ESTROGEN- folliculinum, oophorinum, 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:

Folliculinum 8X, 12X, 30X, 200X, 12C, 30C, 60C, 200C, Oophorinum (Suis) 8X, 12X, 30X, 200X, 12C, 30C, 60C, 200C.

PURPOSE:

Folliculinum - Breast Tenderness, Oophorinum (Suis) - Hot Flashes

USES:

- For the temporary relief of symptoms including:
- breast tenderness
 hot flashes

These statements are based upon homeopathic practice. They have not been reviewed by the Food and Drug Administration.

WARNINGS:

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, Contact a physician or Poison Control Center right away.

Tamper Evident: Sealed for your protection. Do not use if seal is broken or missing.

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.

INACTIVE INGREDIENTS:

Demineralized water, 25% ethanol

QUESTIONS:

Dist. By: Deseret Biologicals, Inc.

469 W. Parkland Drive

Sandy, UT 84070

www.desbio.com

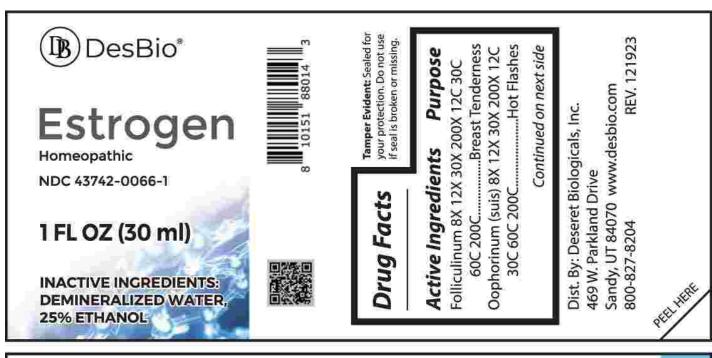
800-827-8204

PACKAGE LABEL DISPLAY:

DesBio

Estrogen

Homeopathic NDC 43742-0066-1 1 FL OZ (30 ml)



Deadener, No

cts (continued)

temporary relief of

temporary relief of symptoms ing: tenderness •hot flashes Continued on next side

Deadener, No text

(continued) **Drug Facts**

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Directions

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Other information

homeopathic principles. They have not been reviewed by the Food and Drug These statements are based upon Administration.

Demineralized water, 25% ethanol Inactive Ingredients

Basis of

Questions or Comments?

800-827-8204

ESTROGEN

folliculinum, oophorinum, liquid

Product Type HUMAN OTC DRUG **Item Code (Source)** NDC:43742-0066

ORAL **Route of Administration**

Active Ingredient/Active Moiety Ingredient Name

Strength Strength **ESTRONE** (UNII:

SUS SCROFA OV UNII:S7YTV04R8O

2DI9HA706A) (ESTRONE - UNII:2DI9HA706A)	ESTRONE	8 [hp_X] in 1 mL
VARY (UNII: S7YTV04R8O) (SUS SCROFA OVARY - D)	SUS SCROFA OVARY	8 [hp_X] in 1 mL

Inactive Ingredients

ALCOHOL (UNII: 3K9958V90M)

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Pa	cka	ging

Marketing Start Marketing End

#	item code	Package Description	Date	Date
1	NDC:43742- 0066-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	07/10/2012	
Marketing Information				
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	approved meopathic		07/10/2012	

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment				
Name	Address	ID/FEI	Business Operations	
Apotheca Company		844330915	manufacture(43742-0066) , api manufacture(43742-0066) , label(43742-0066) , pack(43742-0066)	

Revised: 4/2024 Deseret Biologicals, Inc.