

BORRELIA BABESIA REMEDY- babesia microti, borrelia burgdorferi nosode 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 INGREDIENTS:

Babesia Microti 15X, 20X, 30X, 60X, 90X, 120X, 150X, 200X, 500X, 1000X, Borrelia Burgdorferi Nosode 15X, 20X, 30X, 60X, 90X, 120X, 150X, 200X, 500X, 1000X

HOMEOPATHIC INDICATIONS:

For the temporary relief of symptoms related to Lyme Disease including rash, fever, chills, fatigue, and migratory joint pain.**

**These statements are based upon traditional 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 a Poison Control Center right away.

If pregnant or breast-feeding, seek advice of 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 a 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 43742-0654-1

HOMEOPATHIC

BORRELIA-

BABESIA

REMEDY

1 FL OZ (30 ml)

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-0654
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BABESIA MICROTI (UNII: 1948X6KEG3) (BABESIA MICROTI - UNII:1948X6KEG3)	BABESIA MICROTI	15 [hp_X] in 1 mL
BORRELIA BURGDO RFERI (UNII: 0J8NV9V5Q8) (BORRELIA BURGDO RFERI - UNII:0J8NV9V5Q8)	BORRELIA BURGDORFERI	15 [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:43742-0654-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	11/11/2015	08/01/2021
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic			11/11/2015	08/01/2021

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0654) , api manufacture(43742-0654) , label(43742-0654) , pack(43742-0654)

Revised: 10/2016

Deseret Biologicals, Inc.