MEDIX NASAL DECONGESTANT OXYMETAZOLINE HYDROCHLORIDE- oxymetazoline hydrochloride liquid
Medic -33, Inc.

Medix Nasal Decongestant Oxymetazoline Hydrochloride

Drug Facts

Active ingredient

Oxymetazoline hydrochloride 0.05%

Purpose

nasal decongestant

Uses

- for the temporary relief of nasal congestion
- temporarily restores freer breathing through the nose

Warnings

Do not use

- this product for more than 3 days. Use only as directed. Frequent or prolonged use may cause nasal congestion to recur or worsen. If symptoms persist, consult a doctor.

Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- difficulty in urination due to enlargement of the prostate gland

When using this product

- do not exceed recommended dosage.
- 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.

If pregnant or breastfeeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact 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 in any 24-hour period.
- Children under 6 years of age: consult a doctor

Other information

Do not use if the tamper-proof seal is damaged or broken.

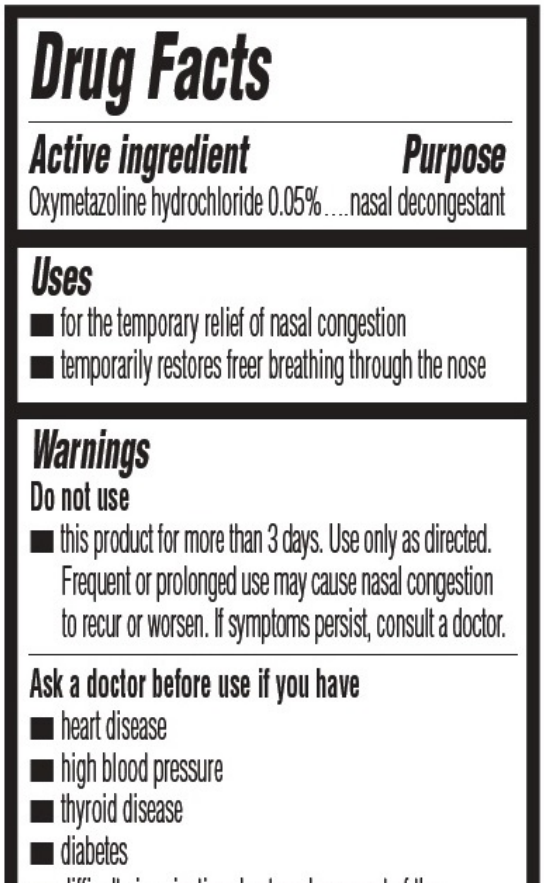
Inactive ingredients

Glutathione, menthol oil, pyrroloquinoline quinine, sodium nitrate, wintergreen oil, water.

Questions or Comments?

305-861-1457

Package Labeling:



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DISTRIBUTED BY:
MEDIC -33, INC.
10185 COLLINS AVE.
BAL HARBOUR, FL 33154

MEDIX NASAL DECONGESTANT OXYMETAZOLINE HYDROCHLORIDE

oxymetazoline hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83522-001
Route of Administration	TOPICAL		

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
GLUTATHIONE (UNII: GAN16C9B8O)	
PYRROLOQUINOLINEDIONE TRICARBOXYLIC ACID (UNII: 47819QGH5L)	
SODIUM NITRATE (UNII: 8M4L3H2ZVZ)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83522-001-01	1 in 1 CARTON	09/10/2023	
1		120 mL in 1 BOTTLE; 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	09/10/2023	

Labeler - Medic -33, Inc. (116663992)

Revised: 9/2024 Medic -33, Inc.