URTICA U- urtica u liquid
Rxhomeo Private Limited d.b.a. Rxhomeo, 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.

ACTIVE INGREDIENT

URTICA U HPUS 1X and higher

USES

USES: Temporary Relief - Minor Burns And Sun Burn*

* Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

INDICATIONS

Condition listed above or as directed by the physician

DOSAGE

Adults: Mix 4-6 drops in 1/2 cup of water, take orally three times daily or as directed by a physician. Children (2 years and older): Take half the adult dose.

WARNINGS

This product is to be used for self-limiting conditions

If symptoms do not improve in 4 days, or worsen, discontinue use and seek assistance of health professional

As with any drug, if you are pregnant, or nursing a baby, seek professional advice before taking this product

Keep this and all medication out of reach of children

Do not use if capseal is broken or missing.

Close the cap tightly after use.

INACTIVE INGREDIENTS

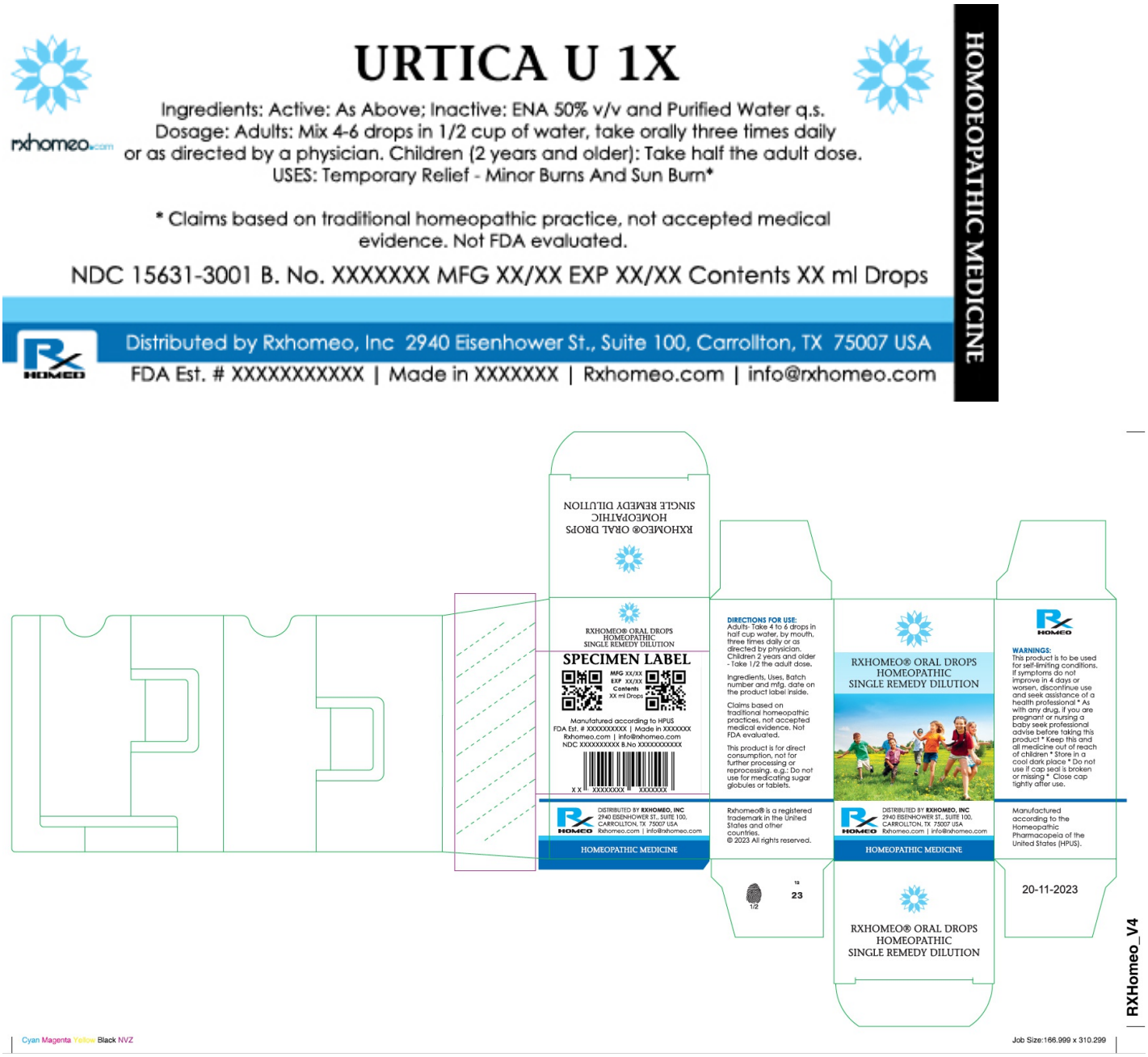
Active: As Above; Inactive: ENA 50% v/v and Purified Water q.s.

STORAGE

Store in a cool dark place

QUESTIONS OR COMMENTS

www.Rxhomeo.com | 1.888.2796642 | info@rxhomeo.com - Rxhomeo, Inc 2940 Eisenhower St., Suite 100, Carrollton, TX 75007 USA



URTICA U

urtica u liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:15631-3001
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
URTICA URENS (UNII: IHN2NQ5OF9) (URTICA URENS - UNII:IHN2NQ5OF9)	URTICA URENS	1 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:15631-3001-0	10 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/23/2024	
2	NDC:15631-3001-1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/23/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/23/2024	

Labeler - Rxhomeo Private Limited d.b.a. Rxhomeo, Inc (650833994)

Establishment

Name	Address	ID/FEI	Business Operations
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc		650833994	manufacture(15631-3001) , label(15631-3001)

Revised: 6/2024 Rxhomeo Private Limited d.b.a. Rxhomeo, Inc