

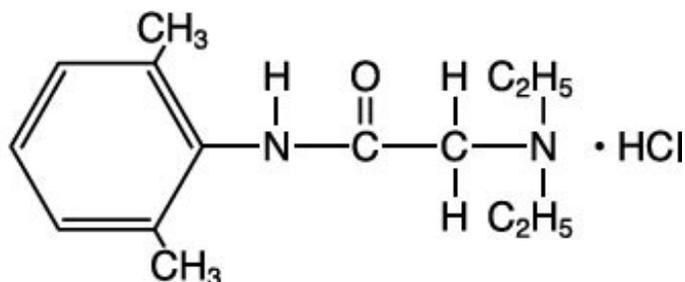
LIDOCAINE- lidocaine hydrochloride injection, solution
Fresenius Kabi USA, LLC

Lidocaine Hydrochloride Injection USP

INTRAVENOUS FOR CARDIAC ARRHYTHMIAS

DESCRIPTION

Lidocaine hydrochloride, chemical name: acetamide, 2-(diethylamino)-N-(2,6-dimethylphenyl)-, monohydrochloride has the following structural formula:



Lidocaine Hydrochloride Injection, USP intravenous for cardiac arrhythmias, is a sterile, nonpyrogenic solution prepared from lidocaine with the aid of hydrochloric acid in Water for Injection.

Each mL contains: Lidocaine HCl 20 mg; Sodium Chloride 6 mg; Water for Injection q.s. Hydrochloric acid and/or sodium hydroxide may have been added for pH adjustment (pH 5.0 to 7.0). The container is for single use; solution contains no preservative.

CLINICAL PHARMACOLOGY

Lidocaine hydrochloride exerts its antiarrhythmic effect by raising the electrical stimulation threshold of the ventricle during diastole. In usual therapeutic doses, lidocaine hydrochloride produces no change in myocardial contractility, systemic arterial pressure or in absolute refractory period.

Onset of action is rapid after an intravenous bolus with a duration of action of approximately 15 to 20 minutes.

Approximately 90% of an administered dose of lidocaine hydrochloride is metabolized in the liver. The remainder (10%) of the drug is excreted unchanged in the urine. Pharmacokinetic data indicate reduced elimination of lidocaine after prolonged (24 hours) infusion with resultant prolongation of the half-life to approximately three times that seen following a single administration.

INDICATIONS AND USAGE

Lidocaine Hydrochloride Injection, USP administered intravenously is specifically indicated in the acute management of 1) ventricular arrhythmias occurring during cardiac manipulation such as cardiac surgery, and 2) life-threatening arrhythmias, particularly those which are ventricular in origin, such as those which occur during acute myocardial

infarction.

CONTRAINDICATIONS

Lidocaine Hydrochloride Injection, USP is contraindicated in patients with a known history of hypersensitivity to local anesthetics of the amide type. Lidocaine Hydrochloride Injection, USP should not be used in patients with STOKES-ADAMS syndrome, Wolff-Parkinson-White syndrome, or with severe degrees of sinoatrial, atrioventricular or intraventricular block.

WARNINGS

Constant monitoring of the electrocardiogram and blood pressure is essential in the proper administration of Lidocaine Hydrochloride Injection, USP intravenously. If hypotension or excessive depression of cardiac conductivity (such as prolongation of the PR interval and QRS complex and the appearance of aggravation of arrhythmias) is seen, administration of Lidocaine Hydrochloride Injection, USP should be discontinued.

Emergency resuscitative equipment and drugs must be immediately available to manage possible adverse reactions involving the cardiovascular, respiratory, or central nervous system.

Occasional acceleration of ventricular rate may occur when Lidocaine Hydrochloride Injection, USP is administered to patients with atrial fibrillation.

Toxicity may be manifest as central nervous system depression (sedation) or irritability (twitching), which may progress to frank convulsions accompanied by respiratory depression and/or arrest. Early recognition of premonitory signs, assurance of adequate oxygenation and (where necessary) establishment of artificial airway with ventilatory support are essential to management of the problem. Should convulsions persist despite ventilatory support, *small doses* of anticonvulsant drugs may be used intravenously. Examples of such drugs include benzodiazepines (e.g., diazepam, lorazepam), or an ultrashort acting barbiturate (e.g., thiopental or thiamylal). If the patient is under anesthesia, a short-acting muscle relaxant (succinylcholine) may be used. Longer acting drugs should be used only when recurrent convulsions are evidenced.

PRECAUTIONS

The safe use of Lidocaine Hydrochloride Injection, USP requires careful ECG observation in an environment equipped and by persons trained for resuscitation.

General

Caution should be employed in the repeated use of Lidocaine Hydrochloride Injection, USP in patients with severe liver or renal disease because accumulation may occur and lead to toxic phenomena since Lidocaine Hydrochloride Injection, USP is metabolized mainly in the liver and excreted by the kidneys. This drug should also be used with caution in patients with hypovolemia and shock and all forms of heart block (see **CONTRAINDICATIONS** and **WARNINGS**).

