

**MDFARMA COLD AND FLU RELIEF- atropine, naja naja venom, magnesium chloride, potassium hydroxide spray
Green Earth Health Inc.**

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

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.

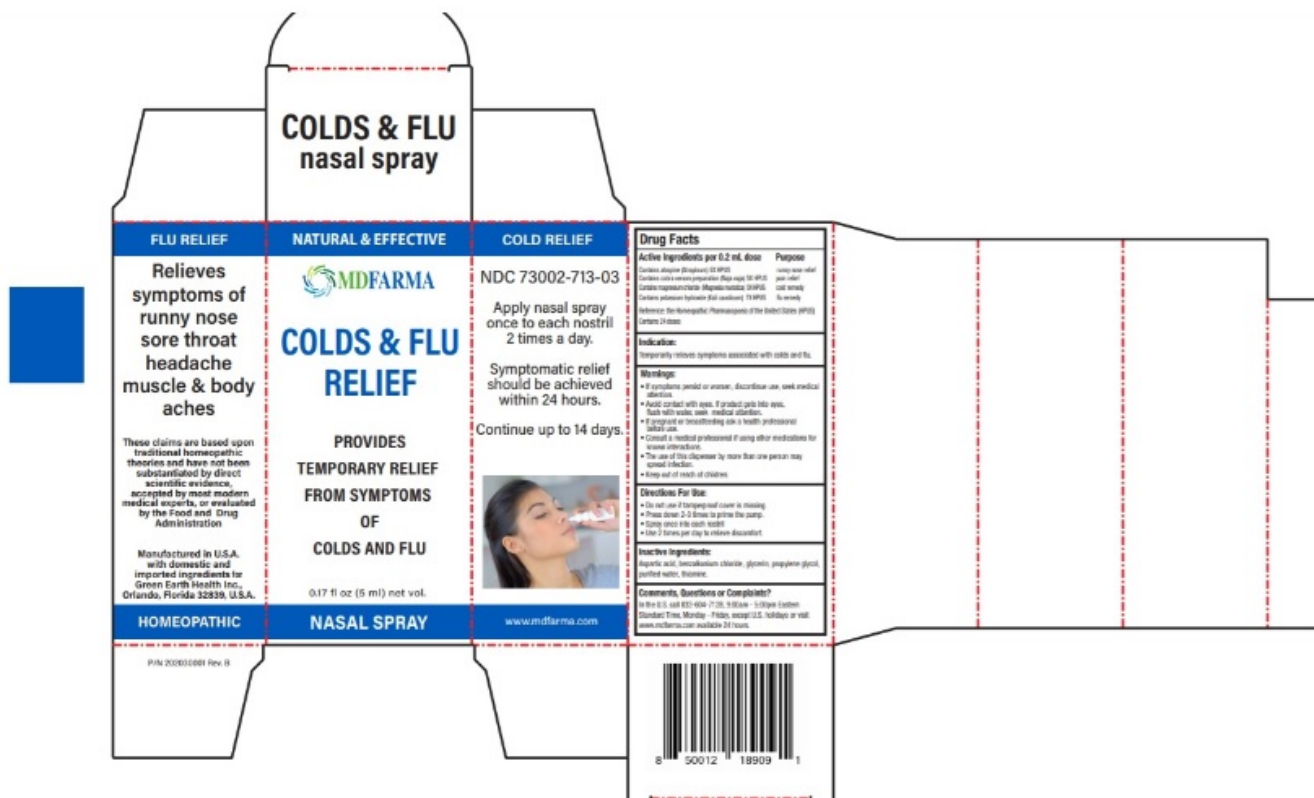
Inactive ingredients

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

Comments, Questions or Complaints?

In the U.S. call 833-604-7128, 9:00am - 5:00pm Eastern Standard Time, Monday - Friday, except U.S. holidays or visit www.mdfarma.com available 24 hours.

Product label



MDFARMA COLD AND FLU RELIEF

atropine, naja naja venom, magnesium chloride, potassium hydroxide spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73002-713
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ATROPINE (UNII: 7C0697DR9I) (ATROPINE - UNII:7C0697DR9I)	ATROPINE	5 [hp_X] 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 CHLORIDE (UNII: 02F3473H9O) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM CATION	5 [hp_X] in 0.2 mL
POTASSIUM HYDROXIDE (UNII: WZ H3C48M4T) (HYDROXIDE ION - UNII:9159UV381P)	POTASSIUM HYDROXIDE	7 [hp_X] in 0.2 mL

Inactive Ingredients

Ingredient Name	Strength
ASPARTIC ACID (UNII: 30KYC7MIAI)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
THIAMINE (UNII: X66NSO3N35)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73002-713-03	5 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/23/2020	

Labeler - Green Earth Health Inc. (116983264)

Registrant - Green Earth Health Inc. (116983264)

Revised: 12/2023

Green Earth Health Inc.