HERPES ZOSTER REMEDY- herpes zoster 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:

Herpes Zoster 15X, 20X, 30X, 60X, 90X, 120X, 150X, 200X, 500X, 1000X.

HOMEOPATHIC INDICATIONS:

For temporary relief of symptoms related to Herpes Zoster infection including itching, tingling, burning, or painful rash usually on one side of the body or one side of the face, liver pain and mouth ulcers.**

**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, seek advice of 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.

KEEP OUT OF REACH OF CHILDREN:

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

HOMEOPATHIC INDICATIONS:

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QUESTIONS:

Dist. By: Deseret Biologicals, Inc.
469 W. Parkland Drive
Sandy, UT 84070 www.desbio.com

PACKAGE LABEL DISPLAY:

DES**BIO**

NDC 43742-0174-1

HOMEOPATHIC

HERPES ZOSTER

REMEDY

1 FL OZ (30 ml)

HERPES ZOSTER REMEDY

herpes zoster nosode liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43742-0174
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HUMAN HERPESVIRUS 3 (UNII: 9885M7D6JP) (HUMAN HERPESVIRUS 3 - UNII:9885M7D6JP)	HUMAN HERPESVIRUS 3	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:43742-0174-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	01/20/2016	09/25/2020

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		11/06/2012	09/25/2020

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0174) , api manufacture(43742-0174) , label(43742-0174) , pack(43742-0174)

Revised: 10/2016 Deseret Biologicals, Inc.