

D-59- fucus vesiculosus, spongia officinalis skeleton, roasted, graphite, oyster shell calcium carbonate, crude, sodium sulfate, and croton oil 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-59

NDC 58264-0059-1

INDICATIONS

Tendency to put on weight, slimming effect.

INGREDIENTS

ACTIVE

Fucus vesiculosus 3x, Spongia tosta 3x, Graphites 12x, Calcarea carbonica 12x, Natrum sulfuricum 2x, Croton tiglium 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 59

HOMEOPATHIC

STRESS FORMULA

1 FL. OZ.

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.

Rev. 3/18



NDC 58264-0059-1

INDICATIONS:

Tendency to put on weight, slimming effect.

INGREDIENTS:

ACTIVE: Fucus vesiculosus 3x, Spongia tosta 3x, Graphites 12x, Calcarea carbonica 12x, Natrum sulfuricum 2x, Croton tiglium 4x

INACTIVE: 20% alcohol in purified water.

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

D-59

fucus vesiculosus, spongia officinalis skeleton, roasted, graphite, oyster shell calcium carbonate, crude, sodium sulfate, and croton oil solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0059
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Fucus vesiculosus (UNII: 535G2ABX9M) (Fucus vesiculosus - UNII:535G2ABX9M)	Fucus vesiculosus	3 [hp_X] in 1 mL
Spongia officinalis skeleton, roasted (UNII: 1PIP394IID) (Spongia officinalis skeleton, roasted - UNII:1PIP394IID)	Spongia officinalis skeleton, roasted	3 [hp_X] in 1 mL
Graphite (UNII: 4QQN74LH4O) (Graphite - UNII:4QQN74LH4O)	Graphite	12 [hp_X] in 1 mL
Oyster shell calcium carbonate, crude (UNII: 2E32821G6I) (Oyster shell calcium carbonate, crude - UNII:2E32821G6I)	Oyster shell calcium carbonate, crude	12 [hp_X] in 1 mL
Sodium sulfate (UNII: 0YPR65R21J) (Sodium sulfate anhydrous - UNII:36KCS0R750)	Sodium sulfate	2 [hp_X] in 1 mL
Croton oil (UNII: WK97EQG57S) (Croton oil - UNII:WK97EQG57S)	Croton oil	4 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Alcohol (UNII: 3K9958V90M)	
Water (UNII: 059QF0KO0R)	

Packaging

Marketing Start Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58264-0059-1	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: 6/2018

DNA Labs, Inc.