SUNMARK NASAL- oxymetazoline hcl spray
Strategic Sourcing Services LLC

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.

McKesson Nasal Spray Drug Facts

Active ingredient

Oxymetazoline hydrochloride 0.05%

Purpose

Nasal decongestant

Uses

- temporarily relieves nasal congestion due to:
- common cold
- hay fever
- upper respiratory allergies
- temporarily relieves sinus congestion and pressure
- shrinks swollen nasal membranes so you can breathe more freely

Warnings

Ask a doctor before use if you have

- heart disease
- high blood pressure
- diabetes
- thyroid disease
- trouble urinating due to an enlarged prostate gland

When using this product

- do not use more than directed
- do not use for more than 3 days. Use only as directed. Frequent or prolonged use may cause nasal congestion to recur or worsen.
- temporary discomfort such as burning, stinging, sneezing or an increase in nasal discharge may occur
- use of this container by more than one person may spread infection

Stop use and ask a doctor if

symptoms persist

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

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

Directions

- adults and children 6 to under 12 years of age (with adult supervision): 2 or 3 sprays in each nostril not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24-hour period.
- children under 6 years of age: ask a doctor

To spray, squeeze bottle quickly and firmly. Do not tilt head backward while spraying. Wipe nozzle clean after use. Replace cap tightly to maintain child resistance.

Other information

- store at 20-25°C (68-77°F)
- retain carton for future reference on full labeling

Inactive ingredients

benzalkonium chloride solution, benzyl alcohol, dibasic sodium phosphate, edetate disodium, glycerin, monobasic sodium phosphate, polyethylene glycol, povidone, propylene glycol, purified water

Questions or comments?

1-800-719-9260

Principal Display Panel

sunmark®

COMPARE TO AFRIN® NASAL SPRAY ACTIVE INGREDIENT

See New Directions

nasal spray

oxymetazoline HCl 0.05%

nasal decongestant

Extra Moisturizing

Fast, powerful congestion relief

For colds & allergies

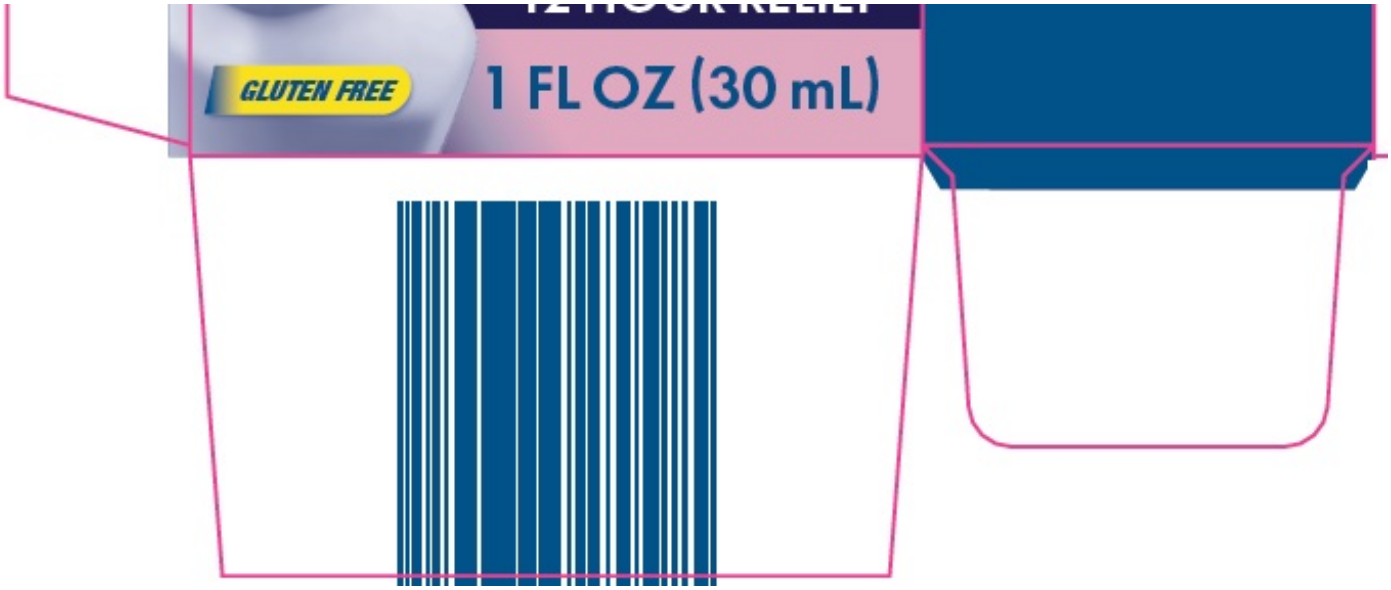
Soothes and helps rehydrate dry noses

12 HOUR RELIEVE

GLUTEN FREE

1 FL OZ (30 mL)


sunmark®
COMPARE TO
AFRIN® NASAL SPRAY
ACTIVE INGREDIENT*
NDC 49348-230-27
See New Directions
nasal
spray
oxymetazoline HCl 0.05% -
nasal decongestant
Extra Moisturizing
Fast, powerful
congestion relief
For colds & allergies
Soothes and helps
rehydrate dry noses
12 HOUR RELIEF
sunmark®
nasal
spray
oxymetazoline HCl 0.05% -
nasal decongestant
Extra Moisturizing


GLUTEN FREE
1 FL OZ (30 mL)

sunmark®

nasal spray

oxymetazoline HCl 0.05% -
nasal decongestant

Drug Facts

Active ingredient — Purpose

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Drug Facts (continued)

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Questions or comments?

1-800-719-9260

*This product is not manufactured or distributed by MSD Consumer Care, Inc., distributor of Afrin® Nasal Spray.

DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING

McKESSON

Another Quality Product
Distributed By McKesson
One Post Street,
San Francisco, CA 94104
Money Back Guarantee
Please visit us at
www.sunmarkbrand.com

SUNMARK NASAL

oxymetazoline hcl spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49348-230
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.05 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
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**
	NDC:49348			

1	NDC:49348-230-27	1 in 1 CARTON	09/19/2003	
1		30 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	part341	09/19/2003	

Labeler - Strategic Sourcing Services LLC (116956644)

Revised: 11/2022 Strategic Sourcing Services LLC