

MEIJER SALINE- nasal spray

MEIJER

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Saline Nasal Spray

Drug Facts

Active Ingredients

Sodium Chloride 0.65%

Purpose

Moisturizer

Uses

provides instant, soothing relief to 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

Aloe barbadensis leaf juice (aloe vera gel), 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, owners of the registered trademark Ocean®.

DIST. BY MEIJER DISTRIBUTION INC.

GRAND RAPIDS, MI 49544

Saline Nasal Spray

NDC: 41250-636-30

saline Nasal Spray

Sodium Chloride 0.65%

+ Soothing Aloe

Instantly relieves dry nasal passages caused by sinus, cold and allergy medications and dry air

Gentle enough for infants

Relief for stuffy noses

3 FL O.Z. (89 mL)

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

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LOT: 75930MJRLR

EXP:

13733 80175 1

PID: 45505688

DIST. BY MEIJER DISTRIBUTION, INC.
ONE AND A HALF CENTERS
www.meijer.com

MEIJER SALINE

nasal spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-636
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
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
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:41250-636-30	1 in 1 CARTON	02/17/2021	
1		89 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 final	part349	02/17/2021	

Labeler - MEIJER (006959555)

Revised: 1/2023

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