# D-44- crataegus laevigata fruit, prunus laurocerasus leaf, nerium oleander leaf, and cytisus scoparius 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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### **D-44**

# NDC 58264-0044-1

### INDICATIONS

Circulatory disorders, low blood pressure.

### INGREDIENTS

# **ACTIVE**

Crataegus oxyacantha 1x, Laurocerasus 3x, Oleander 6x, Spartium scoparium 6x

# **INACTIVE**

20% alcohol in purified water.

# **Rx CAUTION**

Federal law prohibits dispensing without prescription.

### SUGGESTED DOSAGE

One dropper under tongue two times daily. Acute symptoms  $\frac{1}{2}$  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 44

# HOMEOPATHIC STRESS FORMULA

1 FL. OZ.

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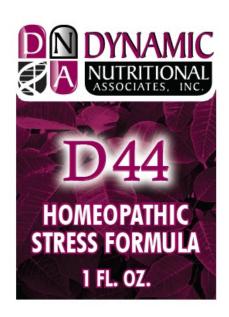
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Rev. 3/18



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

# **D-44**

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Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58264-0044	
Route of Administration	SUBLINGUAL			

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
CRATAEGUS LAEVIGATA FRUIT (UNII: D5RZ7MF1YF) (CRATAEGUS LAEVIGATA FRUIT - UNII:D5RZ7MF1YF)	CRATAEGUS LAEVIGATA FRUIT	1 [hp_X] in 1 mL		
<b>PRUNUS LAUROCERASUS LEAF</b> (UNII: F26P2D4757) (PRUNUS LAUROCERASUS LEAF - UNII:F26P2D4757)	PRUNUS LAUROCERASUS LEAF	3 [hp_X] in 1 mL		
NERIUM OLEANDER LEAF (UNII: 7KV510R6H6) (NERIUM OLEANDER LEAF - UNII: 7KV510R6H6)	NERIUM OLEANDER LEAF	6 [hp_X] in 1 mL		
CYTISUS SCOPARIUS WHOLE (UNII: V99S891QU7) (CYTISUS SCOPARIUS WHOLE - UNII: V99S891QU7)	CYTISUS SCOPARIUS WHOLE	6 [hp_X] in 1 mL		

# **Inactive Ingredients**

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

l	Packaging				
	# Item C	ode	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:5826		7 mL in 1 BOTTLE, GLASS; Type 0: Not a bination Product	01/01/1990	

Marketing Information				
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
	01/01/1990			
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date		

# **Labeler -** DNA Labs, Inc. (031784339)

Revised: 6/2022 DNA Labs, Inc.