KALI IODATUM 1X- kali iodatum. 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 Kali Iodatum 1X

ACTIVE INGREDIENT (HPUS*): Kali Iodatum 1X.

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

USES: Temporarily relieves dysfunctional thyroid metabolism.**

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

WARNINGS: THIS PRODUCT CONTAINS IODIDE. DO NOT TAKE IF YOU ARE HIPERSENSITIVE OR ALLERGIC TO IODIDE. IF YOU ARE 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 over 12 years of age: add 1 drop (3.83 mg iodide and 1.18 mg Potassium) to 1 oz of purified water. Take 1 times per day or as directed by a health care professional. Children under 12: contact a doctor.

Do not use if cap seal is broken.

INACTIVE INGREDIENTS: Ethanol Alcohol 5% (Non-GMO, Kosher Certified, Corn, Gluten Allergen Free), Glycerin, Purified Water.

Mfg. By: OHM Pharma, INC. Mineral Wells, TX 76067

www.ohmpharma.com

HOMEOPATHIC MEDICINE

NDC: 66096-166-02

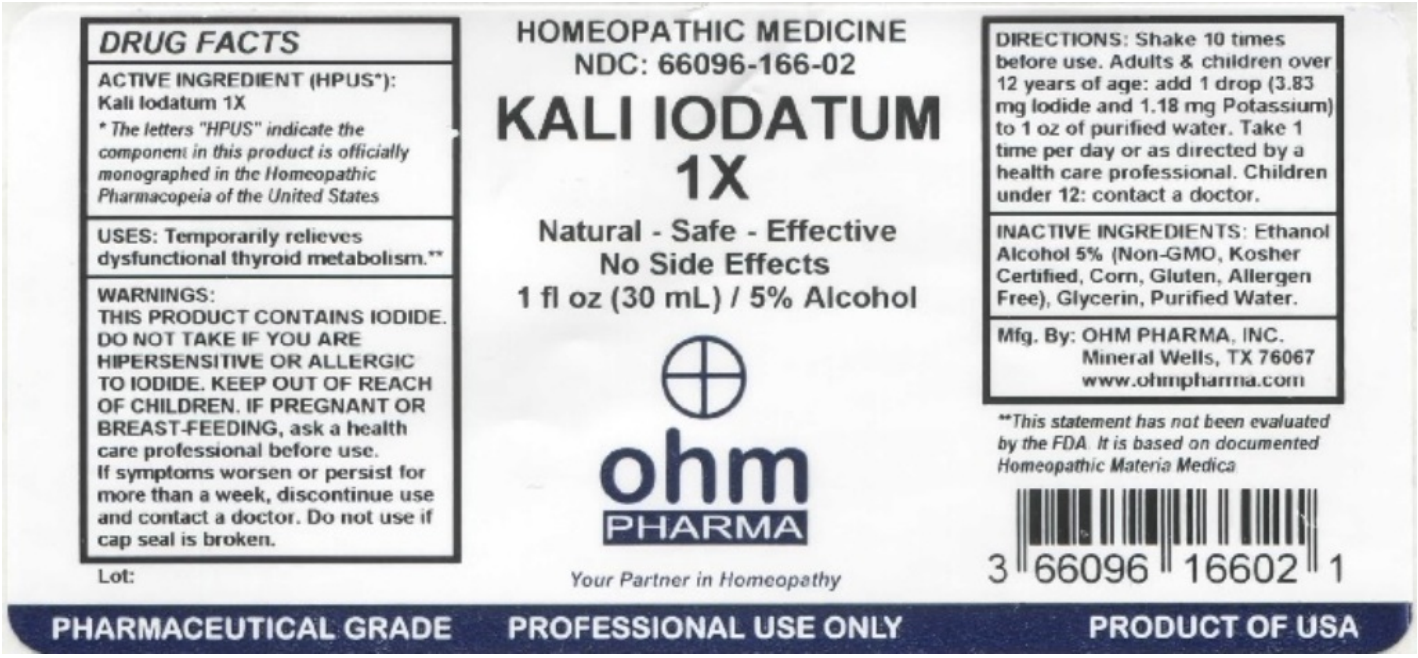
KALI IODATUM 1X

Natural - Safe - Effective

No Side Effects

1 fl oz (30mL) / 5% Alcohol

PRODUCT OF USA



Temporarily relieves dysfunctional thyroid metabolism.

KALI IODATUM 1X

kali iodatum. liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66096-166
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POTASSIUM IODIDE (UNII: 1C4QK22F9J) (IODIDE ION - UNII:09G4I6V86Q)	IODIDE ION	1 [hp_X] in 30 mL

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

Packaging				
			Marketing Start	**Marketing End**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66096-166-02	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	02/10/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		02/10/2016	

Labeler - OHM PHARMA INC. (030572478)

Revised: 12/2021 OHM PHARMA INC.