

D-10- sepia officinalis whole, sanguinaria canadensis root, sulfuric acid, black cohosh, and lachesis muta venom 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-10

NDC 58264-0010-1

INDICATIONS

Flushes of heat.

INGREDIENTS

ACTIVE

Sepia officinalis 4x, Sanguinaria canadensis 4x, Acidum sulfuricum 4x, Cimicifuga 4x, Lachesis mutus 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 10

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-10

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0010
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SEPIA OFFICINALIS WHOLE (UNII: 48GD5780QF) (SEPIA OFFICINALIS WHOLE - UNII:48GD5780QF)	SEPIA OFFICINALIS WHOLE	4 [hp_X] in 1 mL
SANGUINARIA CANADENSIS ROOT (UNII: N9288CD508) (SANGUINARIA CANADENSIS ROOT - UNII:N9288CD508)	SANGUINARIA CANADENSIS ROOT	4 [hp_X] in 1 mL
SULFURIC ACID (UNII: O40UQP6WCF) (SULFURIC ACID - UNII:O40UQP6WCF)	SULFURIC ACID	4 [hp_X] in 1 mL
BLACK COHOSH (UNII: K73E24S6X9) (BLACK COHOSH - UNII:K73E24S6X9)	BLACK COHOSH	4 [hp_X] in 1 mL
LACHESIS MUTA VENOM (UNII: VSW71SS07I) (LACHESIS MUTA VENOM - UNII:VSW71SS07I)	LACHESIS MUTA VENOM	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:58264-0010-1	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.