In patients with sinus bradycardia or incomplete heart block, the administration of

Lidocaine Hydrochloride Injection, USP intravenously or the elimination of ventricular ectopic beats without prior acceleration in heart rate may promote more frequent and serious ventricular arrhythmias or complete heart block. (See **CONTRAINDICATIONS**.)

Many potent anesthetic drugs, neuromuscular blocking agents and possibly amide local anesthetics may serve as triggering agents for the fulminant hypermetabolic process termed malignant hyperthermia. Key to successful outcome of fulminant hypermetabolism is early recognition of premonitory signs, i.e., unexplained or unexpected tachycardia and increased metabolic rate as evidenced by respiratory and/or metabolic acidosis. Treatment includes administration of oxygen and discontinuation of lidocaine hydrochloride administration and where necessary administration of dantrolene sodium. For additional information on management, see prescribing information for dantrolene sodium.

Drug Interaction

The effect of Lidocaine Hydrochloride Injection, USP for the management of cardiac arrhythmias is potentiated with the simultaneous or prior administration of propranolol.

Lidocaine Hydrochloride Injection, USP should be used with caution in patients with digitalis toxicity accompanied by atrioventricular block. (See **CONTRAINDICATIONS**.)

Drug interaction was studied with several beta-blockers (propranolol, metoprolol, nadolol, and pindolol). Pretreatment with propranolol, metoprolol and nadolol increased the serum concentration of lidocaine. Pretreatment with pindolol did not affect the serum concentration of lidocaine.

Carcinogenesis, Mutagenesis, Impairment of Fertility

Studies of Lidocaine Hydrochloride Injection, USP in animals to evaluate the carcinogenic and mutagenic potential or the effect on fertility have not been conducted.

Pregnancy Category B

Reproductive studies have been performed in the rat at doses up to 5 times the human dose and have revealed no evidence of impaired fertility or harm to the fetus due to Lidocaine Hydrochloride Injection, USP. There are, however, no adequate and well controlled studies in pregnant women. Because animal reproduction studies are not always predictive of human response, this drug should be used in pregnancy only if clearly needed.

Labor and Delivery

The effects of Lidocaine Hydrochloride Injection, USP solution on the mother and the fetus, when used in the management of cardiac arrhythmias during labor and delivery, are not known. Lidocaine Hydrochloride Injection, USP readily crosses the placental barrier.

Nursing Mothers

It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when Lidocaine Hydrochloride Injection, USP is administered to a nursing woman.

Pediatric Use

Safety and effectiveness in children have not been established by controlled clinical studies. (See **DOSAGE AND ADMINISTRATION.**)

ADVERSE REACTIONS

To report SUSPECTED ADVERSE REACTIONS, contact Fresenius Kabi USA, LLC at 1-800-551-7176 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

Most adverse reactions accompanying administration of Lidocaine Hydrochloride Injection, USP result from systemic toxicity affecting primarily the central nervous system and the cardiovascular system.

Systemic reactions of the following types have been reported:

1. Central Nervous System

Light-headedness; drowsiness; dizziness; apprehension; euphoria; tinnitus; blurred vision or double vision; vomiting; sensations of heat; cold or numbness; twitching; tremors; convulsions; unconsciousness; respiratory depression and arrest.

2. Cardiovascular System

Hypotension; cardiovascular collapse; and bradycardia which may lead to cardiac arrest. Continuous monitoring of blood pressure and the electrocardiogram are essential to prevent these events.

3. Allergic reactions may occur but are infrequent. There have been no reports of cross-sensitivity between lidocaine hydrochloride and procainamide or between lidocaine and quinidine.

OVERDOSAGE

Management of Adverse Reactions—

1. In the case of severe reaction, discontinue the use of Lidocaine Hydrochloride Injection, USP.
2. Institute emergency resuscitative procedures and administer the emergency drugs necessary to manage the severe reaction. For severe convulsions, small increments of diazepam or an ultra short-acting barbiturate (thiopental or thiamylal); or if those are not available a short-acting barbiturate (pentobarbital or secobarbital); or if the patient is under anesthesia, a short-acting muscle relaxant (succinylcholine) may be given intravenously. Muscle relaxants and intravenous medications should only be used by those familiar with their use. Patency of the airway and adequacy of ventilation must be assured. (See **WARNINGS.**)
3. Should circulatory depression occur, vasopressors, such as ephedrine, or metaraminol may be used. Should cardiac arrest occur, immediate initiation of standard CPR procedures should commence.

DOSAGE AND ADMINISTRATION

Adult

For Direct Injection—The usual dose is 50 to 100 mg administered intravenously under ECG monitoring. This dose may be administered at the rate of approximately 25 to 50 mg/min. Sufficient time should be allowed to enable a slow circulation to carry the drug to the site of action. If the initial injection of 50 to 100 mg does not produce a desired response, a second dose may be repeated after 5 minutes.

