# AXIV ALLERGY- diphenhydramine hydrochloride capsule, liquid filled VIVUNT LLC

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### **AXIV Allergy**

#### **Drug Facts**

Active Ingredient (in each softgel)	Purpose
Diphenhydramine HCl 25 mg	Antihistamine

#### Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat
- temporarily relieves these symptoms due to the common cold:
- runny nose
- sneezing

## Warnings

#### Do not use

- to make a child sleepy
- with any other product containing diphenhydramine, even one used on skin

### Ask a doctor before use if you have

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- trouble urinating due to an enlarged prostate gland

## Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers

## When using this product

- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

## Keep out of reach of children

Keep out of reach of children.

#### If pregnant or breast-feeding

ask a health professional before use.

#### **Overdose Warning**

In case of overdose, get medical help or contact a Poison Control Center immediately.

#### **Directions**

- Take every 4 to 6 hours, or as directed by a doctor
- Do not take more than 12 Softgels in 24 hours for Adults and children 12 years and over. Do not take more than 6 Softgels for children 6 to under 12 years of age in 24 hours.

adults and children 12 years and over	1 or 2 Softgels
children 6 to under 12 years	1 Softgel
children under 6 years	do not use

#### Other information

- Store at 20° 25 °C (68 °- 77 °F)
- Read all product information before using
- Tamper Evident: Do not use if the foil printed on the blister is torn or ripped.
- Tamper Evident: Do not use if carton or pouch is open

## **Inactive ingredients**

Gelatin, Glycerin, Methylparaben, Polyethylene Glycol 400, Propylparaben, Purified Water, Sorbitol, Titanium Dioxide

#### PRINCIPAL DISPLAY PANEL 24

Compare to Benadryl® Allergy Liqui-Gels®

active ingredients\*

NDC 82706-019-01

**AXIV** Antihistamine

- Runny Nose
- Sneezing
- Itchy, Watery Eyes
- Itchy Throat

**DYE-FREE** 

Diphenhydramine HCl

24 SOFTGELS\*\* \*\*Liquid-filled capsules



Sorbitol, Titanium Dioxide.

#### PRINCIPAL DISPLAY PANEL 6

Compare to Benadryl® Allergy Liqui-Gels®

active ingredients\*

NDC 82706-019-02

**AXIV** Antihistamine

Runny Nose Sneezing Itchy, Watery Eyes Itchy Throat

**DYE-FREE** 

Diphenhydramine HCl

6 SOFTGELS\*\* \*\*Liquid-filled capsules





Active ingredient (in each caplet)
Diphenhydramine HOI 25 mg ......

Drug Facts

Antihistamine

Purpose

## **AXIV ALLERGY**

diphenhydramine hydrochloride capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82706-019
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients		
Ingredient Name	Strength	
PROPYLPARABEN (UNII: Z8IX2SC10H)		
GELATIN (UNII: 2G86QN327L)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		
SORBITOL (UNII: 506T60A25R)		
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

Product Characteristics			
Color	white (Transparent)	Score	no score
Shape	OVAL	Size	22mm
Flavor		Imprint Code	AXIV
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82706- 019-01	2 in 1 CARTON	03/08/2024	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:82706- 019-02	3 in 1 CARTON	03/08/2024	
2		2 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/08/2024	

## Labeler - VIVUNT LLC (045829437)

Revised: 3/2024 VIVUNT LLC