SODIUM CHLORIDE- sodium chloride injection, solution
Cardinal Health 107, LLC

0.9% Sodium Chloride

Injection, USP

Carpuject™ with Luer Lock
Fliptop Plastic Vial
LifeShield® Fliptop Plastic Vial

Preservative-Free

Rx only

DESCRIPTION

This preparation is designed solely for parenteral use only after addition of drugs that require dilution or must be dissolved in an aqueous vehicle prior to injection.

0.9% Sodium Chloride Injection, USP is a sterile, nonpyrogenic, isotonic solution of sodium chloride and water for injection. Each mL contains sodium chloride 9 mg. It contains no bacteriostat, antimicrobial agent or added buffer and is supplied only in single-dose containers to dilute or dissolve drugs for injection. 0.308 mOsmol/mL (calc.). 0.9% Sodium Chloride Injection, USP contains no preservatives. The solution may contain hydrochloric acid and/or sodium hydroxide for pH adjustment. pH 5.3 (4.5 to 7.0).

Sodium Chloride, USP is chemically designated NaCl, a white crystalline compound freely soluble in water.

The glass container is a Type I borosilicate glass and meets the requirements of the powdered glass test according to the USP standards.

The semi-rigid vial is fabricated from a specially formulated polyolefin. It is a copolymer of ethylene and propylene. The safety of the plastic has been confirmed by tests in animals according to USP biological standards for plastic containers. The container requires no vapor barrier to maintain the proper drug concentration.

CLINICAL PHARMACOLOGY

Sodium chloride in water dissociates to provide sodium (Na^+) and chloride (Cl^-) ions. These ions are normal constituents of the body fluids (principally extracellular) and are essential for maintaining electrolyte balance.

The distribution and excretion of sodium (Na^+) and chloride (Cl^-) are largely under the control of the kidney which maintains a balance between intake and output.

The small volume of fluid and amount of sodium chloride provided by 0.9% Sodium Chloride Injection, USP when used only as an isotonic vehicle for parenteral injection of drugs, is unlikely to exert a significant effect on fluid and electrolyte balance except possibly in neonates and very small infants.

Water is an essential constituent of all body tissues and accounts for approximately 70% of total body weight. Average normal adult daily requirement ranges from two to three liters (1.0 to 1.5 liters each for insensible water loss by perspiration and urine production).

Water balance is maintained by various regulatory mechanisms. Water distribution depends primarily on the concentration of electrolytes in the body compartments and sodium (Na^+) plays a major role in maintaining physiologic equilibrium.

INDICATIONS AND USAGE

This parenteral preparation is indicated only for diluting or dissolving drugs for intravenous, intramuscular or subcutaneous injection, according to instructions of the manufacturer of the drug to be administered.

PRECAUTIONS

Consult the manufacturer's instructions for choice of vehicle, appropriate dilution or volume for dissolving the drugs to be injected, including the route and rate of injection.

Inspect reconstituted (diluted or dissolved) drugs for clarity (if soluble) and freedom from unexpected precipitation or discoloration prior to administration.

Pregnancy: Animal reproduction studies have not been conducted with 0.9% Sodium Chloride Injection, USP. It is also not known whether sodium chloride injection containing additives can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. Sodium chloride injection containing additives should be given to a pregnant woman only if clearly needed.

Pediatric Use: The safety and effectiveness in the pediatric population are based on the similarity of the clinical conditions of the pediatric and adult populations. In neonates or very small infants the volume of fluid may affect fluid and electrolyte balance.

Drug Interactions

Some drugs for injection may be incompatible in a given vehicle, or when combined in the same vehicle or in a vehicle containing benzyl alcohol. Consult with pharmacist, if available.

Use aseptic technique for single or multiple entry and withdrawal from all containers.

When diluting or dissolving drugs, mix thoroughly and use promptly.

Do not store reconstituted solutions of drugs for injection unless otherwise directed by the manufacturer of the solute.

