D-33- atropa belladonna, bufo bufo cutaneous gland, copper, anemone pratensis, silicon dioxide, and zinc 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-33

NDC 58264-0033-1

INDICATIONS

Epilepsy, irritations of the nerve.

INGREDIENTS

ACTIVE

Belladonna 30x, Bufo rana 200x, Cuprum metallicum 12x, Pulsatilla 30x, Silicea 30x, Zincum metallicum 12x

INACTIVE

20% alcohol in purified water.

Rx CAUTION

Federal law prohibits dispensing without a prescription.

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 33

HOMEOPATHIC
STRESS FORMULA

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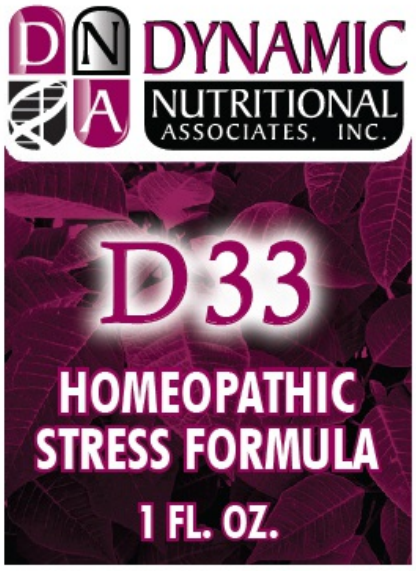
Rev. 3/18



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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:58264-0033
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ATROPA BELLADONNA (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA - UNII:WQZ3G9PF0H)	ATROPA BELLADONNA	30 [hp_X] in 1 mL
BUFO BUFO CUTANEOUS GLAND (UNII: Q59QU6N72Q) (BUFO BUFO CUTANEOUS GLAND - UNII:Q59QU6N72Q)	BUFO BUFO CUTANEOUS GLAND	200 [hp_X] in 1 mL
COPPER (UNII: 789U1901C5) (COPPER - UNII:789U1901C5)	COPPER	12 [hp_X] in 1 mL
ANEMONE PRATENSIS (UNII: 8E272251DI) (ANEMONE PRATENSIS - UNII:8E272251DI)	ANEMONE PRATENSIS	30 [hp_X] in 1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	30 [hp_X] in 1 mL

ZINC (UNII: J41CSQ7QDS) (ZINC - UNII:J41CSQ7QDS)	ZINC	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:58264-0033-1	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.