

**NAT MUR- natrum muriaticum 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.*

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**DRUG FACTS:**

**ACTIVE INGREDIENT:**

**(in each drop):** 100% of Natrum Muriaticum 30C.

**INDICATIONS:**

May temporarily relieve runny nose with watery discharge and much sneezing.\*\*

\*\*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:**

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

### HOMEOPATHIC REMEDY

### NAT MUR 30C

**1 fl. oz. (30 ml)**

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LOT: XXXXXX MFD: MM/YY



## NAT MUR

natrum muriaticum liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:44911-0159
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X) (CHLORIDE ION -	SODIUM CHLORIDE	30 [mg Cl] in 1 mL

UNII:Q32ZN48698)		SODIUM CHLORIDE	30 [mL] 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:44911-0159-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/19/2015	11/28/2028
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic			05/19/2015	11/28/2028

**Labeler -** Energique, Inc. (789886132)

**Registrant -** Apotheca Company (844330915)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Apotheca Company		844330915	manufacture(44911-0159) , api manufacture(44911-0159) , label(44911-0159) , pack(44911-0159)