

MAJOR NASAL DECONGESTANT- oxymetazoline hydrochloride spray
Preferred Pharmaceuticals Inc.

Major Nasal Decongestant Spray No Drip

Drug facts

Active Ingredient

Oxymetazoline 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 membranes so you can breathe more freely

Warnings

Ask a Doctor before use if you have

- heart disease
- high blood pressure
- diabetes
- thyroid disease
- trouble urinating due to enlarged prostate gland.

When using this product

Do not use more than directed

- do not use for more than three 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 the reach of children

if swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Before using for the first time, remove the protective cap from the tip and prime the 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 and even stroke, and sniff deeply.
- Wipe nozzle clean after use.
- Adults and children 6 to under 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 consult a doctor.

Other information

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

Inactive ingredients

Benzalkonium Chloride, Disodium EDTA, Glycerin, Microcrystalline Cellulose and Carboxymethylcellulose Sodium, Polyethylene Glycol 300, Povidone, Propylene Glycol, Purified Water, Sodium Phosphate Dibasic, Sodium Phosphate Monobasic

Questions? Call 800-222-1807

Principal Display Panel

Major ® No Drip Nasal Mist NDC 68788-8744-3
Nasal Spray Oxymetazoline HCL 0.05%
12 hour Nasal Spray
1 FL OZ (30 mL)

Preferred Pharmaceuticals Inc.

Nasal Decongestant 12-Hour Spray



CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed.

Nasal Decongestant 12-Hour Spray
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Active Ingredient Oxymetazoline HCl, 0.05%
Pkg Size: Exp Date: #####/#####
Lot#: Batch#:



Directions English
Take _____ sprays in each nostril every _____ hours
Use as directed by your doctor



GTIN

SN #####
EXP #####

Instrucciones Espanol:
La Toma _____ rocia cada _____ horas
Uso según lo dirigido por su doctor

Nasal Decongestant 12-Hour Spray
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Nasal Decongestant 12-Hour Spray
Qty:
Insurance NDC:
Lot: Bat:

Nasal Decongestant 12-Hour Spray
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Ins:
Mfg: Major Pharmaceuticals
Prod#:
Warning
Uses: For the temporary relief of nasal congestion due to the common cold, hay fever, or other upper respiratory allergies; shrinks swollen membranes so you can breathe more freely. Ask a doctor before use if you have heart disease, high blood pressure, thyroid disease, diabetes, trouble urinating due to enlarged prostate gland. When using this product, may cause temporary discomfort such as burning, stinging, sneezing or an increase in nasal discharge; the use of this container by more than one person may spread infection. Frequent or prolonged use may cause nasal congestion to recur or worsen. Stop use and ask a doctor if symptoms persist. See package for additional information. Store between 20°-25°C (68°-77°F).

Log
Chart
Billing
Patient

MAJOR NASAL DECONGESTANT				
oxymetazoline hydrochloride spray				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-8744(NDC:0904-7427)	
Route of Administration	NASAL			
Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.5 mg in 1 mL		
Inactive Ingredients				
Ingredient Name	Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
GLYCERIN (UNII: PDC6A3C0OX)				
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JH2SW)				
CROSPVIDONE, UNSPECIFIED (UNII: 2S7830E561)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8744-3	1 in 1 CARTON	09/26/2024	
1		30 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 Drug	M012	09/26/2024	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment			
Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	RELABEL(68788-8744)

Revised: 8/2025

Preferred Pharmaceuticals Inc.