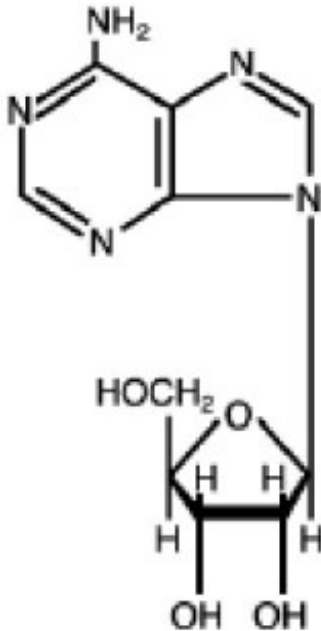


**ADENOSINE- adenosine injection, solution**  
**HF Acquisition Co LLC, DBA HealthFirst**

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Rx only

FOR RAPID BOLUS INTRAVENOUS USE

Adenosine is an endogenous nucleoside occurring in all cells of the body. It is chemically 6-amino-9-β-D-ribofuranosyl-9-H-purine and has the following structural formula:



C<sub>10</sub>H<sub>13</sub>N<sub>5</sub>O<sub>4</sub> M.W. 267.24

Adenosine is a white crystalline powder. It is soluble in water and practically insoluble in alcohol. Solubility increases by warming and lowering the pH. Adenosine is not chemically related to other antiarrhythmic drugs. Adenosine injection, USP is a sterile, nonpyrogenic solution for rapid bolus intravenous injection. Each mL contains 3 mg adenosine and 9 mg sodium chloride in water for injection. The pH of the solution is between 4.5 and 7.5.

The TopPac® plastic syringe is molded from a specially formulated cyclo olefin copolymer. Water permeates from inside the container at an extremely slow rate which will have an insignificant effect on solution concentration over the expected shelf life.

Solutions in contact with the plastic container may leach out certain chemical components from the plastic in very small amounts which have been toxicologically assessed and are within the safety margins.

#### Mechanism of Action

Adenosine injection, USP slows conduction time through the A-V node, can interrupt the reentry pathways through the A-V node, and can restore normal sinus rhythm in patients with paroxysmal supraventricular tachycardia (PSVT), including PSVT associated with Wolff-Parkinson-White Syndrome.

Adenosine injection, USP is antagonized competitively by methylxanthines such as caffeine and theophylline, and potentiated by blockers of nucleoside transport such as

dipyridamole. Adenosine injection, USP is not blocked by atropine.

## Hemodynamics

The intravenous bolus dose of 6 mg or 12 mg adenosine injection, USP usually has no systemic hemodynamic effects. When larger doses are given by infusion, adenosine decreases blood pressure by decreasing peripheral resistance.

## Pharmacokinetics

Intravenously administered adenosine is rapidly cleared from the circulation via cellular uptake, primarily by erythrocytes and vascular endothelial cells. This process involves a specific transmembrane nucleoside carrier system that is reversible, nonconcentrative, and bidirectionally symmetrical. Intracellular adenosine is rapidly metabolized either via phosphorylation to adenosine monophosphate by adenosine kinase, or via deamination to inosine by adenosine deaminase in the cytosol. Since adenosine kinase has a lower  $K_m$  and  $V_{max}$  than adenosine deaminase, deamination plays a significant role only when cytosolic adenosine saturates the phosphorylation pathway. Inosine formed by deamination of adenosine can leave the cell intact or can be degraded to hypoxanthine, xanthine, and ultimately uric acid. Adenosine monophosphate formed by phosphorylation of adenosine is incorporated into the high-energy phosphate pool. While extracellular adenosine is primarily cleared by cellular uptake with a half-life of less than 10 seconds in whole blood, excessive amounts may be deaminated by an ecto-form of adenosine deaminase. As adenosine injection, USP requires no hepatic or renal function for its activation or inactivation, hepatic and renal failure would not be expected to alter its effectiveness or tolerability.

