ADRENALINUM- adrenalinum pellet OHM PHARMA 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.

Adrenalinum

DRUG FACTS: ACTIVE INGREDIENTS: ADRENALINUM

CONTENT: Approx. 100 Pellets

USE: To be used according to standard homeopathic indications.

WARNINGS: Keep out of reach of children. If pregnant or breast-feeding, ask a health care professional before use. Stop use and ask a health care professional if symptoms persist for more than 3 days or worsen.

Keep out of reach of children.

DIRECTIONS: Dissolve 3-5 pellets under the tongue 3 times a day or as directed by professional.

OTHER INFORMATION: Store at room temperature. **Do not use** if pellet dispenser seal is broken.

INACTIVE INGREDIENTS: Organic sucrose, lactose free.

Manufactured according to the Homeopathic Pharmacopoeia of the United States (HPUS).

PRODUCT OF USA.

Mfg. By: OHM PHARMA, INC. Mineral Wells, TX 76067

www.ohmpharma.com FDA Est # 3003231743

ADRENALINUM

The OTC Potency range from Adrenalinum 6X-30X, 3C-30C, 200C, 1M.

Standard bottle sizes for dilution-form can range from 30mL to 60mL.



To be used according to standard homeopathic indications.

ADRENALINUM

adrenalinum pellet

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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66096-778
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Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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EPINEPHRINE (UNII: YKH834O4BH) (EPINEPHRINE - UNII:YKH834O4BH) EPINEPHRINE 6 [hp_C] in 6 [hp_C]

Inactive Ingredients

Ingredient Name	Strength
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SUCROSE (UNII: C151H8M554)

Product Characteristics

Color	white	Score	
Shape	ROUND	Size	4mm
Flavor		Imprint Code	
Contains			

Packaging

#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66096-778- 01	6 [hp_C] in 1 TUBE; Type 0: Not a Combination Product	11/01/2019	

Marketing Information			
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
	11/01/2019		
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date	

Labeler - OHM PHARMA INC. (030572478)

Revised: 12/2021 OHM PHARMA INC.