#### 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.

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#### 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%

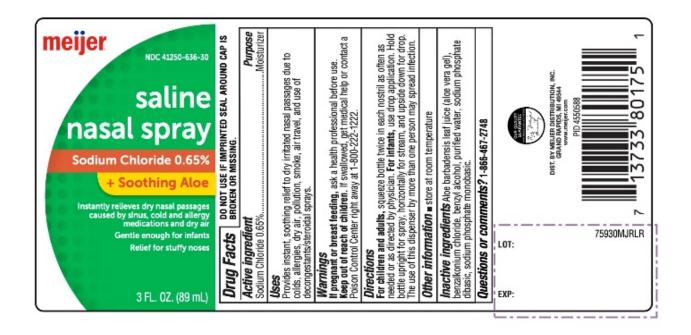
+ Soothing Aloe

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.

Distributed by:



March Andrews to be the state

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			<b>Basis of Strength</b>		Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698			SODIUM CHLORIDE		6.5 mg in 1 mL
Institute Ingradiante					
Inactive Ingredients					
	Ingredient Name	•			Strength
ALOE VERA LEAF (UNII: ZY81Z83	-				
BENZALKONIUM CHLORIDE (UNI	,				
BENZYL ALCOHOL (UNII: LKG8494	4WBH)				
WATER (UNII: 059QF0KO0R)					
		CD696LDA74)			
SODIUM PHOSPHATE, DIBASIC,	UNSPECIFIED FORM (UNI	I. GRUOULDA/4)			

Pa	ackaging			
#	ltem 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		
Μ	larketing l	nformation		
M	l <b>arketing</b>   Marketing Category	nformation Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

Labeler - MEIJER (006959555)

Revised: 1/2023

MEIJER