CLEAR MIST- sodium chloride liquid
Leosons Overseas Corp

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.

Clear Mist (Sheffield Pharmaceutical)

Drug Facts

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Active ingredient & Purpose

Active ingredients	***Purpose***
Sodium chloride 0.65%	Moisturizer

Uses

- For dry nasal membranes

Warnings

Do not use

- if seal is broken or missing.

Keep out of reach of children.

The use of this dispenser by more than one person may spread infection.

Directions

- squeeze twice in each nostril as needed
- upright delivers a spray, horizontally a stream, upside down a drop

Inactive Ingredients

benzalkonium chloride, disodium phosphate, phenylcarbinol, monosodium phosphate, water.

Questions or comments?

Call toll-free at **1-855-452-9500** or email at info@Leosonsintl.com

Distributed by:

LEOSONS

10 Maryland Avenue
Albany, New York 12205

PRINCIPAL DISPLAY PANEL

Clear Mist ®

Saline

Nasal

Spray

Contains Sodium Chloride 0.65%

Alcohol Free

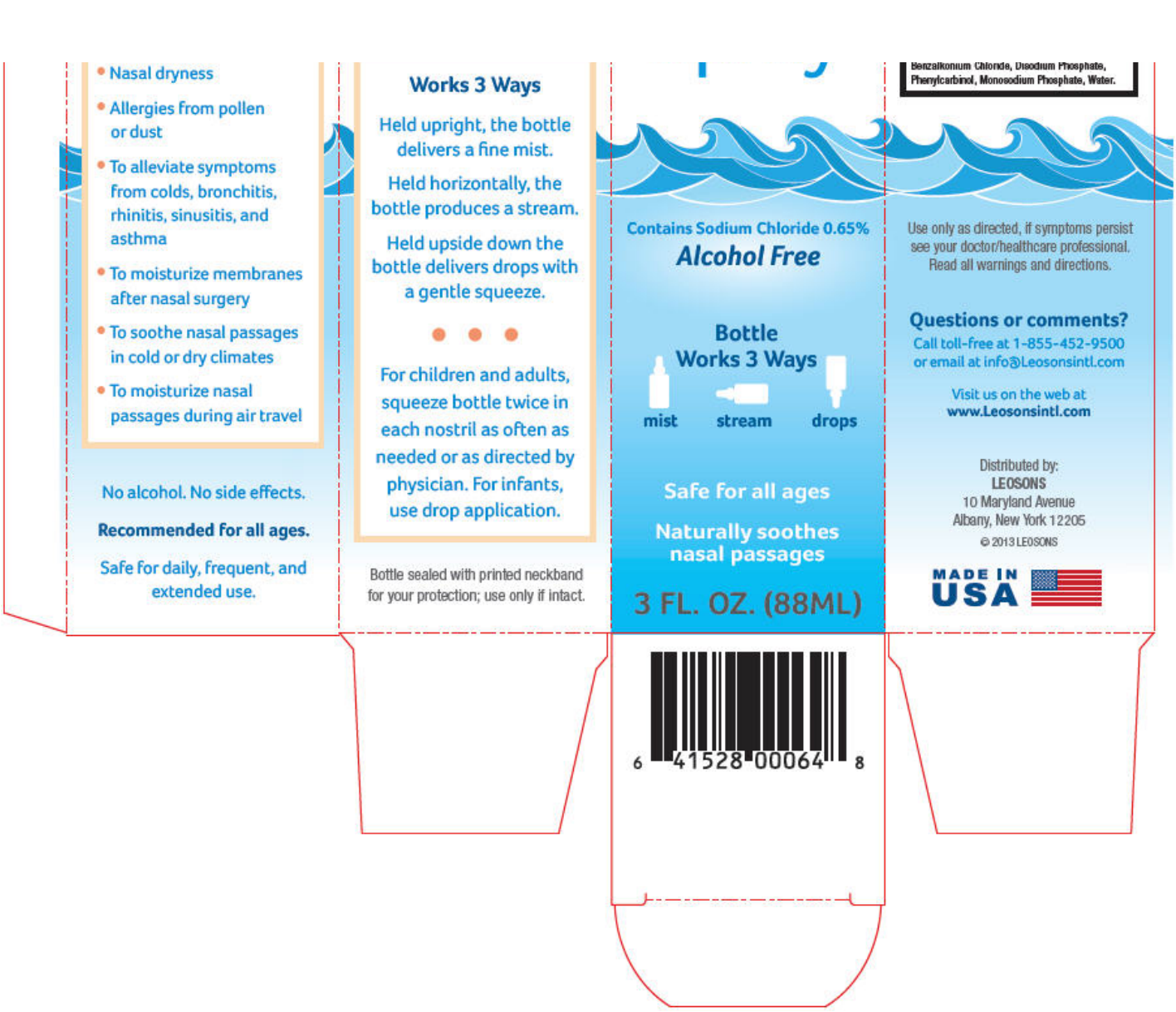
Bottle

Works 3 Ways

mist stream drops

Safe for all ages

Naturally soothes nasal passages



CLEAR MIST

sodium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69626-0064
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698, SODIUM CATION - UNII:LYR4M0NH37)	SODIUM CHLORIDE	6.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPUU)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69626-0064-8	1 in 1 CARTON	01/01/2016	
1		88 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
2	NDC:69626-0064-5	1 in 1 CARTON	01/01/2016	
2		44 mL in 1 BOTTLE, SPRAY; 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	01/01/2016	

Labeler - Leosons Overseas Corp (148605470)

Revised: 12/2021 Leosons Overseas Corp