through its action on beta-adrenergic receptors, adrenaline causes bronchial smooth muscle relaxation that helps alleviate bronchospasm, wheezing and

dyspnoea that may occur during anaphylaxis. Other major effects are increased systolic blood pressure, reduced diastolic pressure, tachycardia, hyperglycaemia and hypokalaemia. It is a powerful cardiac stimulant. It has vasopressor properties, an antihistaminic action and is a bronchodilator.

Adrenaline also alleviates pruritus, urticaria, and angioedema and may be effective in relieving gastrointestinal and genitourinary symptoms associated with anaphylaxis because of its relaxant effects on the smooth muscle of the stomach, intestine, uterus, and urinary bladder.

The onset of action is rapid and of short duration. After intravenous infusion the half life is approximately 5 to 10 minutes. Adrenaline is rapidly distributed to the heart, spleen, several glandular tissues and adrenergic nerves. It is approximately 50% bound to plasma proteins. Adrenaline is rapidly metabolised in the liver and tissues. Up to 90% of the intravenous dose is excreted as metabolites in the urine. It crosses the placenta and is excreted in breast milk.

Indications

For the emergency treatment of anaphylaxis (acute severe allergic reactions) due to insect stings, or bites, foods, drugs or other allergens as well as idiopathic or exercise induced anaphylaxis.

Such 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, diarrhoea and abdominal cramps, involuntary voiding, wheezing, dyspnoea due to laryngeal spasm, pruritus, rashes, urticaria or angioedema.

For these reasons, auto-injectors should always be carried by such persons in situations of potential risk.

Contraindications

There are no known absolute contraindications to the use of EpiPen® Jr. during an acute, life-threatening situation..

Clinical conditions where special precautions are advised and interactions with other medicines are described in the '**PRECAUTIONS**' section.

Precautions

EpiPen® Jr. is not intended as a substitute for immediate medical care. In conjunction with the administration of adrenaline, the patient should seek immediate medical or hospital care. More than two sequential doses of adrenaline should only be administered under direct medical supervision.

The presence of anaphylactic shock should be confirmed before administering EpiPen® Jr., as EpiPen® Jr. is only indicated for the treatment of anaphylaxis.

EpiPen® Jr. Adrenaline Auto-Injector contains sodium metabisulfite, a sulfite, which may itself cause allergic-type reactions in certain susceptible persons. The alternatives to using adrenaline in a life-threatening situation may not be satisfactory. The presence of a sulfite in this product should not deter administration for serious allergic reactions even if the patient is sulfite-sensitive.

DO NOT INJECT INTRAVENOUSLY as cerebral haemorrhage may occur due to a sharp rise in blood pressure. Rapidly acting vasodilators can counteract the marked pressor effects of adrenaline if there is such inadvertent administration.

Use with caution in patients with ventricular fibrillation, cerebral arteriosclerosis, prebrillatory rhythm, tachycardia, myocardial infarction, phenothiazine-induced circulatory collapse and prostatic hypertrophy.

Adrenaline causes ECG changes including a decrease in T-wave amplitude in all leads of normal persons. Caution should be taken when administering in the presence of cardiac dilation.

Adrenaline can cause potentially fatal ventricular arrhythmias including fibrillation, especially in patients with organic heart disease or those receiving other drugs that sensitize the heart to arrhythmias (see **Interactions with Other Medicines**).

Use with caution in patients with pre-existing conditions whereby the use of vasopressor drugs is contraindicated (e.g. thyrotoxicosis).

Anginal pain may be induced by adrenaline in patients with coronary insufficiency.

Administer with caution to the elderly, and to individuals with diabetes, cardiovascular disease, hypertension, organic brain damage, narrow angle glaucoma, hyperthyroidism and psychoneurosis. In patients with Parkinsonism the drug increases rigidity and tremor.

Syncope has occurred following administration to asthmatic children.

EpiPen® Jr should not be injected into the hands, feet, ears, nose, buttocks or the genitalia as it may result in loss of blood flow to the affected area and may not provide effective treatment of anaphylaxis. Treatment should be directed at vasodilatation in addition to further treatment of anaphylaxis. If an accidental injection into one of these areas occurs, specialist medical advice must be sought immediately. Ensure the product is kept well clear of the face.

Despite these concerns, adrenaline is essential for the treatment of anaphylaxis. Therefore, patients with these conditions, and/or any other person who might be in a position to administer EpiPen® Auto-Injector to a patient experiencing anaphylaxis should be carefully instructed in regard to the circumstances under which adrenaline should be used.

