

HERPES SIMPLEX REMEDY- herpes simplex i, herpes simplex ii 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

Herpes Simplex I 15X, 20X, 30X, 60X, 90X, 120X, 150X, 200X, 500X, 1000X, Herpes Simplex II 15X, 20X, 30X, 60X, 90X, 120X, 150X, 200X, 500X, 1000X

HOMEOPATHIC INDICATIONS:

For temporary relief of symptoms related to Herpes Simplex infection including mouth sores, genital lesions, blisters or ulcers on mouth, lips and gums, fever blisters, fever and enlarged lymph nodes in the neck or groin.**

**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 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.

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.

INACTIVE INGREDIENTS

Demineralized water, 25% Ethanol.

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QUESTIONS

Dist. By: Deseret Biologicals, Inc.

469 W. Parkland Drive

Sandy, UT 84070

www.desbio.com

PACKAGE LABEL DISPLAY:

DESBIO

NDC 43742-0182-1

HOMEOPATHIC

HERPES SIMPLEX

REMEDY

1 FL OZ (30 ml)

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LOT:

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HERPES SIMPLEX REMEDY

herpes simplex i, herpes simplex ii liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-0182
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HUMAN HERPESVIRUS 1 (UNII: 22G38P19RL) (HUMAN HERPESVIRUS 1 - UNII:22G38P19RL)	HUMAN HERPESVIRUS 1	15 [hp_X] in 1 mL
HUMAN HERPESVIRUS 2 (UNII: 74J6DNH49U) (HUMAN HERPESVIRUS 2 -	HUMAN HERPESVIRUS	15 [hp_X]

UNII:74J6DNH49U)

2

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:43742-0182-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	10/12/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/12/2012	01/21/2021

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

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0182) , api manufacture(43742-0182) , label(43742-0182) , pack(43742-0182)

Revised: 5/2016

Deseret Biologicals, Inc.