

FAMILY CARE NASAL RELIEF- oxymetazoline hydrochloride spray
United Exchange Corp

Family Care Nasal Spray Severe Congestion 0.5 oz 571 ZDP

Active ingredient

Purpose

Oxymetazoline HCl 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 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 (1-800-222-1222) right away.

Directions

- adults and children 6 to 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. Shake well before use. Remove safety seal. 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 onto bottle.

Other information

- store between 20° and 25°C (66° and 77°F)

Inactive ingredients benzalkonium chloride, benzyl alcohol, camphor, edetate disodium, eucalyptol, menthol, polysorbate 80, propylene glycol, purified water, sodium phosphate dibasic, sodium phosphate monobasic monohydrate

Distributed by:

United Exchange Corp

Cypress, CA 90630 USA

1-888-645-8204

Made in China

FAMILY CARE™
NASAL RELIEF
 Oxymetazoline HCl 0.05%
 (Nasal Decongestant)
SEVERE

Compare to **AFRIN®** Severe Congestion active ingredient*
CHILD SAFETY CAP

FAMILY CARE™
NASAL RELIEF
 Oxymetazoline HCl 0.05%
 (Nasal Decongestant)

Pump Mist Spray

Features:

- Fast & Powerful Congestion Relief
- Maximum Strength with Menthol

SEVERE

UP TO
12
 HOUR RELIEF
0.5 FL OZ (15 ML)



FAMILY CARE™

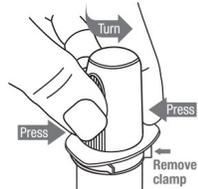
FULL DIRECTIONS IN DRUG FACTS PANEL

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.



Drug Facts (continued)

Other information

- store at room temperature 20-25°C (68-77°F)

Inactive ingredients

benzalkonium chloride, benzyl alcohol, camphor, edetate disodium, eucalyptol, menthol, polysorbate 80, propylene glycol, purified water, sodium phosphate dibasic, sodium phosphate monobasic monohydrate

Questions or comments?

Call 1-888-645-8204
 Monday-Friday 9AM-5PM (PST)

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

TAMPER EVIDENT: DO NOT USE IF PRINTED SEAL AROUND PUMP AND CAP IS BROKEN OR MISSING.

*This product is not manufactured or distributed by Bayer®, distributor of Afrin® Severe Congestion.

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 UNITED EXCHANGE CORP.
 Cypress, CA 90630 USA
 1-888-645-8204

Made In China

571-ZDP-BX-R2
 NAS-MD-ZDP101-R2

Drug Facts

Active ingredient Oxymetazoline hydrochloride 0.05%.....
Purpose Nasal decongestant

Uses

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Warnings

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 - 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.

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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: consult a doctor
- shake well before use. Remove safety seal. 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 onto bottle.

Lot#:

Exp:

FAMILY CARE NASAL RELIEF

oxymetazoline hydrochloride spray

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:65923-571

Route of Administration	NASAL			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	5 g in 100 mL	
Inactive Ingredients				
	Ingredient Name	Strength		
	SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)			
	BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)			
	EUCALYPTOL (UNII: RV6J6604TK)			
	WATER (UNII: 059QF0KO0R)			
	BENZYL ALCOHOL (UNII: LKG8494WBH)			
	CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)			
	POLYSORBATE 80 (UNII: 6OZP39ZG8H)			
	PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
	MENTHOL (UNII: L7T10EIP3A)			
	SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)			
	EDETATE DISODIUM (UNII: 7FLD91C86K)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-571-15	1 in 1 CARTON	05/25/2018	
1		15 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 Drug	M012	05/31/2013		

Labeler - United Exchange Corp (840130579)

Revised: 10/2025

United Exchange Corp