## Clinical Trial Results

In controlled studies in the United States, bolus doses of 3 mg, 6 mg, 9 mg, and 12 mg were studied. A cumulative 60% of patients with paroxysmal supraventricular tachycardia had converted to normal sinus rhythm within one minute after an intravenous bolus dose of 6 mg adenosine injection, USP (some converted on 3 mg and failures were given 6 mg), and a cumulative 92% converted after a bolus dose of 12 mg. Seven to sixteen percent of patients converted after 1 to 4 placebo bolus injections. Similar responses were seen in a variety of patient subsets, including those using or not using digoxin, those with Wolff-Parkinson-White Syndrome, males, females, blacks, Caucasians, and Hispanics.

Adenosine is not effective in converting rhythms other than PSVT, such as atrial flutter, atrial fibrillation, or ventricular tachycardia, to normal sinus rhythm.

Intravenous adenosine injection, USP is indicated for the following:

Conversion to sinus rhythm of paroxysmal supraventricular tachycardia (PSVT), including that associated with accessory bypass tracts (Wolff-Parkinson-White Syndrome). When clinically advisable, appropriate vagal maneuvers (e.g., Valsalva maneuver), should be attempted prior to adenosine injection, USP administration.

It is important to be sure the adenosine injection, USP solution actually reaches the systemic circulation (see DOSAGE AND ADMINISTRATION).

Adenosine injection, USP does not convert atrial flutter, atrial fibrillation, or ventricular tachycardia to normal sinus rhythm. In the presence of atrial flutter or atrial fibrillation, a transient modest slowing of ventricular response may occur immediately following

adenosine injection, USP administration.

Intravenous adenosine injection, USP is contraindicated in:

Second- or third-degree A-V block (except in patients with a functioning artificial pacemaker).

Sinus node disease, such as sick sinus syndrome or symptomatic bradycardia (except in patients with a functioning artificial pacemaker).

Known hypersensitivity to adenosine.

#### Heart Block

Adenosine injection, USP exerts its effect by decreasing conduction through the A-V node and may produce a short lasting first-, second- or third-degree heart block. Appropriate therapy should be instituted as needed. Patients who develop high-level block on one dose of adenosine injection, USP should not be given additional doses. Because of the very short half-life of adenosine, these effects are generally self-limiting. Appropriate resuscitative measures should be available.

Transient or prolonged episodes of asystole have been reported with fatal outcomes in some cases. Rarely, ventricular fibrillation has been reported following adenosine injection, USP administration, including both resuscitated and fatal events. In most instances, these cases were associated with the concomitant use of digoxin and, less frequently with digoxin and verapamil. Although no causal relationship or drug-drug interaction has been established, adenosine injection, USP should be used with caution in patients receiving digoxin or digoxin and verapamil in combination.

#### Arrhythmias at Time of Conversion

At the time of conversion to normal sinus rhythm, a variety of new rhythms may appear on the electrocardiogram. They generally last only a few seconds without intervention, and may take the form of premature ventricular contractions, atrial premature contractions, atrial fibrillation, sinus bradycardia, sinus tachycardia, skipped beats, and varying degrees of A-V nodal block. Such findings were seen in 55% of patients.

#### Bronchoconstriction

Adenosine injection, USP is a respiratory stimulant (probably through activation of carotid body chemoreceptors) and intravenous administration in man has been shown to increase minute ventilation ( $V_e$ ) and reduce arterial  $PCO_2$  causing respiratory alkalosis.

Adenosine administered by inhalation has been reported to cause bronchoconstriction in asthmatic patients, presumably due to mast cell degranulation and histamine release. These effects have not been observed in normal subjects. Adenosine injection, USP has been administered to a limited number of patients with asthma and mild to moderate exacerbation of their symptoms has been reported. Respiratory compromise has occurred during adenosine infusion in patients with obstructive pulmonary disease. Adenosine injection, USP should be used with caution in patients with obstructive lung disease not associated with bronchoconstriction (e.g., emphysema, bronchitis, etc.) and should be avoided in patients with bronchoconstriction or bronchospasm (e.g., asthma). Adenosine injection, USP should be discontinued in any patient who develops severe respiratory difficulties.

