BERBERIS VULGARIS 30C- berberis vulgaris root bark liquid
Natural Creations, 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.

BERBERIS VULGARIS 30C

ACTIVE INGREDIENT (HPUS*):

Berberis Vulgaris 30C

USES: Temporarily relieves minor discomfort in the extremities.**

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DIRECTIONS: Adults & children above 12 years: 10 drops orally 3 times daily, or as directed by a health care professional.

KEEP OUT OF THE REACH OF CHILDREN. In case of overdose (or accidental ingestion) get medical help or contact a Poison Control Center right away.

WARNINGS:

- Consult a physician for use in children under 12 years of age.
- **IF PREGNANT OR BREAST-FEEDING**, ask a health care professional before use.
- **KEEP OUT OF THE REACH OF CHILDREN**. In case of overdose (or accidental ingestion) get medical help or contact a Poison Control Center right away.
- Do not use if **TAMPER EVIDENT** seal is broken or missing.

INACTIVE INGREDIENTS: Ethyl Alcohol USP, Purified Water.

QUESTIONS & COMMENTS?

Natural Creations, Inc. / Woodbine, IA 51579 / 712-647-1600

**The letters "HPUS" indicate the component in the product are officially monographed in the Homeopathic Pharmacopeia of the United States.*

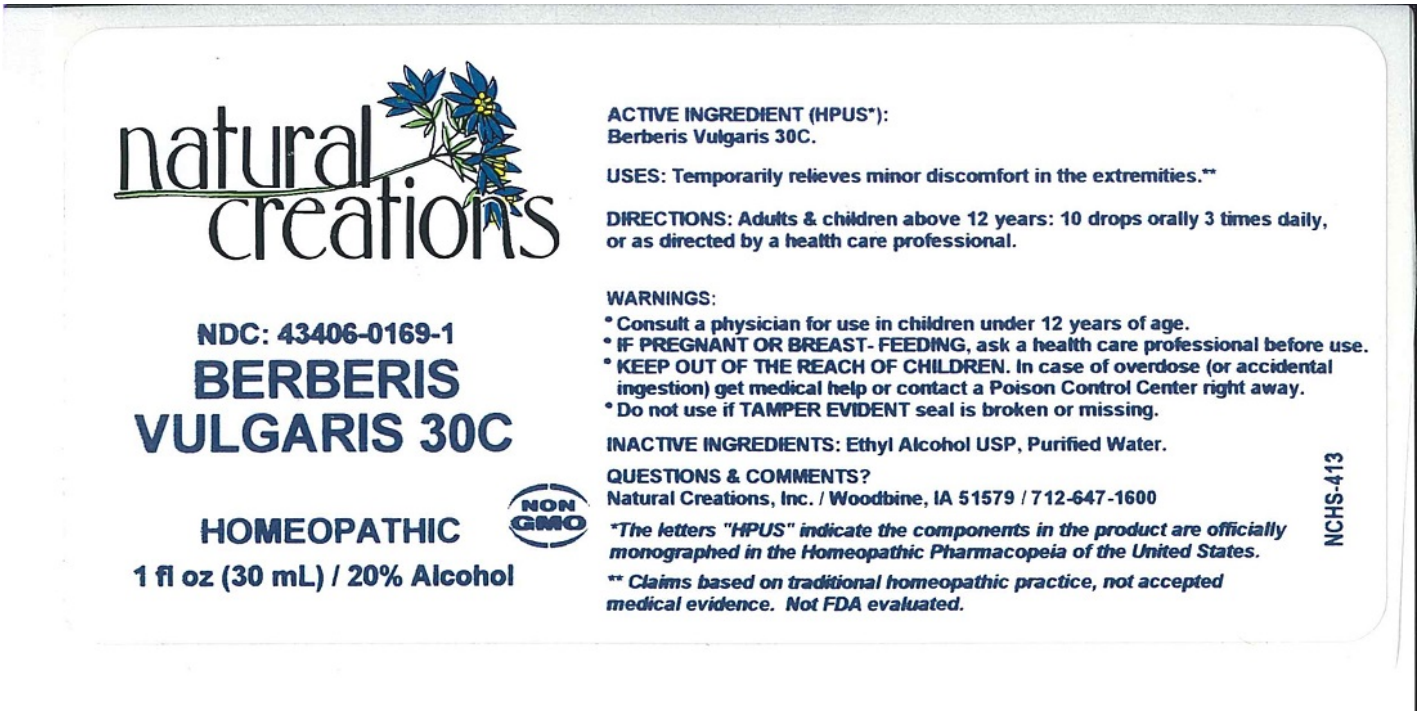
***Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.*

NDC: 43406-0169-1

BERBERIS VULGARIS 30C

HOMEOPATHIC

1 fl oz (30 mL) / 20% Alcohol



BERBERIS VULGARIS 30C

berberis vulgaris root bark liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43406-0169
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	30 [hp_C] in 1 mL

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:43406-0169-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/29/2007	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/29/2007	

Labeler - Natural Creations, Inc. (018022074)

Establishment

Name	Address	ID/FEI	Business Operations
OHM Pharma, Inc.		030572478	manufacture(43406-0169)

Revised: 12/2022 Natural Creations, Inc.