

SODIUM CHLORIDE - sodium chloride injection, solution
Fresenius Kabi USA, LLC

0.9% Sodium Chloride Injection, USP

Rx only



DESCRIPTION

0.9% Sodium Chloride Injection, USP solution is sterile and nonpyrogenic. It is a parenteral solution containing sodium chloride in water for injection intended for intravenous administration.

For 0.9% Sodium Chloride Injection, USP, each 100 mL contains 900 mg sodium chloride in water for injection. Electrolytes per 1,000 mL: sodium 154 mEq; chloride 154 mEq. The osmolarity is 308 mOsmol/L (calc.).

The pH in the 100 mL and smaller containers is 6.0; for the 250 mL and larger containers, the pH is 5.6. The pH range is 4.5 to 7.0 for all containers.

The solution contains no bacteriostat, antimicrobial agent or added buffer and is intended only as a single-dose injection. When smaller doses are required the unused portion should be discarded.

The solution is a parenteral fluid and electrolyte replenisher.

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

Water for injection, USP is chemically designated H₂O.

The flexible container is fabricated from a specially formulated non-plasticized, film containing polypropylene and thermoplastic elastomers (**freeflex**® bag). The **freeflex**® + bag has a needle-free injection port and can accept standard luer lock syringes to add medication. The amount of water that can permeate from the container into the overwrap is insufficient to affect the solution significantly. Solutions in contact with the flexible container can leach out certain of the container's chemical components in very small amounts within the expiration period. The suitability of the container material has been confirmed by tests in animals according to USP biological tests for plastic containers.

CLINICAL PHARMACOLOGY

When administered intravenously, the solution provides a source of water and electrolytes.

Solutions which provide combinations of hypotonic or isotonic concentrations of sodium chloride are suitable for parenteral maintenance or replacement of water and electrolyte requirements.

Isotonic concentrations of sodium chloride are suitable for parenteral replacement of chloride losses that exceed or equal the sodium loss. Hypotonic concentrations of sodium chloride are suited for parenteral maintenance of water requirements when only small quantities of salt are desired. A hypertonic concentration of sodium chloride may be used to repair severe salt depletion syndrome.

Sodium chloride in water dissociates to provide sodium (Na^+) and chloride (Cl^-) ions. Sodium (Na^+) is the principal cation of the extracellular fluid and plays a large part in the therapy of fluid and electrolyte disturbances. Chloride (Cl^-) has an integral role in buffering action when oxygen and carbon dioxide exchange occurs in the red blood cells. 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.

Water is an essential constituent of all body tissues and accounts for approximately 70% of total body weight. Average normal adult daily requirements range 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

Intravenous solutions containing sodium chloride are indicated for parenteral replenishment of fluid and sodium chloride as required by the clinical condition of the patient.

CONTRAINDICATIONS

None known.

WARNINGS

Sodium Chloride Injection, USP should be used with great care, if at all, in patients with congestive heart failure, severe renal insufficiency and in clinical states in which there exists edema with sodium retention.

The intravenous administration of Sodium Chloride Injection, USP can cause fluid and/or solute overloading resulting in dilution of serum electrolyte concentrations, overhydration, congested states or pulmonary edema.

The risk of dilutive states is inversely proportional to the electrolyte concentration of the injections. The risk of solute overload causing congested states with peripheral and pulmonary edema is directly proportional to the electrolyte concentrations of the injections.

In patients with diminished renal function, administration of Sodium Chloride Injection, USP may result in sodium retention.

PRECAUTIONS

General

Do not use plastic containers in series connections. Such use could result in air embolism due to residual air being drawn from the primary container before administration of the fluid from the secondary container is completed.

Pressurizing intravenous solutions contained in flexible plastic containers to increase flow rates can result in air embolism if the residual air in the container is not fully evacuated prior to administration.

Use of a vented intravenous administration set with the vent in the open position could result in air embolism. Vented intravenous administration sets with the vent in the open position should not be used with flexible plastic containers.

Laboratory Tests

Clinical evaluation and periodic laboratory determinations are necessary to monitor changes in fluid balance, electrolyte concentrations and acid-base balance during prolonged parenteral therapy or whenever the condition of the patient warrants such evaluation.

