PSORINUM- psorinum 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:

PSORINUM 7X

INDICATIONS:

To be used according to standard homeopathic indications.**

**These statements are based upon traditional homeopathic practice. They have not been reviewed by the Food and Drug Administration.

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.

RX ONLY:

Caution: Federal law prohibits dispensing without prescription.

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. If symptoms persist, 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

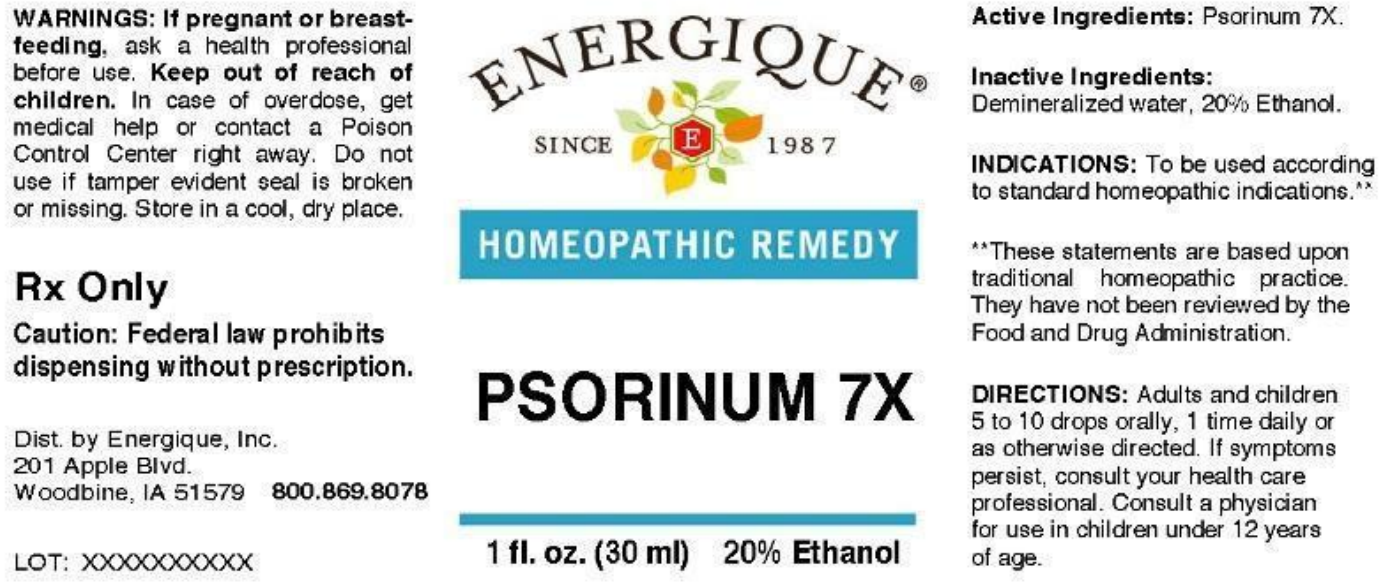
HOMEOPATHIC REMEDY

PSORINUM 7X

1fl. oz. (30 ml)

RX ONLY:

Caution: Federal law prohibits dispensing without prescription.



PSORINUM psorinum liquid
Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:44911-0097
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SCABIES LESION LYSATE (HUMAN) (UNII: 5UAU16Z1U4) (SCABIES LESION LYSATE (HUMAN) - UNII:5UAU16Z1U4)	SCABIES LESION LYSATE (HUMAN)	7 [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:44911-0097-1	30 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/09/2014	09/22/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/09/2014	09/22/2025

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0097) , api manufacture(44911-0097) , label(44911-0097) , pack(44911-0097)

Revised: 12/2021 Energique, Inc.