Use in Pregnancy (Category A)

Adrenaline has been given to a large number of pregnant women and women of childbearing age without any proven increase in the frequency of malformations or other direct or indirect harmful effects on the foetus having been observed.

Adrenaline may delay the second stage of labour by inhibiting contractions of the uterus.

Use with caution in pregnant women whose maternal blood pressure is in excess of 130/80.

Use in Lactation

Adrenaline is excreted in breast milk.

Carcinogenesis, Mutagenesis and Impairment on Fertility

Adrenaline and other catecholamines have been shown to have mutagenic potential *in vitro* and to be an oxidative mutagen in a *WP2* bacterial reverse mutation assay. Adrenaline had a moderate degree of mutagenicity, and was positive in the DNA Repair test with *B. Subtilis* (REC) assay, but was not mutagenic in the *Salmonella* bacterial reverse mutation assay.

Studies of adrenaline after repeated exposure in animals to evaluate the carcinogenic and mutagenic potential or the effect on fertility have not been conducted. This should not prevent the use of adrenaline under the conditions noted under the '**INDICATIONS**' section.

Effects on Ability to Drive and Operate Machinery

The patients' ability to drive and use machinery may be affected by the anaphylactic reaction, as well as by possible adverse effects to adrenaline.

Interactions with Other Medicines

Central Nervous System and Other Drugs

The effects of adrenaline may be potentiated by tricyclic antidepressants, levothyroxine sodium, thyroid hormones, monoamine oxidase inhibitors and some antihistamines (eg. diphenhydramine, dexchlorpheniramine).

Other Sympathomimetic Agents

Adrenaline should not be administered with other sympathomimetic agents because of the danger of additive effects and increased toxicity.

Alpha-adrenergic Blocking Agents

Alpha-adrenergic blocking agents such as ergot alkaloids and phentolamine can reverse the pressor response to adrenaline.

Beta-adrenergic Blocking Agents

Patients taking non-selective beta-blocking drugs when administered adrenaline for the treatment of an anaphylactic reaction may experience severe hypertension and bradycardia. Propranolol inhibits the bronchodilator effect of adrenaline. The risk of cardiac arrhythmias is higher when adrenaline is given to patients receiving digoxin or quinidine.

General Anaesthetics

Halothane and other anaesthetics such as cyclopropane and trichlorethylene increase the risk of adrenaline-induced ventricular arrhythmias and acute pulmonary oedema if hypoxia is present.

Hypoglycaemic Agents

Adrenaline-induced hyperglycaemia may lead to loss of blood sugar control in diabetic patients treated with hypoglycaemic agents.

Incompatibilities

Adrenaline is physically incompatible with alkalis, metals, oxidising agents, sodium warfarin, hyaluronidase and many other drugs; it forms polymers with sodium bicarbonate.

Adverse Effects

Common symptomatic adverse events include anxiety, apprehension, restlessness, tachycardia, respiratory difficulty, tremor, weakness, dizziness, headache, dyspnoea, cold extremities, pallor, sweating, nausea, vomiting, sleeplessness, hallucinations, palpitations, fear and flushing or redness of face and skin. Psychomotor agitation, disorientation, impaired memory and psychosis may occur.

Potentially fatal ventricular arrhythmias, including ventricular fibrillation may occur and severe hypertension may lead to cerebral haemorrhage and pulmonary oedema.

The potential for adrenaline to produce these types of adverse effects does not contraindicate its use in an acute life-threatening allergic reaction.

Accidental injection into the hands, fingers or feet may result in loss of blood flow to the affected area (see **PRECAUTIONS**). Adverse events experienced as a result may include increased heart rate, local reactions including injection site pallor, coldness or hypoaesthesia or injury at the injection site resulting in bruising, bleeding, discolouration, erythema or skeletal injury.

Dosage and Administration

Selection of the appropriate dosage strength is determined according to patient body weight and this decision should be based on careful assessment of the individual patient and recognition of the life-threatening nature of reactions for which EpiPen is prescribed. A dosage of 0.01 mg/kg body weight is recommended. The EpiPen[®] Jr Auto-Injector delivers 0.15 mg adrenaline injection (0.3 mL, 1:2000) and is intended for use in patients who weigh 15 -30 kg. EpiPen, which provides a dosage of 0.3 mg, may be more appropriate for patients weighing more than 30 kg (please refer to EpiPen Product Information). The prescribing physician may choose to prescribe more or less than this amount*. With severe persistent anaphylaxis, repeat injections with an additional EpiPen[®] Auto-Injector may be necessary.

Since the dose of adrenaline delivered from an EpiPen[®] Jr. Auto-Injector is fixed, consider using other forms of injectable adrenaline if doses lower than 0.15 mg are deemed necessary.

