HEB SALINE- nasal spray
HEB

Saline Nasal Spray

Active Ingredients

Sodium Chloride 0.65%

Purpose

Moisturizer

Uses

provides instant, soothing relief dry irritated nasal passages due to colds, allergies, dry air, pollution, smoke, air travel and use of decongestants/steroidal sprays.

Warnings

If pregnant or breast-feeding, ask a healthcare professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away at 1-800-222-1222

Directions

For children and adults,squeeze bottle twice in each nostril as often as needed or as directed by physician. **For infants,**use drop application. Hold bottle upright for spray, horizontally for stream, and upside down for drop. The use of this dispenser by more than one person may spread infection.

Other Information

- store at room temperature

Inactive Ingredients

benzalkonium chloride, benzyl alcohol, purified water, sodium phosphate dibasic, sodium phosphate monobasic

Questions or comments?

1-866-467-2748

*This product is not manufactured or distributed by Valeant Pharmaceuticals North America LLC, owner of the registered trademark Ocean®.

Saline Nasal Spray

Compare to the active ingredient in Ocean®*

NDC 37808-845-88

Saline Nasal Spray

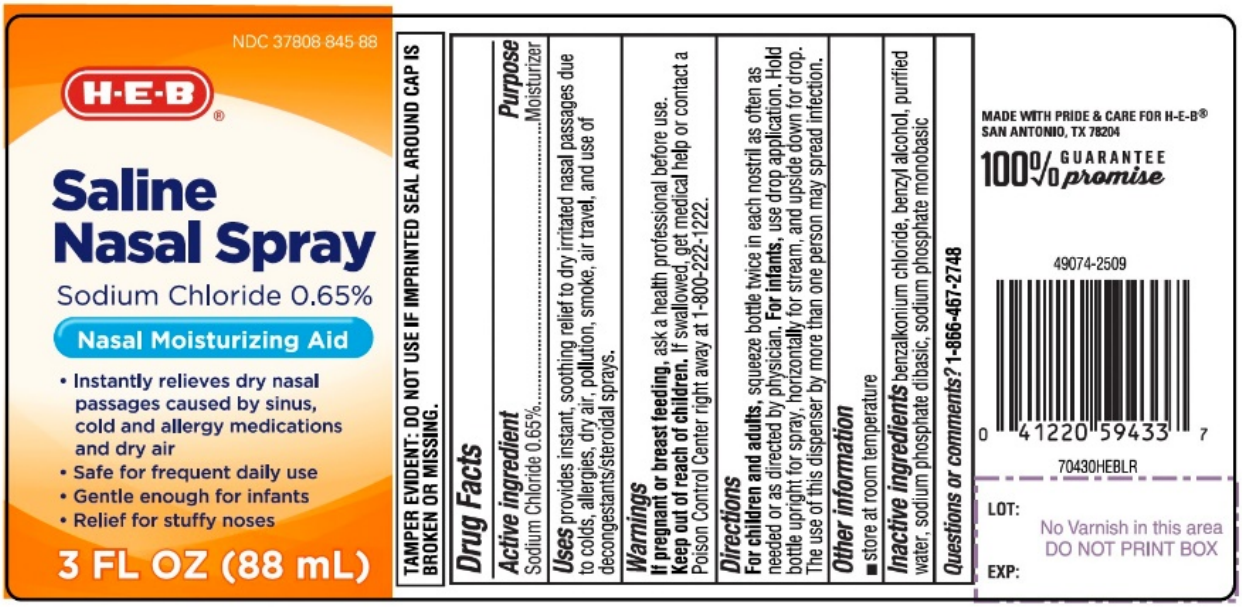
Sodium Chloride 0.65%

- Instantly relieves dry nasal passages caused by sinus, cold and allergy medications and dry air
- Safe for frequent daily use
- Gentle enough for infants
- Relief for stuffy noses

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL AROUND CAP IS BROKEN OR MISSING.

Distributed by:

Package Label for 88 mL



HEB SALINE
nasal spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-845
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	6.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
WATER (UNII: 059QF0KO0R)	
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JIH2SW)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-845-88	1 in 1 CARTON	04/11/2025	
1		88 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/27/2021	

Labeler - HEB (007924756)

Revised: 10/2025 HEB