D-79- urtica urens, toxicodendron pubescens leaf, and echinacea angustifolia solution
DNA Labs, 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.

D-79

NDC 58264-0080-1

INDICATIONS

Allergy to poison oak or ivy.

INGREDIENTS

ACTIVE

Urtica urens 3x, Rhus toxicodendron 3x, Echinacea angustifolia 2x

INACTIVE

20% alcohol in purified water.

SUGGESTED DOSAGE

One dropper under tongue two times daily. Acute symptoms ½ dropper under tongue every 30 minutes for two hours.

SHAKE WELL

Warnings

- Use only if cap seal is unbroken.
- If pregnant or breastfeeding, ask a healthcare professional before use.
- Keep this and all medication out of the reach of children.

To be used according to standard homeopathic indications.

PRINCIPAL DISPLAY PANEL - 1 FL. OZ. Bottle Label

DYNAMIC
NUTRITIONAL
ASSOCIATES, INC.

D 79

HOMEOPATHIC
STRESS FORMULA

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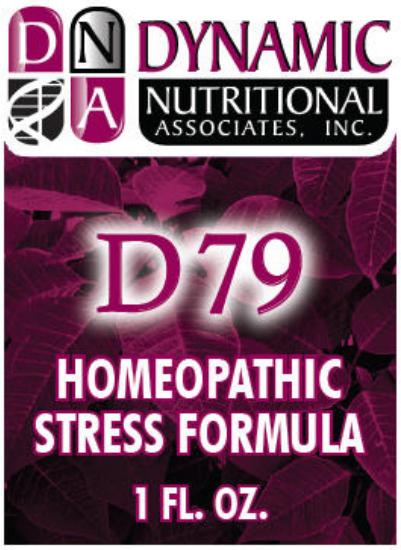
Rev. 3/18



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MANUFACTURED FOR:
DNA LABORATORIES, INC.
Coeur d'Alene, ID 83814

D-79
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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58264-0080
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
URTICA URENS (UNII: IHN2NQ5OF9) (URTICA URENS - UNII:IHN2NQ5OF9)	URTICA URENS	3 [hp_X] in 1 mL
TOXICODENDRON PUBESCENS LEAF (UNII: 6IO182RP7A) (TOXICODENDRON PUBESCENS LEAF - UNII:6IO182RP7A)	TOXICODENDRON PUBESCENS LEAF	3 [hp_X] in 1 mL
ECHINACEA ANGUSTIFOLIA (UNII: VB06AV5US8) (ECHINACEA ANGUSTIFOLIA - UNII:VB06AV5US8)	ECHINACEA ANGUSTIFOLIA	2 [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:58264-0080-1	29.57 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	01/01/1990	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/01/1990	

Labeler - DNA Labs, Inc. (031784339)

Revised: 6/2022 DNA Labs, Inc.