

**0.9% SODIUM CHLORIDE INJECTION, 1000ML, BLUE PHARMA, TOTALCARE-
sodium chloride 0.9% w/v for intravenous infusion injection, solution
0.9% SODIUM CHLORIDE INJECTION, 1000ML, DRE BIOLAB, TOTALCARE-
sodium chloride 0.9% w/v for intravenous infusion injection, solution
0.9% SODIUM CHLORIDE INJECTION, 500ML, DRE BIOLAB, TOTALCARE-
sodium chloride 0.9% w/v for intravenous infusion injection, solution
0.9% SODIUM CHLORIDE INJECTION, 500ML, BLUE PHARMACEUTICALS,
TOTALCARE- sodium chloride 0.9% w/v for intravenous infusion injection,
solution
AMCO INTERNATIONAL CORP**

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

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. The 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.

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.

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

None known.

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.

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.

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

Carcinogenesis, Mutagenesis, Impairment of Fertility

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.

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.

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.

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.

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.

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.

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

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).

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 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.

Important Prescribing Information

Subject: Marketing of Pending Approval 0.9% Sodium Chloride Injection to address drug shortages

Dear Healthcare Professional,

To prevent a drug shortage of large volume parenteral fluid drug products, we are coordinating with the U.S. Food and Drug Administration (FDA) to temporarily import 0.9% Sodium Chloride Injection (250 mL; 500 mL and 1,000 mL) from specific DRE manufacturing facilities. FDA has not approved these products manufactured by this facility.

You may be provided with additional letters for other imported products you receive. Please read each letter in its entirety because each letter may contain different product specific information.

Effective immediately, and during this temporary period, we will offer the following imported products:

It is important to note the following:

- After opening the carton or box, the bags should be inspected visually to confirm there is no visible particulate matter or bag defects, such as leaks. Container integrity is imperative to ensure sterility of products listed in the table above. Parenteral drug products should be inspected visually for particulate matter and bag defects prior to administration, whenever solution or container permits.
USE A NEW BAG IF PARTICULATES ARE VISIBLE OR IF THE IV BAG CONTAINS A LEAK.

Reporting Adverse Events or Product Quality Issues

To report adverse events associated with these imported products, please call

. Adverse events or quality problems experienced with the use of these imported products may also be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail or by fax:

- Complete and submit the report Online: www.fda.gov/medwatch/report.htm
- Regular mail or Fax: Download form www.fda.gov/MedWatch/getforms.htm or call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178 (1-800-332-0178).

To report product quality issues associated with these imported products, please contact

If you have any questions about the information contained in this letter or the use of the imported products, please contact

To place an order, please contact

PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 1000 mL Bag Label

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

[NDC 84866-6667-1]

DRE HEALTH TotalCare

For intravenous use. Rx only

1 000ml 0.9% Sodium Chloride Injection, USP

Electrolytes per 1,000 ml:

Sodium

Chloride

308 mOsmol/LITER (CALC.)

SingleDoseOnly. DiscardUnusedPortion.

154 mEq

154 mEq pH 4.5to 7.0

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 RoomTemperature].Protectfromfreezing.

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

MFG BY: AMCO INTERNATIONAL CORP

EXP. LOT.

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

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

[NDC 84866-6681-1]

DRE BioLab TotalCare

For intravenous use. Rx only

500ml 0.9% Sodium Chloride Injection, USP

Electrolytes per 1,000 ml:

Sodium

Chloride

308 mOsmol/LITER (CALC.)

SingleDoseOnly. DiscardUnusedPortion.

154 mEq

154 mEq pH 4.5to 7.0

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. °°°°

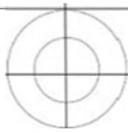
STORE AT: 20 to 25 C (68 to 77 F) [see USP Controlled RoomTemperature].Protectfromfreezing.

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

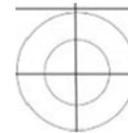
MFG BY: AMCO INTERNATIONAL CORP

EXP. LOT.

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NDC XXXXXX



TotalCare

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.

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MFG BY:
AMCO INTERNATIONAL
CORP

EXP
LOT

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

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

[NDC 84866-6681-2]

DRE BioLab TotalCare

For intravenous use. Rx only

500ml 0.9% Sodium Chloride Injection, USP

Electrolytes per 1,000 ml:

Sodium

Chloride

308 mOsmol/LITER (CALC.)

SingleDoseOnly. DiscardUnusedPortion.

154 mEq

154 mEq pH 4.5to 7.0

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. °°°°

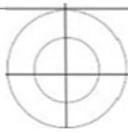
STORE AT: 20 to 25 C (68 to 77 F) [see USP Controlled RoomTemperature].Protectfromfreezing.

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

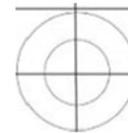
MFG BY: AMCO INTERNATIONAL CORP

EXP. LOT.

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NDC XXXXXX



TotalCare

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.

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MFG BY:
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EXP
LOT

PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 1000 mL Case Label

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

[NDC 84866-6666-2]

DRE BioLab TotalCare

For intravenous use. Rx only

500ml 0.9% Sodium Chloride Injection, USP

Electrolytes per 1,000 ml:

Sodium

Chloride

308 mOsmol/LITER (CALC.)