DRUG INTERACTIONS

Caution must be exercised in the administration of Sodium Chloride Injection, USP to patients receiving corticosteroids or corticotropin.

Carcinogenesis, Mutagenesis, Impairment of Fertility

Studies have not been performed with Sodium Chloride Injection, USP to evaluate the potential for carcinogenesis, mutagenesis or impairment of fertility.

Pregnancy:

Teratogenic Effects

Pregnancy Category C

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

Labor and Delivery

Studies have not been conducted to evaluate the effects of Sodium Chloride Injection, USP on labor and delivery. Caution should be exercised when administering this drug during labor and delivery.

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 Sodium Chloride Injection, USP is administered to a nursing mother.

Pediatric Use

The use of Sodium Chloride Injection, USP in pediatric patients is based on clinical practice.

Plasma electrolyte concentrations should be closely monitored in the pediatric population as this population may have impaired ability to regulate fluids and electrolytes.

Geriatric Use

Clinical studies of Sodium Chloride 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, 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 drug therapy.

This drug is known to be substantially excreted by the kidney, and the risk of toxic reactions to this drug may be greater in patients with impaired renal function. Because elderly patients are more likely to have decreased renal function, care should be taken in dose selection, and it may be useful to monitor renal function.

Do not administer unless solution is clear and container is undamaged. Discard unused portion.

ADVERSE REACTIONS

Reactions which may occur because of the solution or the technique of administration include febrile response, infection at the site of injection, venous thrombosis or phlebitis extending from the site of injection, extravasation and hypervolemia.

If an adverse reaction does occur, discontinue the infusion, evaluate the patient, institute appropriate therapeutic countermeasures and save the remainder of the fluid for examination if deemed necessary.

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.

OVERDOSAGE

In the event of overhydration or solute overload, re-evaluate the patient and institute appropriate corrective measures (see **WARNINGS, PRECAUTIONS, and ADVERSE REACTIONS**).

DOSAGE AND ADMINISTRATION

The dose is dependent upon the age, weight and clinical condition of the patient.

Additives may be incompatible. Consult with pharmacist, if available. When introducing additives, use aseptic technique, mix thoroughly and do not store.

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

INSTRUCTIONS FOR USE

Check flexible container solution composition, lot number, and expiry date.

Do not remove solution container from its overwrap until immediately before use.

Use sterile equipment and aseptic technique.

To Open

1. Turn solution container over so that the text is face down. Using the pre-cut corner tabs, peel open the overwrap and remove solution container.
2. Check the solution container for leaks by squeezing firmly. If leaks are found, or if the seal is not intact, discard the solution.
3. Do not use if the solution is cloudy or a precipitate is present.

To Add Medication

The Additive port of the **freeflex**[®]+ container accepts standard luer lock syringes. **Do not use a needle for additions.**

1. Identify LIGHT BLUE Additive Port with arrow pointing toward solution container.
2. Immediately before injecting additives, break off LIGHT BLUE Additive Port Cap with the arrow pointing toward solution container.
3. Hold base of LIGHT BLUE Additive Port.
4. Attach Luer Lock syringe to the threaded LIGHT BLUE Additive Port. Inject additive.
5. Mix solution container contents thoroughly.

Preparation for Administration

1. Immediately before inserting the infusion set, break off BLUE Infusion Port Cap with the arrow pointing away from container.
2. Use a non-vented infusion set or close the air-inlet on a vented set.
3. Close the roller clamp of the infusion set.
4. Hold the base of BLUE Infusion Port.
5. Insert spike through BLUE Infusion Port by rotating wrist slightly until the spike is inserted.

NOTE: See full directions accompanying administration set.

WARNING: Do not use flexible container in series connections.