#### Drug Interactions

Intravenous adenosine injection, USP has been effectively administered in the presence of other cardioactive drugs, such as quinidine, beta-adrenergic blocking agents, calcium channel blocking agents, and angiotensin converting enzyme inhibitors, without any change in the adverse reaction profile. Digoxin and verapamil use may be rarely associated with ventricular fibrillation when combined with adenosine injection, USP (see WARNINGS). Because of the potential for additive or synergistic depressant effects on the SA and AV nodes, however, adenosine injection, USP should be used with caution in the presence of these agents. The use of adenosine injection, USP in patients receiving digitalis may be rarely associated with ventricular fibrillation (see WARNINGS).

The effects of adenosine are antagonized by methylxanthines such as caffeine and theophylline. In the presence of these methylxanthines, larger doses of adenosine may be required or adenosine may not be effective. Adenosine effects are potentiated by dipyridamole. Thus, smaller doses of adenosine may be effective in the presence of dipyridamole. Carbamazepine has been reported to increase the degree of heart block produced by other agents. As the primary effect of adenosine is to decrease conduction through the A-V node, higher degrees of heart block may be produced in the presence of carbamazepine.

#### Carcinogenesis, Mutagenesis, Impairment of Fertility

Studies in animals have not been performed to evaluate the carcinogenic potential of adenosine injection, USP. Adenosine was negative for genotoxic potential in the Salmonella (Ames Test) and Mammalian Microsome Assay.

Adenosine, however, like other nucleosides at millimolar concentrations present for several doubling times of cells in culture, is known to produce a variety of chromosomal alterations. Fertility studies in animals have not been conducted with adenosine.

#### Pregnancy Category C

Animal reproduction studies have not been conducted with adenosine; nor have studies been performed in pregnant women. As adenosine is a naturally occurring material, widely dispersed throughout the body, no fetal effects would be anticipated. However, since it is not known whether adenosine injection, USP can cause fetal harm when administered to pregnant women, adenosine injection, USP should be used during pregnancy only if clearly needed.

#### Pediatric Use

No controlled studies have been conducted in pediatric patients to establish the safety and efficacy of adenosine injection, USP for the conversion of paroxysmal supraventricular tachycardia (PSVT). However, intravenous adenosine has been used for the treatment of PSVT in neonates, infants, children and adolescents (see DOSAGE AND ADMINISTRATION).

#### Geriatric Use

Clinical studies of adenosine injection, USP 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 elderly and younger patients. In general, adenosine injection, USP in geriatric patients should be used with caution since this population may have a diminished cardiac function, nodal dysfunction, concomitant diseases or drug therapy that may alter hemodynamic function and produce severe bradycardia or AV block.

The following reactions were reported with intravenous adenosine injection, USP used in controlled U.S. clinical trials. The placebo group had a less than 1% rate of all of these reactions.

#### Cardiovascular

Facial flushing (18%), headache (2%), sweating, palpitations, chest pain, hypotension (less than 1%).

#### Respiratory

Shortness of breath/dyspnea (12%), chest pressure (7%), hyperventilation, head pressure (less than 1%).

#### Central Nervous System

Lightheadedness (2%), dizziness, tingling in arms, numbness (1%), apprehension, blurred vision, burning sensation, heaviness in arms, neck and back pain (less than 1%).

#### Gastrointestinal

Nausea (3%), metallic taste, tightness in throat, pressure in groin (less than 1%).

#### Post-Marketing Experience

(see WARNINGS)

The following adverse events have been reported from marketing experience with adenosine injection, USP. Because these events are reported voluntarily from a population of uncertain size, are associated with concomitant diseases and multiple drug therapies and surgical procedures, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure. Decisions to include these events in labeling are typically based on one or more of the following factors: (1) seriousness of the event, (2) frequency of the reporting, (3) strength of causal connection to the drug, or a combination of these factors.

