

**D-18- berberis vulgaris root bark, lytta vesicatoria, equisetum hyemale, eutrochium purpureum root, and solanum dulcamara stem solution/ drops
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-18

NDC 58264-0018-1

INDICATIONS

Pains in the sacral region.

INGREDIENTS

ACTIVE

Berberis vulgaris 4x, Cantharis 4x, Equisetum hyemale 6x, Eupatorium purpureum 3x, Dulcamara 4x,

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 18

HOMEOPATHIC STRESS FORMULA

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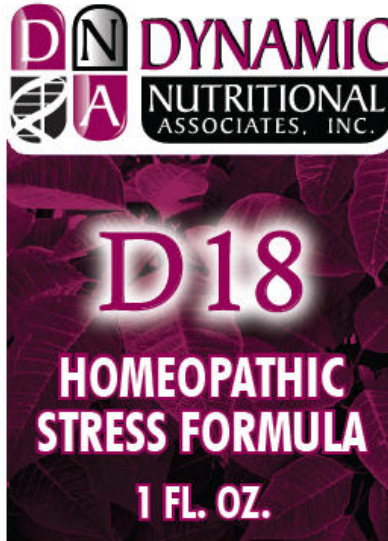
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Rev. 3/18



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

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0018
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	4 [hp_X] in 1 mL
LYTTA VESICATORIA (UNII: 3Q034RO3BT) (LYTTA VESICATORIA - UNII:3Q034RO3BT)	LYTTA VESICATORIA	4 [hp_X] in 1 mL
EQUISETUM HYEMALE (UNII: 59677RXH25) (EQUISETUM HYEMALE - UNII:59677RXH25)	EQUISETUM HYEMALE	6 [hp_X] in 1 mL
EUTROCHIUM PURPUREUM ROOT (UNII: 0VYG5PVN5Q) (EUTROCHIUM PURPUREUM ROOT - UNII:0VYG5PVN5Q)	EUTROCHIUM PURPUREUM ROOT	3 [hp_X] in 1 mL
SOLANUM DULCAMARA STEM (UNII: IR986LE7DF) (SOLANUM DULCAMARA STEM - UNII:IR986LE7DF)	SOLANUM DULCAMARA STEM	4 [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:58264-0018-1	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.