HOW SUPPLIED

0.9% Sodium Chloride Injection, USP is supplied in single-dose flexible plastic containers as follows:

Product Code	Unit of Sale	Strength	Each
232150	NDC 65219-218-50	0.9% (450 mg per 50 mL)	NDC 65219-218-01

	Package of 60 freeflex [®] + bags	(9 mg/mL)	50 mL in a 100 mL freeflex [®] + bag
232200	NDC 65219-220-10 Package of 50 freeflex [®] + bags	0.9% (900 mg per 100 mL) (9 mg/mL)	NDC 65219-220-01 100 mL freeflex [®] + bag
232225	65219-222-25 Package of 30 freeflex [®] + bags	0.9% (2,250 mg per 250 mL) (9 mg/mL)	65219-222-01 250 mL freeflex [®] + bag
232250	65219-224-50 Package of 20 freeflex [®] + bags	0.9% (4,500 mg per 500 mL) (9 mg/mL)	65219-224-01 500 mL freeflex [®] + bag

Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. Protect from freezing. The container closure is not made with natural rubber latex. Non-PVC, Non-DEHP, Sterile.



Lake Zurich, IL 60047

www.fresenius-kabi.com/us

Revised: July 2024

PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 50 mL Bag Label

freeflex[®] + NDC 65219-218-01 **50 mL**
0.9% Sodium Chloride Injection, USP

For intravenous use. Rx only



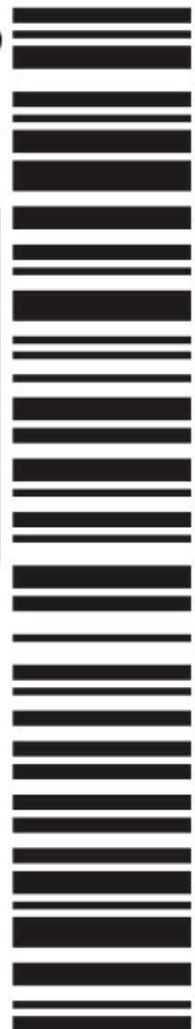
NDC 65219-218-01

50 mL

0.9% Sodium Chloride Injection, USP

For intravenous use.

Rx only



(01)00365219218012

Each 100 mL contains: Sodium Chloride 900 mg in water for injection.	
Electrolytes per 1,000 mL:	
Sodium	154 mEq
Chloride	154 mEq
308 mOsmol/LITER (CALC.)	pH 4.5 to 7.0

Single Dose Only. Discard Unused Portion.

Additives may be incompatible. Consult with pharmacist. When introducing additives, use aseptic technique, mix thoroughly and do not store. Use only if solution is clear and container is undamaged. Must not be used in series connections.

Usual dosage: See package insert.

STORE AT: 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. Protect from freezing.

The container closure is not made with natural rubber latex.

Non-PVC, Non-DEHP, Sterile.

Mfd. for:

 **FRESENIUS KABI**
 Lake Zurich, IL 60047
 Made in Norway

LOT

EXP

403782

www.fresenius-kabi.com/us FPE 1336 01-69-19-014 1 2 3 4 5 6 7 8 9 0

PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 50 mL Case Label

NDC 65219-218-50 232150

0.9% Sodium Chloride Injection, USP

freeflex® +

50 mL x 60

Store at 20° to 25°C (68° to 77°F)

[see USP Controlled Room Temperature].

Protect from freezing.

NDC 65219-218-50

232150



0.9% Sodium Chloride Injection, USP

freeflex®

50 mL x 60

Store at 20° to 25°C (68° to 77°F)

[see USP Controlled Room Temperature].

Protect from freezing.

Manufactured for:



**FRESENIUS
KABI**

Lake Zurich, IL 60047

www.fresenius-kabi.com/us

Made in Norway

EXP: MM-YYYY LOT: 0000000 QTY: 60



(17) YYMMDD (10) 0000000 (30) 60



(01) 30365219218501

63851
FPE 1336 01-89-19-014

PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 100 mL Bag Label

freeflex®+ NDC 65219-220-01 **100 mL**

0.9% Sodium Chloride Injection, USP

For intravenous use. Rx only



NDC 65219-220-01

100 mL

0.9% Sodium Chloride Injection, USP

For intravenous use.

Rx only



Each 100 mL contains: Sodium Chloride 900 mg
in water for injection.

Electrolytes per 1,000 mL:

Sodium	154 mEq
Chloride	154 mEq
308 mOsmol/LITER (CALC.)	pH 4.5 to 7.0

Single Dose Only. Discard Unused Portion.

Additives may be incompatible. Consult with pharmacist. When introducing additives, use aseptic technique, mix thoroughly and do not store. Use only if solution is clear and container is undamaged. Must not be used in series connections.

