

NASAL DECONGESTANT- oxymetazoline hcl spray, metered Proficient Rx LP

Major Pharmaceuticals Nasal Decongestant 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
- 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: Shake well before use. Hold white tabs, SQUEEZE grooved area of cap FIRMLY and turn counter clockwise. Before using the first time, prime metered pump by depressing pump firmly several times. To spray, hold bottle with thumb at base and nozzle between first and second fingers. Without tilting head, insert nozzle into nostril. Fully depress rim with a firm, even stroke and sniff deeply. 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, microcrystalline cellulose and carboxymethylcellulose sodium, monobasic sodium phosphate, polyethylene glycol, povidone, purified water

Questions or comments?

1-800-616-2471

Principal Display Panel

Soothing - 12 Hour

NASAL DECONGESTANT Spray

Original

Oxymetazoline hydrochloride 0.05%

Compare to active ingredient of Afrin® No Drip

1 FL. OZ. (30 mL)

NDC 71205-219-30

Relabeled by:

Proficient Rx LP

Thousand Oaks, CA 91320



Scan Here



NDC 71205-219-30

Relabeled By: Proficient Rx LP
Thousand Oaks, CA 91320

Nasal Decongestant 0.05%
1 FL. OZ. (30 mL) Spray
Lot #:00000 SN# MASTER
NDC 71205-219-30 Exp:00/00/00

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NDC 71205-219-30 Exp:00/00/00



Nasal Decongestant 0.05%

1 FL. OZ. (30 mL) Spray

Each bottle contains: Oxymetazoline hydrochloride 0.05% Nasal decongestant

See Bottle

Product ID: SN021930

Dist. By: Major Pharmaceuticals 17177 N Laurel Park Drive, Suite 233 Livonia, MI 48152

Store at 20°-25°C (68°-77°F)

Keep medication out of the reach of children



GTIN: 00371205219307
SN# MASTER
Exp. 00/00/00
Lot #:00000

NASAL DECONGESTANT

oxymetazoline hcl spray, metered

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71205-219(NDC:0904-6761)
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)	
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JH2SW)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Product Characteristics

Color	WHITE (to off white, viscous)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71205-219-30	1 in 1 CARTON	02/01/2019	03/31/2026
1		30 mL in 1 BOTTLE, SPRAY; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/12/2018	03/31/2026

Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(71205-219) , RELABEL(71205-219)

Revised: 1/2026

Proficient Rx LP