Do not use unless the solution is clear and seal intact. Do not reuse single-dose containers, discard unused portion.

ADVERSE REACTIONS

Reactions which may occur because of this solution, added drugs or the technique of reconstitution or administration include febrile response, local tenderness, abscess, tissue necrosis or infection at the site of injection, venous thrombosis or phlebitis extending from the site of injection and extravasation.

If an adverse reaction does occur, discontinue the infusion, evaluate the patient, institute appropriate countermeasures, and if possible, retrieve and save the remainder of the unused vehicle for examination.

OVERDOSAGE

Use only as a diluent or solvent. This parenteral preparation is unlikely to pose a threat of carbohydrate, sodium chloride or fluid overload except possibly in neonates or very small infants. In the event these should occur, re-evaluate the patient and institute appropriate corrective measures. See **PRECAUTIONS** and **ADVERSE REACTIONS**.

DOSAGE AND ADMINISTRATION

The volume of the preparation to be used for diluting or dissolving any drug for injection, is dependent on the vehicle concentration, dose and route of administration as recommended by the manufacturer. This parenteral should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit.

HOW SUPPLIED

0.9% Sodium Chloride Injection, USP is supplied in the following:

Unit of Sale	Concentration
Overbagged with 5 x 10 mL Single-dose Plastic Fliptop Vials in each bag, NDC 55154-0126-5	0.9% (10 mL)

WARNING: This Unit Dose package is not child resistant and is Intended for Institutional Use Only. Keep this and all drugs out of the reach of children.

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

Instructions for Use of the Syringe Systems

Instructions for using the Carpuject Syringe are available with the reusable Carpuject Holder, List 2049-02.

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

Distributed By:

Cardinal Health

Dublin, OH 43017

L28642700324

LAB-1097-3.0

Revised: 08/2021

Package/Label Display Panel

NDC 55154-0126-5

0.9% SODIUM CHLORIDE

INJECTION, USP

5 x 10 mL SINGLE-DOSE VIALS

T29

NDC 55154-0126-5

0.9% SODIUM CHLORIDE
INJECTION, USP

5 x 10 mL SINGLE-DOSE VIALS

FOR USE AS A STERILE DILUENT

Each mL contains sodium chloride, 9 mg.
May contain HCl and/or NaOH for pH adjustment.

Sterile, nonpyrogenic. 0.308 mOsmol/mL (calc).
Mix thoroughly after dilution. Use only if clear and seal is intact and undamaged.
Preservative-free. Use promptly. Discard unused portion.

Use Aseptic Technique
Remove cover from Fliptop vial and cleanse stopper with antiseptic.
With Sterile Syringe and Needle:
1. Aspirate desired portion of vial contents and add to suitable container.
2. Discard any remaining fluid in Fliptop vial.
With DNTD (Double Needle Transfer Device), List No. 4797:
1. Prepare I.V. solution container.
2. Remove one sheath from DNTD and insert needle through rubber diaphragm of Fliptop vial.
3. Remove sheath from DNTD, invert Fliptop vial, and insert through rubber stopper of I.V. solution container. (Fluid in Fliptop vial will be drawn into I.V. solution container by vacuum.)
4.After additive is delivered, remove DNTD and Fliptop vial simultaneously.
See product insert for prescribing information, precautions and warnings.

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

RX ONLY

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

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Dublin, OH 43017

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SODIUM CHLORIDE

sodium chloride injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:55154-0126(NDC:0409-4888)
Route of Administration	INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37, CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	9 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Other Ingredients		
Ingredient Kind	**Ingredient Name**	**Quantity**
May contain	**HYDROCHLORIC ACID** (UNII: QTT17582CB)	
May contain	**SODIUM HYDROXIDE** (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55154-0126-5	5 in 1 BAG	09/08/2011	
1		10 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	NDA018803	09/08/2011	

Labeler - Cardinal Health 107, LLC (118546603)

Revised: 10/2025 Cardinal Health 107, LLC