Usual dosage: See package insert.

STORE AT: 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. Protect from freezing.

The container closure is not made with natural rubber latex.

Non-PVC, Non-DEHP, Sterile.

Mfd. for:

**FRESENIUS
KABI**
Lake Zurich, IL 60047
Made in Norway

LOT

EXP

403783

www.fresenius-kabi.com/us FPE 1335 01-69-19-015 1 2 3 4 5 6 7 8 9 0

PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 100 mL Case Label

NDC 65219-220-10 232200

0.9% Sodium Chloride Injection, USP

freeflex® +

100 mL x 50

Store at 20° to 25°C (68° to 77°F)

[see USP Controlled Room Temperature].

Protect from freezing.

NDC 65219-220-10

232200



0.9% Sodium Chloride Injection, USP

freeflex[®]

100 mL x 50

Store at 20° to 25°C (68° to 77°F)

[see USP Controlled Room Temperature].

Protect from freezing.

Manufactured for:



**FRESENIUS
KABI**

Lake Zurich, IL 60047

www.fresenius-kabi.com/us

Made in Norway

EXP: MM-YYYY LOT: 0000000 QTY: 50



(17) YYMMDD (10) 0000000 (30) 50



(01) 30365219220108

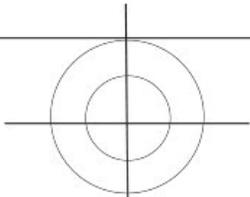
63850
FPE 1335 01-89-19-015

PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 250 mL Bag Label

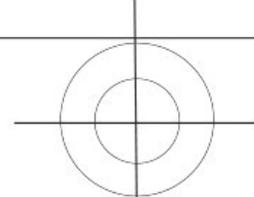
NDC 65219-222-01

freeflex[®] + 250 mL
0.9% Sodium Chloride Injection, USP

For intravenous use. Rx only



NDC 65219-222-01



freeflex[®] 

250 mL

0.9% Sodium Chloride Injection, USP

For intravenous use.

Rx only

50

Each 100 mL contains: Sodium Chloride 900 mg
in water for injection.

Electrolytes per 1,000 mL:

Sodium 154 mEq

Chloride 154 mEq

308 mOsmol/LITER (CALC.) pH 4.5 to 7.0

Single Dose Only. Discard Unused Portion.

100

Additives may be incompatible. Consult with pharmacist. When introducing additives, use aseptic technique, mix thoroughly and do not store. Use only if solution is clear and container is undamaged. Must not be used in series connections.

Usual dosage: See package insert.

150

STORE AT: 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. Protect from freezing.

The container closure is not made with natural rubber latex. Non-PVC, Non-DEHP, Sterile.



200



(01)00365219222019

Mfd. for:



**FRESENIUS
KABI**

LOT

Lake Zurich, IL 60047

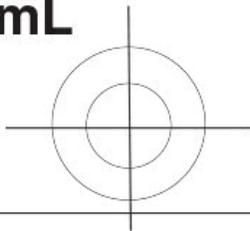
Made in Germany

EXP

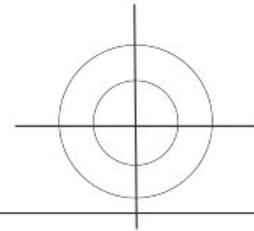
www.fresenius-kabi.com/us

403660

mL



0744331/00 US



PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 250 mL Case Label

NDC 65219-222-25 232225

0.9% Sodium Chloride Injection, USP

freeflex[®] +

250 mL x 30

Store at 20° to 25°C (68° to 77°F)

[see USP Controlled Room Temperature].

Protect from freezing.

NDC 65219-222-25

232225

0.9% Sodium Chloride Injection, USP

freeflex[®]+

250 mL x 30

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

Manufactured for:



**FRESENIUS
KABI**

Lake Zurich, IL 60047

Made in Germany

QTY 30



(01)30365219222256

LOT

EXP

63770

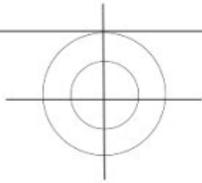
0718661/00 US

PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 500 mL Bag Label

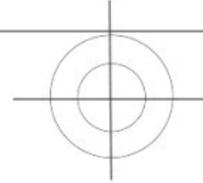
NDC 65219-224-01

freeflex[®]+ **500 mL**
0.9% Sodium Chloride Injection, USP

For intravenous use. Rx only



NDC 65219-224-01



freeflex[®]+

500 mL

0.9% Sodium Chloride Injection, USP

For intravenous use.