Before using, check to make sure the solution in the Auto-Injector is not brown in colour. If it is discoloured or contains a precipitate, do not use, since these changes indicate that the effectiveness of the drug product may be decreased.

The delivered dose of the EpiPen[®] Jr. Auto-Injector should be injected intramuscularly into the anterolateral aspect of the thigh. The delivered dose is 0.3 mL of 1:2,000 USP Adrenaline Injection (150 µg), which is the usual paediatric dose for anaphylactic reactions (allergic emergencies).

DO NOT INJECT INTRAVENOUSLY.

Every effort should be made to avoid inadvertent intravascular administration (see **Overdosage**).

EpiPen[®] Jr should not be injected into the hands, feet, ears, nose, buttocks or the genitalia as it may result in loss of blood flow to the affected area and may not provide effective treatment of anaphylaxis. Treatment should be directed at vasodilatation in addition to further treatment of anaphylaxis. If an accidental injection into one of these areas occurs, specialist medical advice must be sought immediately. Ensure the product is kept well clear of the face.

To manage severe anaphylaxis, repeat EpiPen[®] Jr. injections may be necessary. Each EpiPen[®] Jr. Auto-Injector is used once only. The EpiPen[®] Jr. dose may be repeated after 15 minutes if symptoms recur or have not subsided (see **Overdosage**).

Appropriate steps should be taken to ensure that the patient thoroughly understands the indications and use of this device. The EpiPen[®] Jr. Auto-Injector should not be used for demonstration purposes. An “EpiPen[®] Jr Trainer” injector is available to assist with patient education and practice. The healthcare professional, educator or caregiver should regularly review in detail with the patient, the package leaflet provided inside the EpiPen[®] Jr. Auto-Injector carton,, which includes usage instructions for the EpiPen[®] Jr. Auto-Injector.

The EpiPen[®] Auto-Injector is intended for immediate self-administration. It is designed as emergency supportive therapy only and is not a replacement or substitute for subsequent medical or hospital care.

Patients should be instructed to dispose of the device safely after use by placing the used Auto-Injector in a sharps disposal unit.

Overdosage

Effects

Overdosage or inadvertent intravascular injection of adrenaline may cause cerebral haemorrhage resulting from a sharp rise in blood pressure. Fatalities may also result from pulmonary oedema because of peripheral vascular constriction together with cardiac stimulation.

Cardiac arrhythmias may lead to ventricular fibrillation and death.

Repeated administration of adrenaline can result in severe metabolic acidosis because of elevated blood concentration of lactic acid.

Treatment

Adrenaline is rapidly inactivated in the body and treatment of acute toxicity is mainly supportive. If necessary, the combined alpha and beta mediated effects of adrenaline may be counteracted by labetalol. Individually, alpha mediated effects may be counteracted by phentolamine whilst beta mediated effects may be counteracted by beta blocking agents.

It is advisable to contact the Poisons Information Centre on 0800 POISON (0800 764 766) for advice on the treatment and management of overdosage

Presentation and Storage Conditions

Package containing one EpiPen[®] Jr. Auto-Injector. The EpiPen[®] Jr. Auto-Injector contains 2 mL Adrenaline Injection 1:2,000 USP and delivers a single 150 µg adrenaline dose.

Adrenaline is light sensitive and should be stored in the carrier tube provided. STORE AT 25°C. TEMPERATURE EXCURSIONS TO 15°C PERMITTED. DO NOT REFRIGERATE. PROTECT FROM LIGHT. Before using, check to make sure the solution in the auto-injector is not discoloured. Replace the auto-injector if the solution is discoloured or contains a precipitate.

Medicine Classification

Restricted Medicine

Name and Address

Distributed in New Zealand by:
Mylan New Zealand Ltd
PO Box 11-183
Ellerslie
AUCKLAND
Telephone: 09-579-2792

Supplied by:
DEY[®], an affiliate of Mylan Inc, Napa, CA 94558, USA

Produced for:
DEY[®], Napa, California, 94558
by Meridian Medical Technologies, Inc.,
a subsidiary of King Pharmaceuticals Inc.,
Columbia, MD 21046, USA

EpiPen[®] is a registered trademark of Mylan Inc.

Date of Preparation

1 December 2009

*Australasian Society of Clinical Immunology and Allergy Anaphylaxis Working Party.

Guidelines for EpiPen prescription. Sydney: ASCIA, 2004.
http://www.allergy.org.au/anaphylaxis/epipen_guidelines.htm (accessed October 2009).