

FAMILY CARE NASAL RELIEF- oxymetazoline hydrochloride spray

United Exchange 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.

Active ingredient

Purpose

Oxymetazoline HCl 0.05%.....Nasal Decongestant

Uses

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

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.

Directions

- adults and children 6 to 12 years of age (with adult supervision): 2 or 3 sprays in each nostril not more 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. Shake well before use. To open, rotate cap to align the marks. Squeeze cap on both sides in a counter-clockwise turn and pull off to remove. To spray, remove clamp and hold bottle with thumb at base and nozzle between first and second fingers. Without tilting the head, insert nozzle into nostril. Fully depress rim with a firm, even stroke and sniff deeply. Wipe nozzle clean after use and snap cap back onto bottle.

Other information

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

Inactive ingredients benzalkonium chloride, benzyl alcohol, dibasic sodium phosphate hydrate, disodium EDTA, distilled water, monobasic sodium phosphate dihydrate, PEG 1450, PVP K30

Distributed By:

United Exchange Corp
17211 Valley View Ave.
Cerritos, CA 90703 USA
Made in Korea

Drug Facts

Active ingredient Oxymetazoline HCl 0.05%.....**Purpose** Nasal decongestant

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EXP:
LOT:

FAMILY CARE

NASAL RELIEF

Pump Mist Spray

Our extra strength formulation prevents dripping from the nose to the throat, working almost instantly to give 12 hours of nasal relief.

ORIGINAL

SAFETY SEALED: Do not use if printed seal on bottle is broken or missing.

Drug Facts (continued)

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Inactive ingredients

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Questions?

call 1-800-814-8028

*This product is not manufactured or distributed by Bayer, owner of the registered trademark Afrin® Original.

FAMILY CARE

NASAL RELIEF

Pump Mist Spray

ORIGINAL

Compare to AFRIN® Original active ingredient*

CHILD SAFETY CAP

FAMILY CARE

NASAL RELIEF

Pump Mist Spray

ANTI-DRIP

Features:

- Fast & Powerful Congestion Relief
- Long Lasting Relief

ORIGINAL

12

HOUR RELIEF
0.5 FL OZ (15 mL)



Compare to AFRIN® Original active ingredient*

FAMILY CARE

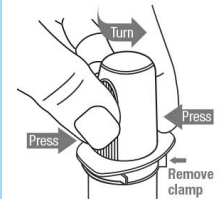
FULL DIRECTIONS IN DRUG FACTS PANEL

DIRECTIONS FOR USE:

STEP 1: To open, rotate cap to align the marks. Squeeze cap on both sides in a counter-clockwise turn and pull off to remove. To spray, remove clamp.

STEP 2: Use as directed.

STEP 3: Wipe nozzle clean after use and snap cap back onto bottle.



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United Exchange Corp.
17211 Valley View Ave.
Cerritos, CA 90703 U.S.A.
www.ueccorp.com
Toll Free: 1 800 814 8028
Made in Korea



FAMILY CARE NASAL RELIEF

oxymetazoline hydrochloride spray

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:65923-551

| | | | | |
|--|--|--|-----------------------------|---------------------------|
| 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 | | |
| | BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) | | | |
| | BENZYL ALCOHOL (UNII: LKG8494WBH) | | | |
| | SODIUM PHOSPHATE, MONOBASIC, DIHYDRATE (UNII: 5QWK665956) | | | |
| | EDETATE DISODIUM (UNII: 7FLD91C86K) | | | |
| | WATER (UNII: 059QF0KO0R) | | | |
| | POLYETHYLENE GLYCOL 1450 (UNII: OJ4Z5Z32L4) | | | |
| | POVIDONE K30 (UNII: U725QWY32X) | | | |
| Packaging | | | | |
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:65923-551-30 | 1 in 1 CARTON | 11/02/2016 | |
| 1 | | 15 mL in 1 BOTTLE, PUMP; 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 | 06/30/2013 | | |

Labeler - United Exchange Corp (840130579)

Revised: 6/2017

United Exchange Corp