FERRUM PHOS- ferrum phosphoricum 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 INGREDIENTS

Ferrum Phosphoricum 30C

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.

DIRECTIONS

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

INACTIVE INGREDIENTS

Demineralized water, 20% Ethanol.

KEEP OUT OF REACH OF CHILDREN

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

INDICATIONS AND USAGE

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.

QUESTIONS

Dist. by Energique, Inc.

201 Apple Blvd

Woodbine, IA 51579

800-869-8078

ENERGIQUE

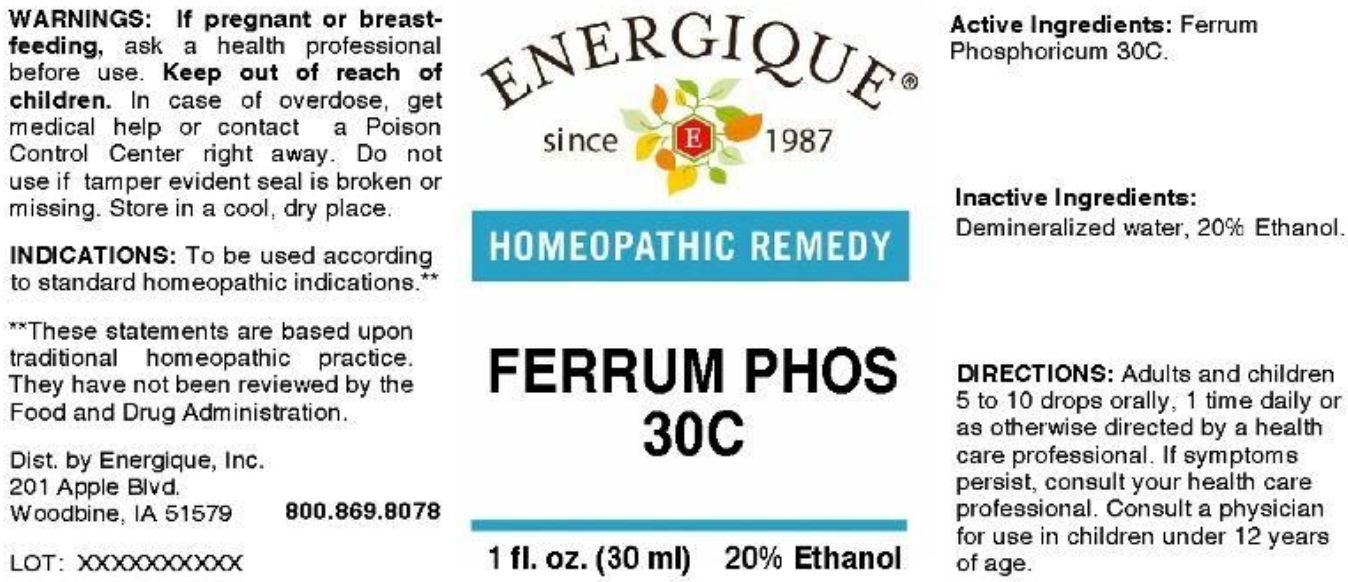
since 1987

HOMEOPATHIC REMEDY

FERRUM PHOS

30C

1 fl oz (30 ml)



FERRUM PHOS

ferrum phosphoricum liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44911-0042
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
FERRUM PHOSPHORICUM (UNII: 91GQH8I5F7) (FERROSOFERRIC PHOSPHATE - UNII:91GQH8I5F7)	FERRUM PHOSPHORICUM	30 [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:44911-0042-1	30 mL in 1 BOTTLE, DROPPER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/02/2012	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0042)

Revised: 10/2012 Energique, Inc.