

**AM-9- spigelia anthelmia, silicon dioxide, and corallium rubrum whole 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.

AM-9

NDC 58264-0348-1

INDICATIONS

Regulates energy flow in the triplewarmer meridian.

INGREDIENTS

ACTIVE

Spigelia anthelmia 8x, Silicea 8x, Corallium rubrum 10x

INACTIVE

20% alcohol in purified water.

SUGGESTED DOSAGE

5 to 10 drops twice a day, meridian energy excess 1 to 5 drops t.i.d, meridian lacking in energy 10 to 20 drops t.i.d.

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.

AM-9

TRIPLEWARMER

ACUPUNCTURE MERIDIAN

1 FL. OZ.

SUGGESTED DOSAGE:
5 to 10 drops twice a day, meridian
energy excess 1 to 5 drops t.i.d,
meridian lacking in energy
10 to 20 drops t.i.d.

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



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

AM-9

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0348
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SPIGELIA ANTHELMIA (UNII: WYT05213GE) (SPIGELIA ANTHELMIA - UNII:WYT05213GE)	SPIGELIA ANTHELMIA	8 [hp_X] in 1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	8 [hp_X] in 1 mL
CORALLIUM RUBRUM WHOLE (UNII: 22Z32PWR7R) (CORALLIUM RUBRUM WHOLE - UNII:22Z32PWR7R)	CORALLIUM RUBRUM WHOLE	10 [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:58264-0348-1	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.