

ROCKY MOUNTAIN SPOTTED FEVER HOMOECHORD- rickettsia prowazekii liquid
Deseret Biologicals, 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:

Rickettsia Nosode 10C, 12C, 15C, 30C, 45C, 60C, 75C, 100C, 250C, 500C.

HOMEOPATHIC INDICATIONS:

For temporary relief of symptoms related to Rocky Mountain Spotted Fever infection including rash, fever, nausea, muscle pain and lack of appetite, abdominal and joint pain, and occasional diarrhea.**

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

WARNINGS:

Keep out of reach of children. In case of overdose, contact physician or Poison Control Center right away.

If pregnant or breast-feeding, ask a health professional before use.

Tamper seal: "Sealed for your protection." Do not use if seal is broken or missing.

KEEP OUT OF REACH OF CHILDREN:

Keep out of reach of children. In case of overdose, contact physician or Poison Control Center right away.

DIRECTIONS:

1-10 drops under the tongue, 3 times a day or as directed by a health professional. Consult a physician for use in children under 12 years of age.

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INACTIVE INGREDIENTS:

Demineralized Water, 25% Ethanol.

QUESTIONS:

Dist. By:

Deseret Biologicals, Inc.

469 W. Parkland Drive

Sandy, UT 84070 www.desbio.com

PACKAGE LABEL DISPLAY:

DESBIO

NDC 43742-0962-1

HOMEOPATHIC

ROCKY MOUNTAIN

SPOTTED FEVER

HOMOCHORD

1 FL OZ (30 ml)

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LOT:

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ROCKY MOUNTAIN SPOTTED FEVER HOMOCHORD

rickettsia prowazekii liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-0962
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
RICKETTSIA PROWAZEKII (UNII: TVS414L9M5) (RICKETTSIA PROWAZ EKII - UNII:TVS414L9M5)	RICKETTSIA PROWAZ EKII	10 [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:43742-0962-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	01/09/2017	09/02/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/09/2017	09/02/2025

Labeler - Deseret Biologicals, Inc. (940741853)**Registrant** - Apotheca Company (844330915)**Establishment**

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0962) , api manufacture(43742-0962) , label(43742-0962) , pack(43742-0962)