HUMAN PAPILLOMA VIRUS HOMOCHORD- human papilloma 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:

Human Papilloma Nosode 15X, 20X, 30X, 60X, 90X, 120X, 150X, 200X, 500X, 1000X.

INDICATIONS:

For the temporary relief of symptoms related to Human Papilloma Virus including itching in the pelvic area.**

**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, get medical help or contact 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 tamper evident seal is broken or missing.

KEEP OUT OF REACH OF CHILDREN:

Keep out of reach of children. In case of overdose, get medical help or contact 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:

DES**BIO**

NDC 43742-1228-1

HOMEOPATHIC

HUMAN

PAPILLOMA VIRUS

HOMOCHORD

1 FL OZ (30 ml)

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LOT:

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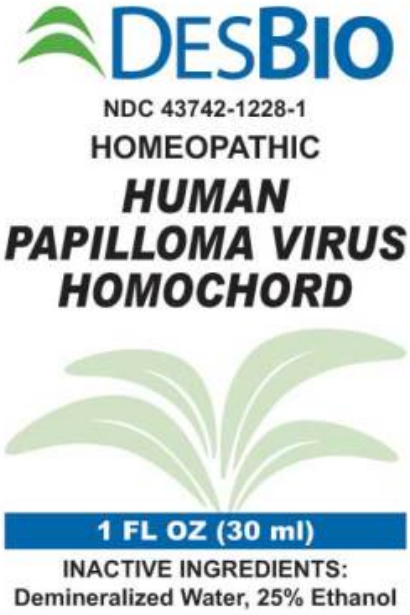

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HUMAN PAPILLOMA VIRUS HOMOCHORD

human papilloma nosode liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43742-1228

Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HUMAN PAPILLOMAVIRUS (UNII: 23CVL7WF4J) (HUMAN PAPILLOMAVIRUS - UNII:23CVL7WF4J)	HUMAN PAPILLOMAVIRUS	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:43742-1228-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	06/01/2018	06/16/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		06/01/2018	06/16/2025

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-1228) , api manufacture(43742-1228) , label(43742-1228) , pack(43742-1228)

Revised: 8/2021 Deseret Biologicals, Inc.