

**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.*

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**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**

**NUTRITIONAL**

ASSOCIATES, INC.

**AM-7**

# LIVER

## ACUPUNCTURE MERIDIAN

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1 to 5 drops t.i.d., meridian lacking in energy  
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Rev. 3/18



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MANUFACTURED FOR:

**DNA LABORATORIES, INC.**

Coeur d'Alene, ID 83814

## AM-7

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### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:58264-0346
<b>Route of Administration</b>	SUBLINGUAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>STRYCHNOS NUX-VOMICA SEED</b> (UNII: 269XH13919) (STRYCHNOS NUX-VOMICA SEED - UNII:269XH13919)	STRYCHNOS NUX-VOMICA SEED	8 [hp_X] in 1 mL
<b>SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE</b> (UNII: 70WT22SF4B) (PHOSPHATE ION - UNII:NK08V8K8HR, SODIUM CATION - UNII:LYR4M0NH37)	SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE	8 [hp_X] in 1 mL
<b>BUFO BUFO WHOLE</b> (UNII: DP1U601YQS) (BUFO BUFO WHOLE - UNII:DP1U601YQS)	BUFO BUFO WHOLE	12 [hp_X] in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>WATER</b> (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.