AM-7- strychnos nux-vomica seed, sodium phosphate, dibasic, heptahydrate, and bufo bufo 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-7

NDC 58264-0346-1

INDICATIONS

Regulates energy flow in the liver meridian.

INGREDIENTS

ACTIVE

Nux vomica 8x, Natrum phosphoricum 8x, Bufo rana 12x

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 NUTRITIONALASSOCIATES, INC.

AM-7

LIVER

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.

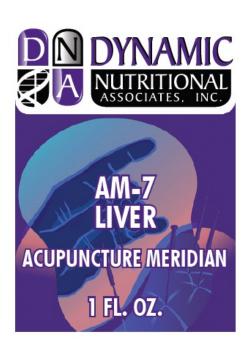
SHAKE WELL

Warnings:

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To be used according to standard homeopathic indications.

Rev. 3/18



NDC 58264-0346-1

INDICATIONS:

Regulates energy flow in the liver meridian.

INGREDIENTS:

ACTIVE: Nux vomica 8x, Natrum phosphoricum 8x, Bufo rana 12x INACTIVE: 20% alcohol in purified water.

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

AM-7

strychnos nux-vomica seed, sodium phosphate, dibasic, heptahydrate, and bufo bufo whole solution

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0346	
Route of Administration	SUBLINGUAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
STRYCHNOS NUX-VOMICA SEED (UNII: 269XH13919) (STRYCHNOS NUX-VOMICA SEED - UNII:269XH13919)	STRYCHNOS NUX-VOMICA SEED	8 [hp_X] in 1 mL		
SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE (UNII: 70WT22SF4B) (PHOSPHATE ION - UNII:NK08V8K8HR, SODIUM CATION - UNII:LYR4M0NH37)	SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE	8 [hp_X] in 1 mL		
BUFO BUFO WHOLE (UNII: DP1U601YQS) (BUFO BUFO WHOLE - UNII:DP1U601YQS)	BUFO BUFO WHOLE	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- 0346-1	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.