PERTUSSIS- aconitum napellus, bryonia (alba), hyoscyamus niger, ipecacuanha, sticta pulmonaria, tabacum, phosphorus, antimonium tartaricum, kreosotum, silicea, pertussinum, tetanus toxin 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:

Aconitum Napellus 4X, Bryonia (Alba) 4X, Hyoscyamus Niger 4X, Ipecacuanha 4X, Sticta Pulmonaria 4X, Tabacum 4X, Phosphorus 5X, Antimonium Tartaricum 6X, Kreosotum 6X, Silicea 8X, Pertussinum 30X, Tetanus Toxin 30X.

HOMEOPATHIC INDICATIONS:

For temporary relief of symptoms related to Pertussis infection including cough, bronchial congestion, fever, and vaccination reactions.**

**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 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 43742-1089-1 HOMEOPATHIC PERTUSSIS 1 FL OZ (30 ml)

WARNINGS:

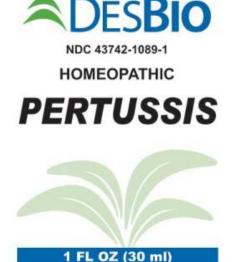
Keep out of reach of children. In case of overdose, contact a physician or Poison Control Center right away. If pregnant or breastfeeding, ask a health professional before use. Tamper seal: "Sealed for Your Protection." Do not use if seal is broken or missing.

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LOT:

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ACTIVE INGREDIENTS:

Aconitum Napellus 4X, Bryonia 4X, Hyoscyamus Niger 4X, Ipecacuanha 4X, Sticta Pulmonaria 4X, Tabacum 4X, Phosphorus 5X, Antimonium Tartaricum 6X, Kreosotum 6X, Silicea 8X, Pertussinum 30X, Tetanus Toxin 30X.

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PERTUSSIS

aconitum napellus, bryonia (alba), hyoscyamus niger, ipecacuanha, sticta pulmonaria, tabacum, phosphorus, antimonium tartaricum, kreosotum, silicea, pertussinum, tetanus toxin liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-1089
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACONITUM NAPELLUS (UNII: U0 NQ8555JD) (ACONITUM NAPELLUS -	ACONITIIM NAPELLIIS	4 [hp_X]	

UNII:U0NQ8555JD)	ACOMITOWINAFELLUS	in 1 mL
BRYONIA ALBA ROOT (UNII: T7J046 YI2B) (BRYONIA ALBA ROOT - UNII:T7J046 YI2B)	BRYONIA ALBA ROOT	4 [hp_X] in 1 mL
HYOSCYAMUS NIGER (UNII: 4WRK2153H3) (HYOSCYAMUS NIGER - UNII:4WRK2153H3)	HYOSCYAMUS NIGER	4 [hp_X] in 1 mL
IPECAC (UNII: 62I3C8233L) (IPECAC - UNII:62I3C8233L)	IPECAC	4 [hp_X] in 1 mL
LOBARIA PULMONARIA (UNII: D1YM0P5Z2T) (LOBARIA PULMONARIA - UNII:D1YM0P5Z2T)	LOBARIA PULMONARIA	4 [hp_X] in 1 mL
TOBACCO LEAF (UNII: 6YR2608RSU) (TOBACCO LEAF - UNII:6YR2608RSU)	TOBACCO LEAF	4 [hp_X] in 1 mL
PHO SPHO RUS (UNII: 27YLU75U4W) (PHO SPHO RUS - UNII:27YLU75U4W)	PHOSPHORUS	5 [hp_X] in 1 mL
ANTIMONY POTASSIUM TARTRATE (UNII: DL6OZ476V3) (ANTIMONY CATION (3+) - UNII:069647RPT5)	ANTIMONY POTASSIUM TARTRATE	6 [hp_X] in 1 mL
WOOD CREOSOTE (UNII: 3JYG22FD73) (WOOD CREOSOTE - UNII:3JYG22FD73)	WOOD CREOSOTE	6 [hp_X] in 1 mL
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4) (SILICON DIO XIDE - UNII: ETJ7Z6 XBU4)	SILICON DIOXIDE	8 [hp_X] in 1 mL
HUMAN SPUTUM, BORDETELLA PERTUSSIS INFECTED (UNII: U364V64HUN) (HUMAN SPUTUM, BORDETELLA PERTUSSIS INFECTED - UNII:U364V64HUN)	HUMAN SPUTUM, BORDETELLA PERTUSSIS INFECTED	30 [hp_X] in 1 mL
CLOSTRIDIUM TETANI (UNII: 751E8J54VM) (CLOSTRIDIUM TETANI - UNII:751E8J54VM)	CLOSTRIDIUM TETANI	30 [hp_X] in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ALCOHOL (UNII: 3K9958V90M)		

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:43742- 1089-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	10/31/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/31/2017	

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-1089, 43742-1089), api manufacture(43742-1089), pack(43742-1089)

Revised: 10/2017 Deseret Biologicals, Inc.