PYRROLE- pyrrole 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 INGREDIENT:

Pyrrole 8X, 12X, 30X, 200X, 12C, 30C, 60C, 200C.

HOMEOPATHIC INDICATIONS:

For temporary relief of sensitivity to wool, including gas, bloating, and headache.**

**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:

QUESTIONS:

Dist. By: Deseret Biologicals, Inc.

469 W. Parkland Drive

Sandy, UT 84070 www.desbio.com

PACKAGE LABEL DISPLAY:

DESBIO

NDC 43742-1323-1

HOMEOPATHIC

PYRROLE

1 FL OZ (30 ml)

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LOT:

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



HOMEOPATHIC





INACTIVE INGREDIENTS: Demineralized Water, 25% Ethanol ACTIVE INGREDIENTS: Pyrrole 8X, 12X, 30X, 200X, 12C, 30C, 60C, 200C.

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PYRROLE

pyrrole liquid

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Prod	luct	Inform	ation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:43742-1323

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRROLE (UNII: 86S1ZD6L2C) (PYRROLE - UNII:86S1ZD6L2C)	PYRROLE	8 [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- 1323-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	03/18/2020	03/26/2025

Marketing Information			
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
	03/18/2020	03/26/2025	
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date	

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-1323) , api manufacture(43742-1323) , label(43742-1323) , pack(43742-1323)

Revised: 6/2021 Deseret Biologicals, Inc.