Rx only

100

Each 100 mL contains: Sodium Chloride 900 mg
in water for injection.

Electrolytes per 1,000 mL:

Sodium	154 mEq
Chloride	154 mEq
308 mOsmol/LITER (CALC.)	pH 4.5 to 7.0

200

Single Dose Only. Discard Unused Portion.

Additives may be incompatible. Consult with pharmacist. When introducing additives, use aseptic technique, mix thoroughly and do not store. Use only if solution is clear and container is undamaged. Must not be used in series connections.

300

Usual dosage: See package insert.

STORE AT: 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. Protect from freezing.

The container closure is not made with natural rubber latex. Non-PVC, Non-DEHP, Sterile.

400



(01)00365219224013

Mfd. for:



**FRESENIUS
KABI**

Lake Zurich, IL 60047

LOT

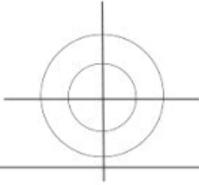
Made in Germany

EXP

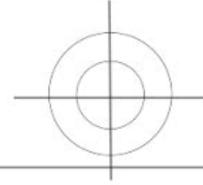
www.fresenius-kabi.com/us

403661

mL



0744341/00 US



PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 500 mL Case Label

NDC 65219-224-50 232250

0.9% Sodium Chloride Injection, USP

freeflex® +

500 mL x 20

Store at 20° to 25°C (68° to 77°F)

[see USP Controlled Room Temperature].

Protect from freezing.

NDC 65219-224-50	232250	QTY 20
0.9% Sodium Chloride Injection, USP		
freeflex® +		Manufactured for:
500 mL x 20		 FRESENIUS KABI
Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. Protect from freezing.		Lake Zurich, IL 60047 Made in Germany
	LOT	
	EXP	
(01)30365219224502	63771	0718671/00 US

SODIUM CHLORIDE

sodium chloride injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65219-220
Route of Administration	INTRAVENOUS		

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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65219-220-10	50 in 1 CASE	11/17/2021	
1	NDC:65219-220-01	100 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207310	09/19/2017	

SODIUM CHLORIDE

sodium chloride injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65219-218
Route of Administration	INTRAVENOUS		

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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65219-218-50	60 in 1 CASE	11/17/2021	
1	NDC:65219-218-01	50 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207310	09/19/2017	

SODIUM CHLORIDE

sodium chloride injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65219-222
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37, CHLORIDE ION - UNII:Q32Z N48698)	SODIUM CHLORIDE	9 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65219-222-25	30 in 1 CASE	11/17/2021	
1	NDC:65219-222-01	250 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207310	09/19/2017	

SODIUM CHLORIDE

sodium chloride injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65219-224
Route of Administration	INTRAVENOUS		

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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65219-224-50	20 in 1 CASE	11/17/2021	
1	NDC:65219-224-01	500 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207310	09/19/2017	

Labeler - Fresenius Kabi USA, LLC (013547657)

Establishment

Name	Address	ID/FEI	Business Operations
HP Halden		247747272	ANALYSIS(65219-220, 65219-218) , MANUFACTURE(65219-220, 65219-218) ,

Pharma AS		541141313	PACK(65219-220, 65219-218)
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Establishment

Name	Address	ID/FEI	Business Operations
Fresenius Kabi Deutschland GmbH		506719546	ANALYSIS(65219-222, 65219-224) , MANUFACTURE(65219-222, 65219-224) , API MANUFACTURE(65219-222, 65219-224)

Establishment

Name	Address	ID/FEI	Business Operations
Fresenius Kabi USA, LLC		964475045	ANALYSIS(65219-220, 65219-218, 65219-222, 65219-224) , MANUFACTURE(65219-220, 65219-218, 65219-222, 65219-224)

Revised: 1/2026

Fresenius Kabi USA, LLC