NO MORE THAN 200 TO 300 MG OF LIDOCAINE HYDROCHLORIDE INJECTION, USP SHOULD BE ADMINISTERED DURING A ONE HOUR PERIOD.

This product is for **DIRECT INJECTION ONLY**. For continuous infusion protocol, see prescribing information for lidocaine hydrochloride, USP for infusion solution.

Pediatric

Although controlled clinical studies to establish pediatric dosing schedules have not been conducted, the American Heart Association's *Standards and Guidelines* recommends a bolus dose of 1 mg/kg followed by an infusion rate of 30 mcg/kg/min.

This product is for **DIRECT INJECTION ONLY**. For continuous infusion protocol, see prescribing information for lidocaine hydrochloride, USP for infusion solution.

Sterilization, Storage and Technical Procedures

It is recommended that chemical disinfection be accomplished by wiping the vial thoroughly with cotton or gauze that has been moistened with either 91% isopropyl alcohol or 70% ethyl alcohol, just prior to use. Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature].

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit.

HOW SUPPLIED:

For Direct Injection Only

Lidocaine Hydrochloride Injection, USP

Product Code	Unit of Sale	Strength	Each
20805	NDC 63323-208-05 Unit of 25	2% 100 mg per 5 mL (20 mg per mL)	NDC 63323-208-01 5 mL Single Dose Vial

Preservative Free

Discard Unused Portion.

Use only if solution is clear and seal intact.



**FRESENIUS
KABI**

Lake Zurich, IL 60047

www.fresenius-kabi.com/us

45775D

Revised: May 2022

PACKAGE LABEL - PRINCIPAL DISPLAY - Lidocaine HCl 5 mL Single Dose Vial Label

NDC 63323-208-01 20805

LIDOCAINE HCl

INJECTION, USP

2%

100 mg per 5 mL

(20 mg per mL)

Intravenous For

Cardiac Arrhythmias

Rx only

5 mL Single Dose Vial

NDC 63323-208-01 20805

LIDOCAINE HCl
INJECTION, USP

2% **100 mg per 5 mL**
(20 mg per mL)

Intravenous For
Cardiac Arrhythmias

Rx only

5 mL Single Dose Vial

Sterile, Nonpyrogenic
FOR DIRECT INJECTION
Preservative Free

Discard unused portion.

Each mL contains: Lidocaine HCl
20 mg; sodium chloride 6 mg; Water
for injection q.s. HCl and/or NaOH may
have been added for pH adjustment.

Usual Dosage: See insert.
Use only if solution is clear and seal intact.
Store at 20° to 25°C (68° to 77°F)
[see USP Controlled Room Temperature].

Fresenius Kabi
Lake Zurich, IL 60047
www.fresenius-kabi.com/us

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LOT/EXP

PACKAGE LABEL - PRINCIPAL DISPLAY - Lidocaine HCl 5 mL Vial Tray Label

NDC 63323-208-05 20805

LIDOCAINE HCl

INJECTION, USP

2%

100 mg per 5 mL

(20 mg per mL)

Intravenous For
Cardiac Arrhythmias
Rx only

25 Single Dose Vials, 5 mL

NDC 63323-208-05 20805

LIDOCAINE HCl
INJECTION, USP

2% **100 mg per 5 mL**
(20 mg per mL)

Intravenous For
Cardiac Arrhythmias
Rx only

25 Single Dose Vials, 5 mL

Sterile, Nonpyrogenic

FOR DIRECT INJECTION

Preservative Free

Discard unused portion.

Each mL contains: Lidocaine HCl 20 mg;
sodium chloride 6 mg; Water for
Injection q.s. HCl and/or NaOH may have
been added for pH adjustment.

Usual Dosage: See insert.

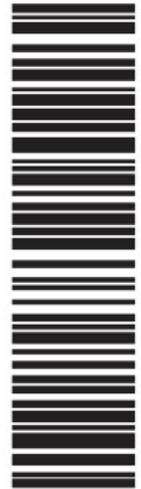
Use only if solution is clear and seal
intact.

Store at 20° to 25°C (68° to 77°F)
[see USP Controlled Room Temperature].



Lake Zurich, IL 60047
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(01)2036333208053

LIDOCAINE

lidocaine hydrochloride injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:63323-208
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	6 mg in 1 mL
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63323-208-05	25 in 1 TRAY	09/05/2000	
1	NDC:63323-208-01	5 mL in 1 VIAL, SINGLE-DOSE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA017584	09/05/2000	

Labeler - Fresenius Kabi USA, LLC (608775388)

Establishment			
Name	Address	ID/FEI	Business Operations
Fresenius Kabi USA, LLC		840771732	ANALYSIS(63323-208) , MANUFACTURE(63323-208)

Establishment			
Name	Address	ID/FEI	Business Operations
Fresenius Kabi USA, LLC		023648251	ANALYSIS(63323-208) , MANUFACTURE(63323-208)

Revised: 2/2025

Fresenius Kabi USA, LLC