# DHEA- dhea (dehydroepiandrosterone) 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.

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#### **DRUG FACTS:**

#### **ACTIVE INGREDIENT:**

DHEA (DEHYDROEPIANDROSTERONE) 6X, 12X, 30X, 200X, 12C, 30C, 60C, 200C

#### **HOMEOPATHIC INDICATIONS:**

For temporary relief of symptoms related to adrenal glands such as fatigue and low energy.\*\*

\*\*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:**

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#### **DIRECTIONS:**

1-10 drops under the tongue, 2 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 43742-0564-1 HOMEOPATHIC

#### **DHEA**

1 FL OZ (30 ml)

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LOT:

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#### DHEA

dhea (dehydroepiandrosterone) liquid

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:43742-0564

#### **Route of Administration**

ORAL

## **Active Ingredient/Active Moiety**

rearro mg. caronqrearro rioloty				
Ingredient Name	<b>Basis of Strength</b>	Strength		
PRASTERONE (UNII: 459AG36T1B) (PRASTERONE - UNII:459AG36T1B)	PRASTERONE	6 [hp X] in 1 mL		

### **Inactive Ingredients**

mucano mg. calema		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ALCOHOL (UNII: 3K9958V90M)		

### **Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43742- 0564-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	01/19/2015	

## **Marketing Information**

- 101 No. 111 g 1111 g 111				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		01/19/2015		

## Labeler - Deseret Biologicals, Inc. (940741853)

## Registrant - Apotheca Company (844330915)

## **Establishment**

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0564) , api manufacture(43742-0564) , label(43742-0564) , pack(43742-0564)

Revised: 1/2024 Deseret Biologicals, Inc.