

METARIVIN SOLN. 0.1%- xylometazoline hydrochloride liquid
Lydia Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Xylometazoline Hydrochloride

■ relieve your cold symptoms cold Rhinitis

Keep out of reach of children

Adults and children 7 years and older: Spray into the nose 3 times a day

■ When using this product Stop use and ask a doctor if nervousness, dizziness, or sleepiness occur. Pain, nasal congestion or cough gets worse or lasts more than 7 days. fever gets worse or lasts more than 3 days New symptoms occur. keep out of reach of children. In case of overdose, get medical help or contact a poison control center right away. If pregnant or breast-feeding, ask a health professional before use.

l-menthol, eucalyptus oil, etc

for nasal use

Metarivin Soln. 0.1%

Drug Facts

Active Ingredients (in 1g)

Xylometazoline Hydrochloride 1mg/mL ----- Purpose
Topical analgesic

Uses

- relieve your cold symptoms
- cold • Rhinitis

Warnings

■ When using this product
Stop use and ask a doctor if nervousness, dizziness, or sleepiness occur. Pain, nasal congestion or cough gets worse or lasts more than 7 days.

fever gets worse or lasts more than 3 days

New symptoms occur.

keep out of reach of children. In case of overdose, get medical help or contact a poison control center right away.

If pregnant or breast-feeding, ask a health professional before use.

Directions

■ Adults and children 7 years and older: Spray into the nose 3 times a day.

Other Information

■ Store below 25°C

Inactive Ingredient

l-menthol, Sodium edetate hydrate, eucalyptus oil

Questions or comments ?

Call weekdays from 9 a.m to 5 p.m EST at 562-273-0692

Distributed By: Bio Mission Group, Inc

9925 Painter Ave Suite #0 Whittier, CA 90605 USA

Made in South Korea

NDC Number:

METARIVIN SOLN. 0.1%

xylometazoline hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72988-0030
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Route of Administration	NASAL
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Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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XYLOMETAZOLINE HYDROCHLORIDE (UNII: X5S84033NZ) (XYLOMETAZOLINE - UNII: WPY40FTH8K)	XYLOMETAZOLINE HYDROCHLORIDE	1 mg in 1 mL
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Inactive Ingredients

Ingredient Name	Strength
COBALT DISODIUM EDETATE (UNII: 3EY1Y2QRLI)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72988-0030-1	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/30/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/30/2022	

Labeler - Lydia Co., Ltd. (695735569)

Registrant - Lydia Co., Ltd. (695735569)

Establishment

Name	Address	ID/FEI	Business Operations
Lydia Co., Ltd.		695735569	manufacture(72988-0030)

Revised: 9/2022

Lydia Co., Ltd.