LEVOTHYROXINE- levothyroxine 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:

Levothyroxine 16X.

HOMEOPATHIC INDICATIONS:

For the temporary relief of symptoms such as backache, constipation, and vomiting.**

**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:

Keep out of reach of children. In case of overdose, contact physician or 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:

DESBIO

NDC 43742-0955-1

HOMEOPATHIC

LEVOTHYROXINE 16X

1 FL OZ (30 ml)

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LEVOTHYROXINE

levothyroxine liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:43742-0955 Route of Administration ORAL

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
LEVOTHYROXINE (UNII: O51BO43MG4) (LEVOTHYROXINE - UNII: O51BO43MG4)	LEVOTHYROXINE	16 [hp X] in 1 mL			

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

Packaging

#	Item Code	tem Code Package Description Marketing Sta		Marketing End Date		
1	NDC:43742- 0955-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	0 1/0 6/20 17			
Marketing Information						
Marketing Category A		ory Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
uı	napproved homeop	athic	01/06/2017			

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment					
Name	Address	ID/FEI	Business Operations		
Apotheca Company		844330915	manufacture(43742-0955), api manufacture(43742-0955), label(43742-0955), pack(43742-0955)		

Revised: 1/2017 Deseret Biologicals, Inc.