

ITCH FORMULA- calendula officinalis, apis mellifica, natrum muriaticum, urtica urens, histaminum hydrochloricum, ledum palustre, sulphur, copaiva officinalis, cantharis, croton tiglium, graphites, mezereum, petroleum, rhus venenata, staphysagria, antipyrinum 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:

Calendula Officinalis 3X, Apis Mellifica 6X, Natrum Muriaticum 6X, Urtica Urens 6X, Histaminum Hydrochloricum 6X, 12X, 30X, Ledum Palustre 6X, 12X, 30X, Sulphur 8X, Copaiva Officinalis 10X, Cantharis 12X, Croton Tiglium 12X, Graphites 12X, Mezereum 12X, Petroleum 12X, Rhus Venenata 12X, Staphysagria 12X, Antipyrinum 16X.

HOMEOPATHIC INDICATIONS:

For the temporary relief of symptoms related to itching, such as hives, itching at night, intolerable scratching, insect bites, poison ivy exposure, skin irritation, dry skin, and lice.**

**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 a 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.

KEEP OUT OF REACH OF CHILDREN:

Keep out of reach of children. In case of overdose, contact a physician or 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 43742-1413-1

HOMEOPATHIC

ITCH

FORMULA

1 FL OZ (30 ml)

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-1413
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALENDULA OFFICINALIS FLOWERING TOP (UNII: 18E7415PXQ) (CALENDULA OFFICINALIS FLOWERING TOP - UNII:18E7415PXQ)	CALENDULA OFFICINALIS FLOWERING TOP	3 [hp_X] in 1 mL
APIS MELLIFERA (UNII: 7S82P3R43Z) (APIS MELLIFERA - UNII:7S82P3R43Z)	APIS MELLIFERA	6 [hp_X] in 1 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	6 [hp_X] in 1 mL
HISTAMINE DIHYDROCHLORIDE (UNII: 3POA0Q644U) (HISTAMINE - UNII:820484N8I3)	HISTAMINE DIHYDROCHLORIDE	6 [hp_X] in 1 mL
RHODODENDRON TOMENTOSUM LEAFY TWIG (UNII: 877L01IZ0P) (LEDUM PALUSTRE TWIG - UNII:877L01IZ0P)	RHODODENDRON TOMENTOSUM LEAFY TWIG	6 [hp_X] in 1 mL
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	6 [hp_X] in 1 mL
COPAIFERA OFFICINALIS RESIN (UNII: 1VH544O5AT) (COPAIFERA OFFICINALIS RESIN - UNII:1VH544O5AT)	COPAIFERA OFFICINALIS RESIN	8 [hp_X] in 1 mL
LYTTA VESICATORIA (UNII: 3Q034RO3BT) (LYTTA VESICATORIA - UNII:3Q034RO3BT)	LYTTA VESICATORIA	10 [hp_X] in 1 mL
CROTON TIGLIUM SEED (UNII: 0HK2GZK66E) (CROTON TIGLIUM SEED - UNII:0HK2GZK66E)	CROTON TIGLIUM SEED	12 [hp_X] in 1 mL
GRAPHITE (UNII: 4QQN74LH4O) (GRAPHITE - UNII:4QQN74LH4O)	GRAPHITE	12 [hp_X] in 1 mL
DAPHNE MEZEREUM BARK (UNII: X2N6E405GV) (DAPHNE MEZEREUM BARK - UNII:X2N6E405GV)	DAPHNE MEZEREUM BARK	12 [hp_X] in 1 mL
KEROSENE (UNII: 1C89KKC04E) (KEROSENE - UNII:1C89KKC04E)	KEROSENE	12 [hp_X] in 1 mL
TOXICODENDRON VERNIX LEAFY TWIG (UNII: Y3VW699H96) (TOXICODENDRON VERNIX LEAFY TWIG - UNII:Y3VW699H96)	TOXICODENDRON VERNIX LEAFY TWIG	12 [hp_X] in 1 mL
DELPHINIUM STAPHISAGRIA SEED (UNII: 00543AP1JV) (DELPHINIUM STAPHISAGRIA SEED - UNII:00543AP1JV)	DELPHINIUM STAPHISAGRIA SEED	12 [hp_X] in 1 mL
ANTIPYRINE (UNII: T3CHA1B51H) (ANTIPYRINE - UNII:T3CHA1B51H)	ANTIPYRINE	16 [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:43742-1413-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	12/06/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/06/2018	

Registrant - Apothea Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apothea Company		844330915	manufacture(43742-1413) , api manufacture(43742-1413) , label(43742-1413) , pack(43742-1413)

Revised: 12/2018

Deseret Biologicals, Inc.