SingleDoseOnly. DiscardUnusedPortion.

154 mEq

154 mEq pH 4.5to 7.0

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 RoomTemperature].Protectfromfreezing.

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

MFG BY: AMCO INTERNATIONAL CORP

EXP. LOT.

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PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 1000 mL Case Label

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

[NDC 84866-6667-2]

DRE BioLab TotalCare

For intravenous use. Rx only

500ml 0.9% Sodium Chloride Injection, USP

Electrolytes per 1,000 ml:

Sodium

Chloride

308 mOsmol/LITER (CALC.)

SingleDoseOnly. DiscardUnusedPortion.

154 mEq

154 mEq pH 4.5to 7.0

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. °°°°

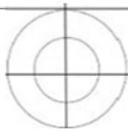
STORE AT: 20 to 25 C (68 to 77 F) [see USP Controlled RoomTemperature].Protectfromfreezing.

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

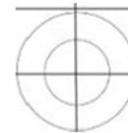
MFG BY: AMCO INTERNATIONAL CORP

EXP. LOT.

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NDC XXXXXX



TotalCare

1000 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.

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EXP
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PACKAGE LABEL - PRINCIPAL DISPLAY - 0.9% Sodium Chloride 1000 mL Bag Label

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

[NDC 84866-6666-1]

DRE HEALTH TotalCare

For intravenous use. Rx only

1 000ml 0.9% Sodium Chloride Injection, USP

Electrolytes per 1,000 ml:

Sodium

Chloride

308 mOsmol/LITER (CALC.)

SingleDoseOnly. DiscardUnusedPortion.

154 mEq

154 mEq pH 4.5to 7.0

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.

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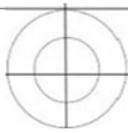
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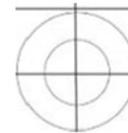
Blue Pharma 1000mL Bag Label

Blue Pharma 1000mL case label

BLUE PHARMA 500mL bag label



NDC XXXXXX



TotalCare

500 mL

0.9% Sodium Chloride Injection, USP

For intravenous use.

Rx only

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 in water for injection.
 Electrolytes per 1,000 mL:

Sodium	154 mEq
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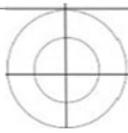
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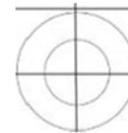
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AMCO INTERNATIONAL
CORP

EXP
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BLUE PHARMA 500ml CASE LABEL



NDC XXXXXX



TotalCare

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

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CORP

EXP
LOT

0.9% SODIUM CHLORIDE INJECTION, 1000ML, BLUE PHARMA, TOTALCARE

sodium chloride 0.9% w/v for intravenous infusion injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:84866-6669
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698, SODIUM CATION - UNII:LYR4M0NH37)	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:84866-6669-2	20 in 1 CASE	10/31/2024	
1	NDC:84866-6669-1	1000 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
unapproved drug other		10/30/2024	

0.9% SODIUM CHLORIDE INJECTION, 1000ML, DRE BIOLAB, TOTALCARE

sodium chloride 0.9% w/v for intravenous infusion injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:84866-6666
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698, SODIUM CATION - UNII:LYR4M0NH37)			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:84866-6666-2	20 in 1 CASE	10/31/2024	
1	NDC:84866-6666-1	1000 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
unapproved drug other			10/30/2024	

0.9% SODIUM CHLORIDE INJECTION, 500ML, DRE BIOLAB, TOTALCARE

sodium chloride 0.9% w/v for intravenous infusion injection, solution

Product Information				
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:84866-6681(NDC:65219-224)	
Route of Administration	INTRAVENOUS			
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698, SODIUM CATION - UNII:LYR4M0NH37)			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:84866-6681-2	20 in 1 CASE	10/31/2024	
1	NDC:84866-6681-1	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	10/30/2024	

0.9% SODIUM CHLORIDE INJECTION, 1000ML, DRE BIOLAB, TOTALCARE

sodium chloride 0.9% w/v for intravenous infusion injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:84866-6667
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698, SODIUM CATION - UNII:LYR4MONH37)	SODIUM CHLORIDE	9 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84866-6667-2	20 in 1 CASE	10/31/2024	
1	NDC:84866-6667-1	1000 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
unapproved drug other		10/30/2024	

0.9% SODIUM CHLORIDE INJECTION, 500ML, BLUE PHARMACEUTICALS, TOTALCARE

sodium chloride 0.9% w/v for intravenous infusion injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:84866-6668
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698, SODIUM CATION - UNII:LYR4MONH37)	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:84866-6668-2	20 in 1 CASE	10/31/2024	
1	NDC:84866-6668-1	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
unapproved drug other		10/30/2024	

Labeler - AMCO INTERNATIONAL CORP (116924852)

Establishment

Name	Address	ID/FEI	Business Operations
AMCO INTERNATIONAL CORP		116924852	manufacture(84866-6666, 84866-6667, 84866-6668, 84866-6669) , relabel(84866-6681) , repack(84866-6681)