

**D-65- arsenic trioxide, berberis aquifolium root bark, oyster shell calcium carbonate, crude, graphite, centella asiatica, and sodium chloride 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-65

NDC 58264-0065-1

INDICATIONS

Dry crusted raised skin rash.

INGREDIENTS

ACTIVE

Arsenicum album 12x, Berberis aquifolium 1x, Calcarea carbonica 30x, Graphites 12x, Hydrocotyle asiatica 3x, Natrum muriaticum 30x

INACTIVE

25% 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 65

HOMEOPATHIC STRESS FORMULA

1 FL. OZ.

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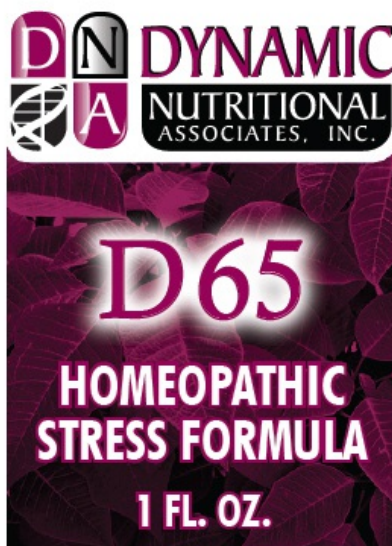
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Rev. 3/18



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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:58264-0065
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARSENIC TRIOXIDE (UNII: S7V92P67HO) (ARSENIC CATION (3+) - UNII:C96613F5AV)	ARSENIC TRIOXIDE	12 [hp_X] in 1 mL
BERBERIS AQUIFOLIUM ROOT BARK (UNII: VZJ9F3C3SB) (BERBERIS AQUIFOLIUM ROOT BARK - UNII:VZJ9F3C3SB)	BERBERIS AQUIFOLIUM ROOT BARK	1 [hp_X] in 1 mL
OYSTER SHELL CALCIUM CARBONATE, CRUDE (UNII: 2E32821G6I) (OYSTER SHELL CALCIUM CARBONATE, CRUDE - UNII:2E32821G6I)	OYSTER SHELL CALCIUM CARBONATE, CRUDE	30 [hp_X] in 1 mL
GRAPHITE (UNII: 4QQN74LH4O) (GRAPHITE - UNII:4QQN74LH4O)	GRAPHITE	12 [hp_X] in 1 mL
CENTELLA ASIATICA (UNII: 7M867G6T1U) (CENTELLA ASIATICA - UNII:7M867G6T1U)	CENTELLA ASIATICA	3 [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

Inactive Ingredients

Ingredient Name		Strength		
ALCOHOL (UNII: 3K9958V90M)				
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58264-0065-1	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.