D-51- atropa belladonna, iodine, lycopus virginicus, sodium chloride, hekla lava, and calcium hexafluorosilicate 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-51

NDC 58264-0051-1

INDICATIONS

Thyroid health.

INGREDIENTS

ACTIVE

Belladonna 30x, Iodium 30x, Lycopus virginicus 12x, Natrum muriaticum 30x, Hekla lava 12x, Lapis albus 12x

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 51

HOMEOPATHIC
STRESS FORMULA

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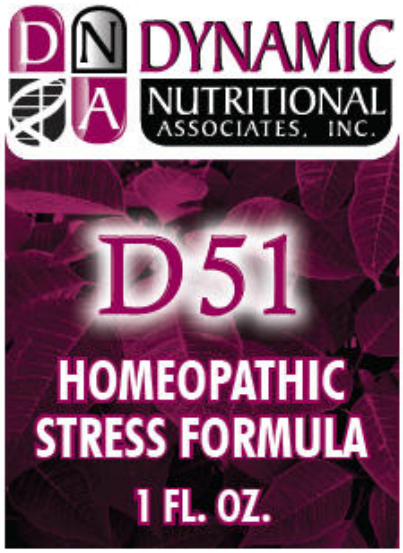
Rev. 3/18



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

D-51

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Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58264-0051
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ATROPA BELLADONNA (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA - UNII:WQZ3G9PF0H)	ATROPA BELLADONNA	30 [hp_X] in 1 mL
IODINE (UNII: 9679TC07X4) (IODINE - UNII:9679TC07X4)	IODINE	30 [hp_X] in 1 mL
LYCOPUS VIRGINICUS (UNII: TWH5125Q6F) (LYCOPUS VIRGINICUS - UNII:TWH5125Q6F)	LYCOPUS VIRGINICUS	12 [hp_X] in 1 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698, SODIUM CATION - UNII:LYR4M0NH37)	SODIUM CHLORIDE	30 [hp_X] in 1 mL
HEKLA LAVA (UNII: C21158IIRK) (HEKLA LAVA - UNII:C21158IIRK)	HEKLA LAVA	12 [hp_X] in 1 mL
CALCIUM HEXAFLUOROSILICATE (UNII: 2NVP93XVQ3) (CALCIUM HEXAFLUOROSILICATE - UNII:2NVP93XVQ3)	CALCIUM HEXAFLUOROSILICATE	12 [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:58264-0051-1	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.