

**N-18- oyster shell calcium carbonate, crude, sepia officinalis juice, silicon dioxide, nitric acid, and sus scrofa pineal gland 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.

N-18

NDC 58264-0216-1

INDICATIONS

Support for the Pineal.

INGREDIENTS

Calcarea carbonica 30x, Sepia 30x, Silicea 30x, Acidum nitricum 12x, Pineal 6/12/30/60/200x, 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.
- As with drugs if you are pregnant or nursing a baby seek professional advice before using this product.
- 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.**

**N-18
PINEAL ENERGIZER**

HOMEOPATHIC ENDOCRINE

1 FL. OZ.

SUGGESTED DOSAGE:

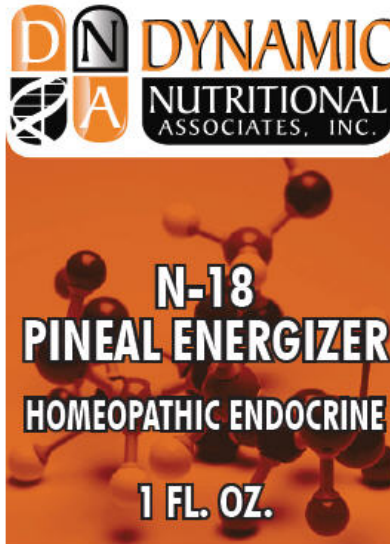
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MANUFACTURED FOR:
DNA LABORATORIES, INC.
Coeur d'Alene, ID 83814

N-18

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0216
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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
SEPIA OFFICINALIS JUICE (UNII: QDL83WN8C2) (SEPIA OFFICINALIS JUICE - UNII:QDL83WN8C2)	SEPIA OFFICINALIS JUICE	30 [hp_X] in 1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	30 [hp_X] in 1 mL
NITRIC ACID (UNII: 411VRN1TV4) (NITRIC ACID - UNII:411VRN1TV4)	NITRIC ACID	12 [hp_X] in 1 mL
SUS SCROFA PINEAL GLAND (UNII: 050QZ2EDK7) (SUS SCROFA PINEAL GLAND - UNII:050QZ2EDK7)	SUS SCROFA PINEAL GLAND	200 [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:58264-0216-1	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.