#### Cardiovascular

Prolonged asystole, ventricular tachycardia, ventricular fibrillation, transient increase in blood pressure, bradycardia, atrial fibrillation, and Torsades de Pointes.

#### Respiratory

Bronchospasm.

#### Central Nervous System

Seizure activity, including tonic clonic (grand mal) seizures, and loss of consciousness.

The half-life of adenosine injection, USP is less than 10 seconds. Thus, adverse effects are generally rapidly self-limiting. Treatment of any prolonged adverse effects should be individualized and be directed toward the specific effect. Methylxanthines, such as caffeine and theophylline, are competitive antagonists of adenosine.

For rapid bolus intravenous use only.

Adenosine injection, USP should be given as a rapid bolus by the peripheral intravenous route. To be certain the solution reaches the systemic circulation, it should be administered either directly into a vein or, if given into an intravenous line, it should be

given as close to the patient as possible and followed by a rapid saline flush.

#### Adult Patients

The dose recommendation is based on clinical studies with peripheral venous bolus dosing. Central venous (CVP or other) administration of adenosine injection, USP has not been systematically studied.

The recommended intravenous doses for adults are as follows:

Initial dose: 6 mg given as a rapid intravenous bolus (administered over a 1 to 2 second period).

Repeat administration: If the first dose does not result in elimination of the supraventricular tachycardia within 1 to 2 minutes, 12 mg should be given as a rapid intravenous bolus.

This 12 mg dose may be repeated a second time if required.

#### Pediatric Patients

The dosages used in neonates, infants, children and adolescents were equivalent to those administered to adults on a weight basis.

Pediatric Patients with a Body Weight less than 50 kg:

Initial dose: Give 0.05 mg/kg to 0.1 mg/kg as a rapid intravenous bolus given either centrally or peripherally. A saline flush should follow.

Repeat administration: If conversion of paroxysmal supraventricular tachycardia does not occur within 1 to 2 minutes, additional bolus injections of adenosine can be administered at incrementally higher doses, increasing the amount given by 0.05 mg/kg to 0.1 mg/kg. Follow each bolus with a saline flush. This process should continue until sinus rhythm is established or a maximum single dose of 0.3 mg/kg is used.

Pediatric Patients with a Body Weight greater than or equal to 50 kg: Administer the adult dose.

Doses greater than 12 mg are not recommended for adult and pediatric patients.

NOTE: Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration.

Adenosine injection, USP is supplied as a sterile nonpyrogenic solution in normal saline, as follows:

Unit of Sale	Strength	Each
NDC 51662-1643-1	12 mg per 4 mL 4 mL fill, in a 5 mL(3 mg per mL) plastic prefilled syringe.	Syringe
NDC 51662-1643-2	12 mg per 4 mL 4 mL fill, in a 5 mL(3 mg per mL) plastic prefilled syringe.	Pouch
NDC 51662-1643-3	12 mg per 4 mL 4 mL fill, in a 5 mL(3 mg per mL) plastic prefilled syringe.	Case of 10 Syringe

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

DO NOT REFRIGERATE as crystallization may occur. If crystallization has occurred, dissolve crystals by warming to room temperature. The solution must be clear at the time of use.

Contains no preservatives. Discard unused portion.

The container closure is not made with natural rubber latex.

May require needle or blunt. To prevent needle-stick injuries, needles should not be recapped, purposely bent or broken by hand.

Paul T, Pfammatter. J-P. Adenosine: an effective and safe antiarrhythmic drug in pediatrics. Pediatric Cardiology 1997; 18:118-126.

