HYPERICUM- hypericum perforatum 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 Hypericum Perforatum 200C.

INDICATIONS:

May temporarily relieve nerve issues, especially after injury.**

**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.

INDICATIONS:

May temporarily relieve nerve issues, especially after injury.**

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

INACTIVE INGREDIENTS:

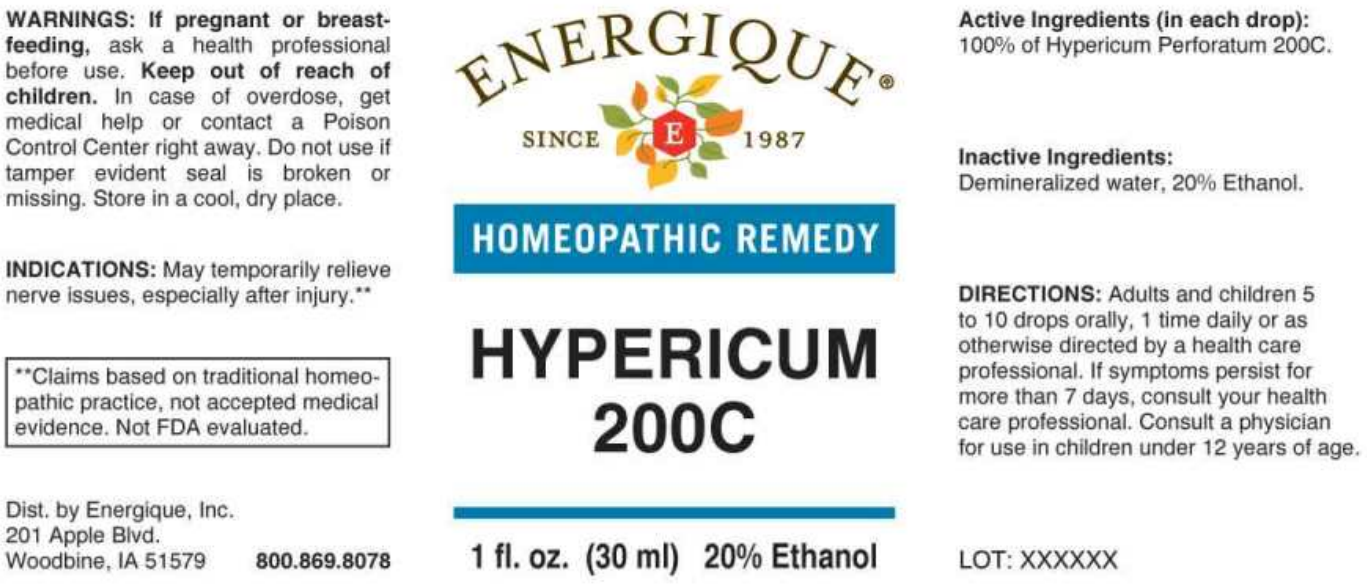
Demineralized water, 20% Ethanol.

QUESTIONS:

Dist. by Energique, Inc.
201 Apple Blvd.
Woodbine, IA 51579 **800.868.8078**

PACKAGE LABEL DISPLAY:

ENERGIQUE
SINCE 1987
HOMEOPATHIC REMEDY
HYPERICUM 200C
1 fl. oz. (30 ml)



HYPERICUM

hypericum perforatum liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44911-0344
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYPERICUM PERFORATUM (UNII: XK4IUX8MNB) (HYPERICUM PERFORATUM - UNII:XK4IUX8MNB)	HYPERICUM PERFORATUM	200 [hp_C] 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:44911-0344-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	04/12/2016	04/20/2026

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		04/12/2016	04/20/2026

Labeler - Energique, Inc (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0344) , api manufacture(44911-0344) , label(44911-0344) , pack(44911-0344)

Revised: 6/2022 Energique, Inc