SUNMARK NASAL ORIGINAL- oxymetazoline hydrochloride spray
Strategic Sourcing Services LLC

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 Use: Push firmly down on cap and turn counter clockwise. To spray, squeeze bottle quickly and firmly. Do not tilt head backward while spraying. Wipe nozzle clean after use. Secure cap after use.

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, monobasic sodium phosphate, polyethylene glycol, povidone, propylene glycol, purified water

Questions or comments?

1-800-719-9260

Principal Display Panel

COMPARE TO AFRIN® ORIGINAL ACTIVE INGREDIENT

nasal spray

oxymetazoline HCl 0.05% -

nasal decongestant

Maximum Strength

Original

Fast, powerful congestion relief

Colds & allergies

12 HOUR RELIEF

GLUTEN FREE

1 FL OZ (30 mL)

sunmark®

nasal spray

oxymetazoline HCl 0.05% - nasal decongestant

Maximum Strength
Original

McKESSON

Another Quality Product
Distributed By McKesson
6555 State Highway 161, Irving, TX 75039
Money Back Guarantee
Please visit us at www.sunmarkbrand.com

Drug Facts

Active ingredient — **Purpose**
Oxymetazoline hydrochloride 0.05%......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
■ thyroid disease ■ diabetes
■ 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 Use: Push firmly down on cap and turn counter clockwise. To spray, squeeze bottle quickly and firmly. Do not tilt head backward while spraying. Wipe nozzle clean after use. Secure cap after use.

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

Drug Facts (continued)

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

Questions or comments?
1-800-719-9260

*This product is not manufactured or distributed by Bayer HealthCare LLC, distributor of Afrin® Original.

How to use:
TO USE: Push firmly down on cap and turn counter clockwise. Secure cap after use.

Push

TURN

DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING

0 10939 41433 5

30410 S1 C6

SUNMARK NASAL ORIGINAL

oxymetazoline hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49348-028
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 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JIH2SW)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	WHITE (Translucent)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49348-028-27	1 in 1 CARTON	06/27/2003	
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	06/27/2003	

Labeler - Strategic Sourcing Services LLC (116956644)

Revised: 2/2020   Strategic Sourcing Services LLC