ECZEMA RELIEF- arsenicum iodatum, kali bromatum, kali muriaticum, natrum bromatum, niccolum sulphuricum, sulphur iodatum, viola tricolor, zincum bromatum. liquid
OHM PHARMA INC.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

OHM Eczema Relief

ACTIVE INGREDIENTS (HPUS*): Equal amounts of : Arsenicum Iodatum 8X, Kali Bromatum 4X, Kali Muriaticum 4X, Natrum Bromatum 4X, Niccolum Sulphuricum 5X, Sulphur Iodatum 14X, Viola Tricolor 12X, Zincum Bromatum 6X.

* The letters "HPUS" indicate that the components in this product are officially monographed in the Homeopathic Pharmacopeia of the United States.

USES: Temporarily relieves eczema - psoriatic skin symptoms.**

**This statement has not been evaluated by the FDA. It is based on documented Homeopathic Materia Medica.

WARNINGS: IF PREGNANT OR BREAST-FEEDING, ask a health care professional before use. If symptoms worsen or persist for more than a week, discontinue use and contact a doctor.

• KEEP OUT OF REACH OF CHILDREN.

DIRECTIONS: Shake 10 times before use. Adults & children above 12 years of age: spray 3-4 times under the tongue 2-3 times a day, or as directed by a health care professional. Children under 12: use half the adult dose.

Do not use if cap seal is broken.

INACTIVE INGREDIENTS: Ethanol Alcohol 15% (Non-GMO, Kosher Certified, Corn, Gluten, Allergen Free), Purified Water, Vegetable Glycerin 15%.

Mfg. By: OHM PHARMA, INC. Mineral Wells, TX 76067 www.ohmpharma.com

HOMEOPATHIC MEDICINE

NDC: 66096-192-05

ECZEMA Relief

Natural - Safe - Effective

No Side Effects

2 fl oz (59 mL) / 15% Alcohol

PRODUCT OF USA



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Your Partner in Homeopathy

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PHARMACEUTICAL GRADE

PROFESSIONAL USE ONLY

PRODUCT OF USA

Temporarily relieves eczema.

ECZEMA RELIEF

arsenicum iodatum, kali bromatum, kali muriaticum, natrum bromatum, niccolum sulphuricum, sulphur iodatum, viola tricolor, zincum bromatum. liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:66096-192 Route of Administration ORAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ARSENIC TRIIODIDE (UNII: 302998802T) (ARSENIC CATION (3+) - UNII: C96613F5AV)	ARSENIC TRIIODIDE	8 [hp_X] in 59 mL
POTASSIUM BROMIDE (UNII: OSD78555ZM) (BROMIDE ION - UNII:952902IX06)	POTASSIUM BROMIDE	4 [hp_X] in 59 mL
POTASSIUM CHLORIDE (UNII: 660YQ98I10) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM CHLORIDE	4 [hp_X] in 59 mL
SODIUM BROMIDE (UNII: LC1V549NOM) (BROMIDE ION - UNII:952902IX06)	SODIUM BROMIDE	4 [hp_X] in 59 mL
NICKEL SULFATE HEXAHYDRATE (UNII: JC9WZ4FK68) (NICKEL CATION - UNII: OIS2CXW7AM)	NICKEL SULFATE HEXAHYDRATE	5 [hp_X] in 59 mL

SULFUR IODIDE (UNII: L6L8KA2AA0) (SULFUR IODIDE - UNII:L6L8KA2AA0)	SULFUR IODIDE	14 [hp_X] in 59 mL
VIOLA TRICOLOR (UNII: 9Q24RAI43V) (VIOLA TRICOLOR - UNII:9Q24RAI43V)	VIOLA TRICOLOR	12 [hp_X] in 59 mL
ZINC BROMIDE (UNII: 007ZBU9703) (ZINC CATION - UNII:13S1S8SF37)	ZINC BROMIDE	6 [hp_X] in 59 mL

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		

ı	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:66096-192- 05	59 mL in 1 PACKAGE; Type 0: Not a Combination Product	05/22/2015	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/22/2015	
потпеораинс			

Labeler - OHM PHARMA INC. (030572478)

Revised: 12/2021 OHM PHARMA INC.