

**HERPES ZOSTER HOMOCORD- 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.*

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**Drug Facts:**

**ACTIVE INGREDIENTS:**

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

**HOMEOPATHIC INDICATIONS:**

For the temporary relief of symptoms related to Herpes Zoster infection including itching, tingling or painful rash usually on one side of the body or one side of the face.\*\*

\*\*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,** 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:**

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**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:**

**DES BIO**

NDC 43742-0731-1

HOMEOPATHIC

**HERPES ZOSTER**

**HOMOCHORD**

1 FL OZ (30 ml)

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**LOT:**

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<b>HERPES ZOSTER HOMOCHORD</b>			
herpes zoster nosode liquid			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:43742-0833
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>HUMAN HERPESVIRUS 3</b> (UNII: 9885M7D6JP) (HUMAN HERPESVIRUS 3 - UNII:9885M7D6JP)	HUMAN HERPESVIRUS 3	15 [hp_X] in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0K00R)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43742-0833-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/18/2016	06/30/2025

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/18/2016	06/30/2025

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

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0833) , api manufacture(43742-0833) , label(43742-0833) , pack(43742-0833)

Revised: 8/2021

Deseret Biologicals, Inc.