COUMARIN PHENOLIC- coumarinum liquid Energique, 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.

Drug Facts:

ACTIVE INGREDIENT:

(in each drop): 20% of Coumarinum 6X, 12X, 30X, 12C, 30C.

INDICATIONS:

May temporarily relieve symptoms associated with reactions to coumarin.**

**Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

WARNINGS:

If pregnant or breastfeeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Do not use if tamper evident seal is broken or missing.

Store in a cool, dry place.

KEEP OUT OF REACH OF CHILDREN:

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

DIRECTIONS:

Adults and children 5 to 10 drops orally, 1 time daily or as otherwise directed by a health care professional. If symptoms persist for more than 7 days, consult your health care professional. Consult a physician for use in children under 12 years of age.

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INACTIVE INGREDIENTS:

Demineralized Water, 20% Ethanol.

QUESTIONS:

Dist. by Energique, Inc.

201 Apple Blvd.

Woodbine, IA 51579 800.869.8078

PACKAGE LABEL DISPLAY:

ENERGIQUE

SINCE 1987

COUMARIN

PHENOLIC

1 fl. oz. (30 ml)

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HOMEOPATHIC REMEDY

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LOT: XXXXXXX

COUMARIN PHENOLIC

coumarinum liquid

Product Information	Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0418		
Route of Administration	ORAL				

ı	Active Ingredient/Active Moiety		
ı	Ingredient Name	Basis of Strength	Strength
ı	COUMARIN (UNII: A4VZ22K1WT) (COUMARIN - UNII:A4VZ22K1WT)	COUMARIN	6 [hp_X] in 1 mL

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

Packaging					
#	Item Code		Package Description	Marketing Start Date	Marketing End Date
1	NDC:44911- 0418-1	30 mL Produc	in 1 BOTTLE, DROPPER; Type 0: Not a Combination et	04/05/2017	
T	Marketing I	nforr	nation		
Marketing Information					
	Marketing Cate	gory	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
uı	napproved homeop	athic		04/05/2017	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address		Business Operations
Apotheca Company		844330915	$manufacture (44911-0418) \ , \ api \ manufacture (44911-0418) \ , \ label (44911-0418) \ , \ pack (44911-0418) \ .$

Revised: 9/2020 Energique, Inc.