



B. Braun Medical Inc.  
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### IMPORTANT PRESCRIBING INFORMATION

**Subject: Temporary Importation of 0.9% Sodium Chloride w/v Intravenous Infusion in Ecoflac Plus Containers**

Dear Health Care Provider,

The purpose of this letter is to inform you of an additional foreign product that B. Braun Medical Inc. (B. Braun) will be providing in the United States (U.S.) to address the critical drug shortage of 0.9% Sodium Chloride for Injection. Due to the shortage of 0.9% Sodium Chloride Injection products in the U.S., B. Braun is coordinating with the U.S. Food and Drug Administration (FDA) to temporarily import foreign Sodium Chloride 0.9% w/v Intravenous Infusion BP in the Ecoflac plus container into the U.S. market. The foreign product is manufactured at an FDA inspected B. Braun sterile injectable facility in Melsungen, Germany which is currently in compliance with FDA regulations.

At this time, no other entity except B. Braun is authorized by the FDA to import or directly/indirectly distribute Sodium Chloride 0.9% w/v Intravenous Infusion BP in the Ecoflac® plus container in the U.S. FDA has not approved B. Braun's Sodium Chloride 0.9% w/v Intravenous Infusion BP in the Ecoflac plus container in the United States.

Effective immediately and during this temporary period, B. Braun will offer the following presentations of B. Braun's Sodium Chloride 0.9% w/v Intravenous Infusion BP in the Ecoflac plus container:

Product Name	Volume	Ingredients
Sodium Chloride 0.9% w/v Intravenous Infusion BP in the Ecoflac plus container	500 mL	Each 500 mL contains: Sodium Chloride 4.5 g, Water for Injections to 500 mL Electrolytes per 500 mL is Sodium 77 mmol, Chloride 77 mmol

**Ecoflac plus container- Not made with natural rubber latex, PVC, or DEHP.**

**Indications and Usage and Dosage Administration**

The foreign unapproved product, packaged in a semi-rigid Ecoflac® (polyethylene plastic) container, contains the same active ingredient in the same concentration as the 0.9% Sodium Chloride Injection products approved in the U. S., packaged in the flexible plastic EXCEL® (ethylene-propylene copolymer) bag. As such, clinical practice pertaining to indication, usage and dosage administration for Sodium Chloride 0.9% w/v Intravenous Infusion BP in the Ecoflac plus container is the same as with the EXCEL® containers.

However, some key differences between the Ecoflac and EXCEL® container packaging and labeling are described below in the Product Comparison Table.

It is also important to note that the Ecoflac plus container is a semi rigid plastic bottle. However, no venting is necessary during infusion. Ecoflac plus collapses completely when emptying.

Ecoflac plus container and carton labeling may include barcodes that may not register accurately in the U.S. scanning systems. Institutions should manually input the product into their systems and confirm that barcode systems do not provide incorrect information when the product is scanned. Alternative procedures should be followed to assure that the correct drug product is being used and administered to individual patients.

**Reporting Adverse Events**

To report adverse events or quality problems with B. Braun's Sodium Chloride 0.9% w/v Intravenous Infusion BP in the Ecoflac plus container, please contact the B. Braun Clinical and Technical Support Department at 1-800-854-6851. Adverse events or quality problems experienced with the use of this product may also be reported to the FDA's MedWatch Adverse Event Reporting Program either online, or regular mail, or by fax:



- Complete and submit the report **Online:** [www.fda.gov/medwatch/report.htm](http://www.fda.gov/medwatch/report.htm)
- Regular mail or Fax: Download form [www.fda.gov/MedWatch/getforms.htm](http://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.

Thank you for your commitment to B.Braun's IV solution products.





Rick Williamson  
Vice President of Pharmaceutical Marketing

**PRODUCT COMPARISON TABLE**

Item	German Product – Ecoflac Container	US Product – EXCEL® Container
Product Drawing	<p style="text-align: center;"><b>Ecoflac® Plus</b></p> 	<p style="text-align: center;"><b>EXCEL®</b></p> 
Product Description	0.9% w/v Sodium Chloride BP	0.9% Sodium Chloride Injection USP
Product Code	9999-10	L8001
Unit Bar Code	No	Yes (NDC and Lot/Exp)
NDC #	0264-9999-10	0264-7800-10
Volume	500 mL	500 mL

**PRODUCT COMPARISON TABLE**

Item	German Product – Ecoflac Container	US Product – EXCEL® Container
Case Quantity	10 units per case	24 units per case
Storage Condition	Do not store above 25°C	Store at room temperature (25°C)
Shelf Life	36 months	30 months
Ingredients	Each 500 mL contains: Sodium Chloride USP 450 mg, Water for Injection USP to 500 mL Total Electrolytes per 500 mL : Sodium 77 mEq, Chloride 77 mEq	Each 500 mL contains: Sodium Chloride USP 450 mg, Water for Injection USP to 500 mL. Total Electrolytes per 500 mL : Sodium 77 mEq, Chloride 77 mEq
Container Type	Ecoflac®Plus, Polyethylene Plastic Containers (semi rigid plastic bottle)	Excel®, Primary plastic container with a clear overwrap (flexible plastic bag)
Container material	Low Density Polyethylene (LDPE)	Copolymer of ethylene and propylene
Container Description	Blow/Fill/Seal (BFS)	Form/Fill/Seal (FFS)
PVC, DEHP, Latex	No	No
Overwrap	No	Yes
Closure Description	Twin cap with peel tab cover	Additive port with elastomeric stopper, Administration port with plastic cover
Pictures of Difference in Container Port System	Twin Port with peel tab cover and two interchangeable access ports with thermoplastic elastomeric septum 	Separate additive and administration ports 

**PRODUCT COMPARISON TABLE**

<b>Item</b>	<b>German Product – Ecoflac Container</b>	<b>US Product – EXCEL® Container</b>
Additive Port Material	Thermoplastic elastomer	Synthetic Isoprene
Needle Size	15 -21 gauge	18-22 gauge
Additive Volume	120 mL	50 mL
Spike Port Resealable	Yes	No
Pressure Infusion	No	Yes, not to exceed 300mm Hg
Sterilization Process	Terminal Steam Sterilization  Sterility Test (Ph. Eur.) Sterility Assurance Level of 10 <sup>-6</sup>	Terminal Steam Sterilization  Parametric Release Sterility Assurance Level of 10 <sup>-6</sup>