ARNICA- arnica montana 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): 100% of Arnica Montana 30C.

INDICATIONS:

May temporarily relieve bruising and stiffness after exertion.**

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

WARNINGS:

If pregnant or breast-feeding, 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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**Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

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

HOMEOPATHIC REMEDY

ARNICA 30C

1 fl. oz. (30 ml)

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LOT: XXXXXX MFD: MM/YY



ARNICA

arnica montana liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:44911-0218

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of
Strength

Strength

ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA -	ARNICA MONTANA	30 wk in 1 mL
UNII: O80TY2087 W	ANNICA MONTANA	JO WK III I IIIL

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

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:44911- 0218-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	07/15/2015	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/15/2015	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	$manufacture (44911-0218) \;,\; api\; manufacture (44911-0218) \;,\; label (44911-0218) \;,\; pack (44911-0218)$

Revised: 7/2022 Energique, Inc.