

EPINEPHRINE- adrenalinum 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:

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

PURPOSE:

Adrenalinum - Exhaustion, Congestion, Stress

USES:

- For the temporary relief of symptoms including:
- exhaustion • congestion • stress

These statements are based upon homeopathic principles. 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
Epinephrine
Homeopathic
NDC 43742-1243-1
1 FL OZ (30 ml)

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INACTIVE INGREDIENTS:
DEMINEALIZED WATER,
25% ETHANOL

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60C 200C.....Exhaustion,
Congestion,
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Continued on next side

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Drug Facts (continued)

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Drug Facts (continued)

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Other Information

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Inactive Ingredients

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Questions or Comments?

800-827-8204

EPINEPHRINE

adrenalinum liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-1243
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
EPINEPHRINE (UNII: YKH834O4BH) (EPINEPHRINE - UNII:YKH834O4BH)	EPINEPHRINE	6 [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:43742-1243-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	07/10/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/10/2018	

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-1243) , api manufacture(43742-1243) , label(43742-1243) , pack(43742-1243)

Revised: 10/2024

Deseret Biologicals, Inc.