

NASAL- oxymetazoline hydrochloride spray
NuCare Pharmaceuticals, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

MAJOR®

Soothing - 12 Hour

NASAL

DECONGESTANT

Spray

Drug Facts

Active ingredient

Oxymetazoline hydrochloride 0.05%

Purpose

Nasal decongestant

Uses

- for the temporary relief of nasal congestion due to the common cold, hay fever or other 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
- thyroid disease
- diabetes
- trouble urinating due to 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.
- temporary discomfort such as burning, stinging, sneezing, or an increase in nasal discharge may occur

- 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 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 to 12 hours
- do not exceed 2 doses within any 24-hour period
- children under 6 years of age: consult a doctor

Before using the first time, remove the protective cap from the tip. To spray, hold bottle between the thumb and first and second fingers. Without tilting head, insert nozzle into nostril. Squeeze the bottle quickly and firmly. Wipe nozzle clean after use.

Other information

- store between 20° to 25° C (68° to 77° F)

Inactive ingredients

benzalkonium chloride, benzyl alcohol, dibasic sodium phosphate, edetate disodium, hydrochloric acid, monobasic sodium phosphate, polyethylene glycol 1450, povidone, propylene glycol, purified water

Questions?

To Report Adverse Drug Event Call: (800) 616-2471

Distributed by:

MAJOR[®] PHARMACEUTICALS
17177 N Laurel Park Drive, Suite 233
Livonia, MI 48152 USA

PRINCIPAL DISPLAY PANEL -

NuCare Pharmaceuticals, Inc.

Use only as directed
by your physician.

Patient Instructions:
Call your doctor for medical advice about side effects.
You may report side effects to FDA at 1-800-FDA-1088.

Distributed by:
Major Pharmaceuticals, Livonia, MI 48152
Packaged by:
NuCare Pharmaceuticals, Inc.
Orange, CA 92867

NDC 68071-4641-1
Lot #: 000000 Exp. Date: 00-00

Nasal Decongestant
1/2oz Spray

See manufacturer's label
for full list of ingredients

Product #: R0400015PED

Nasal Decongestant
1/2oz Spray Exp Date: 00-00
NDC 68071-4641-01 AWP:
Mfg NDC 0904-5711-35
Lot #: 000000 Rx # 23219241

Nasal Decongestant
1/2oz Spray Exp Date: 00-00
NDC 68071-4641-01 AWP:
Mfg NDC 0904-5711-35
Lot #: 000000 Rx # 23219241

Nasal Decongestant
1/2oz Spray Exp Date: 00-00
NDC 68071-4641-01 AWP:
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Lot #: 000000 Rx # 23219241



R0400015PED

Rx # 23219241

WARNING: KEEP OUT OF REACH OF CHILDREN. STORE AT CONTROLLED TEMPERATURE 68-77 °F.

NASAL

oxymetazoline hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-4641(NDC:0904-5711)
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
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JH2SW)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
POLYETHYLENE GLYCOL 1450 (UNII: OJ4Z5Z32L4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:68071-4641-1	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/29/2018	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph final	part341		03/17/2009	

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-4641)

Revised: 2/2021

NuCare Pharmaceuticals, Inc.