COLD AND FLU RELIEF- atropine, naja naja venom, magnesium chloride, potassium hydroxide spray PROTEGE' MEDIA LLC

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.

Active ingredient

Purpose

Contains atropine (Atropinum) 5X HPUS runny nose relief Contains cobra venom preparation (Naja naja) 5X HPUS pain relief Contains magnesium chloride (Magnesia muriatica) 5X HPUS cold remedy Contains potassium hydroxide (Kali causticum) 7X HPUS flu remedy Reference: the Homeopathic Pharmacopoeia of the United States (HPUS)

Contains 24 doses

Use

Temporarily relieves symptoms associated with colds and flu.

Warnings

If symptoms persist or worsen, discontinue use, seek medical attention.

- Avoid contact with eyes. If product gets into eyes, flush with water, seek medical attention.
- If pregnant or breastfeeding ask a health professional before use.
- Consult a medical professional if using other medications for known interactions.
- The use of this dispenser by more than one person may spread infection.

Keep out of reach of children.

Directions

- Do not use if tamperproof cover is missing.
- Press down 2-3 times to prime the pump.
- Spray once into each nostril
- Use 2 times per day to relieve discomfort.

Inactive ingredients

Aspartic acid, benzalkonium chloride, glycerin, propylene glycol, purified water, thiamine.

Comments, Questions or Complaints?

Call 888-735-4009 or info@promedxnow.com

Product label



COLD AND FLU RELIEF

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ATDODINE (LINII: 7C0697DD9D (ATDODINE - LINII:7C0697DD9D

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (So	urce)	NDC:765	524-501
Route of Administration	NASAL				
Active Ingredient/Active Moiety					
Ingredient Name		Basis of Stren	gth	Strength	

5 [hp_X]

ATDODING.

ATROFINE (UNII, /G003/DR31) (ATROFINE - UNII,/G003/DR31)	AIROFINE	in 0.2 mL
NAJA NAJA VENOM (UNII: ZZ4AG7L7VM) (NAJA NAJA VENOM - UNII:ZZ4AG7L7VM)	NAJA NAJA VENOM	5 [hp_X] in 0.2 mL
MAGNESIUM CHLO RIDE (UNII: 02F3473H9O) (MAGNESIUM CATION - UNII:T6 V3LHY838)	MAGNESIUM CATION	5 [hp_X] in 0.2 mL
PO TASSIUM HYDRO XIDE (UNII: WZH3C48 M4T) (HYDRO XIDE ION - UNII:9 159 UV38 1P)	POTASSIUM HYDROXIDE	7 [hp_X] in 0.2 mL

Inactive Ingredients		
Ingredient Name	Strength	
ASPARTIC ACID (UNII: 30 KYC7MIAI)		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
THIAMINE (UNII: X66NSO3N35)		

ı	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:76524-501- 18	5 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/29/2020	

Marketing Infor	rketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		10/29/2020		

Labeler - PROTEGE' MEDIA LLC (117063923)

Revised: 1/2021 PROTEGE' MEDIA LLC