OXYMETHAZOLINE HCL- oxymethazoline hcl spray Proficient Rx LP

12 Hour Decongestant Nasal Spray

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

Ask a doctor before use if you have

heart disease

high blood pressure

thyroid disease

diabetes

trouble urinating due to an enlarged prostate gland

Stop use and ask a doctor if symptoms presist

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 doeses in 24 hours.

children under 6 years of age: ask a doctor.

Instructions for use: Shake well before use. to open, rotate cap to align the marks. Squeeze cap on toh sides in a counter-clockwise turn and pull to remove. To spray, hold bottles 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° to 25° C (68° to 77° F) retain carton for future referance on full labeling

Inactive ingredients

benzalkoium cjloride, dibasic sodium phosphate, edetate disodium dihydrate, monobasic sodium phosphate, polyethylene glycol, propylene glycol, povidone, purified water

Questions or comments?

call **516-341-0666**, 8:30 am - 4:30 pm ET, Monday - Friday

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 prduct

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, sneexing, or an increased nasal discharge may occur

use of this container by more than one person may spread infection

NDC 71205-383-30

Reliabel-1 Laboratories

12 Hour Decongestant Nasal Spray

Oxymetazoline HCL

Pump Mist Anti-Drip

Rapid & Powerful

Congestion Relief

12 Hour Relief

1 FL OZ (30 mL)





NDC 71205-383-30

Relabeled By: Proficient Rx LP Thousand Oaks, CA 91320

Nasal Decongestant 0.05%

1 FL OZ (30 mL) Spray
Lot #:00000 SN# MASTER
NDC 71205-383-30 Exp:00/00/00

Nasal Decongestant 0.05%
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Nasal Decongestant 0.05%
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NDC 71205-383-30 Exp:00/00/00



GTIN: 00371205383305 SN# MASTER Exp. 00/00/00 Lot #:00000

Nasal Decongestant 0.05%

1 FL OZ (30 mL) Spray

Each bottle contains: Oxymetazoline HCI 0.05% Nasal decongestant

See Bottle

Product ID: SN038330

Dist. By: Reliable-1 Laboratories LLC Valley Stream, NY 11580

Store between 20°-25°C (68°-77°F)

Keep medication out of the reach of children

OXYMETHAZOLINE HCL

oxymethazoline hcl spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71205-383(NDC:69618-050)

Route of Administration NASAL

Active Ingredient/Active Moiety

	Ingredient Name	Basis of Strength	Strength
l	OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZ OLINE HYDROCHLORIDE	0.05 mg

Inactive Ingredients Ingredient Name Sodium Phosphate, Monobasic, Anhydrous (UNII: KH7I04HPUU) EDETATE SODIUM (UNII: MP1J8420LU) PROPYLENE GLYCOL (UNII: 6DC9Q167V3)

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
WATER (UNII: 059QF0KO0R)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	

Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:71205-383-30	1 in 1 CARTON	01/10/2019	
	1	1 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	01/01/2019		

Labeler - Proficient Rx LP (079196022)

Establishment			
Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(71205-383), RELABEL(71205-383)

Revised: 1/2024 Proficient Rx LP