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## **NDC 51662-1643-1**

(01)00351662164310

(17)230919

(21)351662141212

(10)789210

SEE MANUFACTURE'S INSERT

DISTRIBUTED BY HF ACQUISITIONS CO., LLC

MUKILTEO, WA 98275

RX ONLY



**NDC 51662-1643-2**



ADENOSINE INJECTION, USP 12mg PER 4mL  
(3mg PER mL) 4mL SYR

NDC: 51662-1643-2

LOT: 789210

EXP: 2023-09-19

(01) 00351662164327

(10) 789210

(17) 230919

(21) 351662141212

FOR RAPID BOLUS INTRAVENOUS USE. PRESERVATIVE FREE. MEDICATION AND FLUID PATH ARE STERILE AND NON-PYROGENIC IF PROTECTIVE COVER IS UNDISTURBED AND PACKAGE IS INTACT. SYRINGES MAY REQUIRE NEEDLE OR BLUNT. EACH mL CONTAINS: ADENOSINE 3 mg AND SODIUM CHLORIDE 9 mg IN WATER FOR INJECTION. USUAL DOSAGE: SEE PACKAGE INSERT. STORE AT 20 TO 25C (68 to 77F) [SEE USP CONTROLLED ROOM TEMPERATURE]. DO NOT REFRIGERATE AS CRYSTALLIZATION MAY OCCUR. IF CRYSTALLIZATION HAS OCCURRED, DISSOLVE CRYSTALS BY WARMING TO ROOM TEMPERATURE. THE SOLUTION MUST BE CLEAR AT THE TIME OF USE. DISCARD UNUSED PORTION. THE CONTAINER CLOSURE IS NOT MADE WITH NATURAL RUBBER LATEX.

See manufacturer's package insert

ORIGINAL MFG NDC: 63323-651-90

RX ONLY

Manufactured by HF Acquisition Co., LLC

Mukilteo, WA 98275



**NDC 51662-1643-3**

ADENOSINE INJECTION, USP 12mg PER 4mL  
(3mg PER mL) 4mL SYR (CASE OF 10)

NDC: 51662-1643-3

LOT: 789210

EXP: 2023-09-19

CASE

(01) 40351662164332

(10) 789210

(17) 230919

(21) 351662141212

FOR RAPID BOLUS INTRAVENOUS USE. PRESERVATIVE FREE. MEDICATION AND FLUID PATH ARE STERILE AND NON-PYROGENIC IF PROTECTIVE COVER IS UNDISTURBED AND PACKAGE IS INTACT. SYRINGES MAY REQUIRE NEEDLE OR BLUNT. EACH mL CONTAINS: ADENOSINE 3 mg AND SODIUM CHLORIDE 9 mg IN WATER FOR INJECTION. USUAL DOSAGE: SEE PACKAGE INSERT. STORE AT 20 TO 25C (68 to 77F) [SEE USP CONTROLLED ROOM TEMPERATURE]. DO NOT REFRIGERATE AS CRYSTALLIZATION MAY OCCUR. IF CRYSTALLIZATION HAS OCCURRED, DISSOLVE CRYSTALS BY WARMING TO ROOM TEMPERATURE. THE SOLUTION MUST BE CLEAR AT THE TIME OF USE. DISCARD UNUSED PORTION. THE CONTAINER CLOSURE IS NOT MADE WITH NATURAL RUBBER LATEX.

See manufacturer's package insert

ORIGINAL MFG NDC: 63323-651-90

RX ONLY

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Mukilteo, WA 98275



(01)40351662164332  
(17)230919  
(21)351662141212  
(10)789210

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MUKILTEO, WA 98275

RX ONLY



## ADENOSINE

adenosine injection, solution

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:51662-1643(NDC:63323- 651)
<b>Route of Administration</b>	INTRAVENOUS		

### Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>ADENOSINE</b> (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	ADENOSINE	3 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51662-1643-3	10 in 1 CASE	04/16/2018	
1	NDC:51662-1643-2	1 in 1 POUCH		
1	NDC:51662-1643-1	2 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	ANDA205568	04/16/2018	

**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(51662-1643)

Revised: 5/2024

HF Acquisition Co LLC, DBA HealthFirst