

OXYMETAZOLINE HYDROCHLORIDE- oxymetazoline hydrochloride spray
CHAIN DRUG MARKETING ASSOCIATION INC.

Nasal Decongestant

Active Ingredient

Purpose

Oxymetazoline HCl 0.05%..... Nasal decongestant

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

Warnings

Ask a doctor before use if you have

- heart disease
- thyroid disease
- high blood pressure
- 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 under 12 years of age (with adult supervision):** 2 or 3 sprays in each nostril not more often than every 10-12 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 and turn in a counter-clockwise direction and pull off to remove. To spray, 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 back onto the bottle.

Other information

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

Questions or comments?

1-800-935-2362, Mon-Fri. 9 am-5 pm EST)

Inactive ingredients

- **No Drip Original Nasal Pump Mist- (NDC-63868-605-01)** avicel, benzalkonium chloride, benzyl alcohol, dibasic sodium phosphate, edetate disodium dihydrate, flavor, monobasic sodium phosphate, polyethylene glycol, povidone, purified water
- **No Drip Extra Moisturizing Nasal Pump Mist- (NDC-63868-676-01)** avicel, benzalkonium chloride, benzyl alcohol, dibasic sodium phosphate, edetate disodium dihydrate, glycerin, monobasic sodium phosphate, polyethylene glycol, povidone, purified water
- Tamper Evident: Do not use the product if the tamper evident seal is broken or missing.
- This product is not manufactured or distributed by Bayer HealthCare LLC distributor of Afrin Original Nasal Spray.

Distributed by C.D.M.A Inc.

43157 W 9 Mile Rd

Novi, MI 48375

www.qualitychoice.com

Question: 800935-2362

PRINCIPAL DISPLAY PANEL

NDC-63868-605-01

COATING
FREE AREA

Drug Facts

Active ingredient Oxymetazoline HCl 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
 - trouble urinating due to an enlarged prostate gland
 - thyroid disease
 - diabetes

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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Drug Facts (continued)

Other information

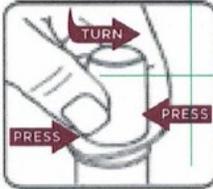
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Questions or comments?

1-800-935-2362 (Mon-Fri 9am-5pm EST)



Tamper Evident: Do not use the product if the tamper evident seal is broken or missing.

*This product is not manufactured or distributed by Bayer HealthCare LLC distributor of Afrin® No Drip Original.



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No Drip Original

Nasal Pump Mist



NDC 63868-605-01

*Compare to Active Ingredient in Afrin® No Drip Original

No Drip Original

Nasal Pump Mist

Oxymetazoline hydrochloride 0.05%
Nasal Decongestant

Relieves:

- ✓Allergy Symptoms
- ✓Common Cold
- ✓Congestion



1 fl.oz. (30 mL)



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Lot #

Exp.

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COATING
FREE AREA

NDC-63868-676-01



OXYMETAZOLINE HYDROCHLORIDE

oxymetazoline hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-605
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.05 g in 100 mL
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Inactive Ingredients

Ingredient Name	Strength
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
WATER (UNII: 059QF0KO0R)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-605-01	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	02/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	02/01/2021	

OXYMETAZOLINE HYDROCHLORIDE

oxymetazoline hydrochloride spray

Product Information

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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
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
glycerin (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
WATER (UNII: 059QF0K0OR)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	

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Packaging

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1	NDC:63868-676-01	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	02/01/2021	

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OTC Monograph Drug	M012	02/01/2021	

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Labeler - CHAIN DRUG MARKETING ASSOCIATION INC. (011920774)

Registrant - Seaway Pharma Inc. (117218785)

Establishment

Name	Address	ID/FEI	Business Operations
117218785		117218785	manufacture(63868-605, 63868-676)

Revised: 12/2024

CHAIN DRUG MARKETING ASSOCIATION INC.