# DIPHENHYDRAMINE HYDROCHLORIDE 25MG- diphenhydramine hydrochloride 25mg tablet

**Strive Pharmaceuticals Inc** 

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## **RIGHT REMEDIES Allergy Relief**

## **Drug Facts**

## Active ingredient (in each tablet)

Diphenhydramine HCl 25 mg

## **Purpose**

**Antihistamine** 

#### Uses

temporarily relieves these symptoms due to hay fever or 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

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

Keep out of reach of children.

**Overdose warning:** In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as children even if you do not notice any signs or symptoms.

#### **Directions**

- take every 4 to 6 hours, or as directed by doctor
- do not take more than 6 times in 24 hours

adults and children 12 years	1 to 2
and older	tablets
children 6 to under 12 years	1 tablet
children under 6 years old	do not use

#### Other information

- each tablet contains: calcium 18.64 mg
- store between 20-25°C (68-77°F)
- protect from light.

## Inactive ingredients

croscarmellose sodium, dicalcium phosphate, D&C red #27, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, silicon dioxide, talc, titanium dioxide

### Questions or comments?

**1-888-577-8033** Monday-Friday 8am-4pm EST

#### RIGHT REMEDIES

Compare to the active ingredient of Benadryl® Allergy ULTRATAB®\*

\*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Benadryl<sup>®</sup> Allergy ULTRATAB<sup>®</sup>

Distributed by:

Strive Pharmaceuticals Inc.

East Brunswick, NJ 08816

Product of India

Packaged and Quality Assured in the USA

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

## **Packaging**

Packaging: 100 tablets

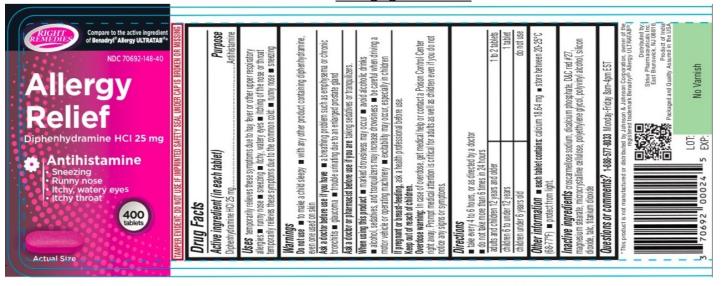


Packaging: 72 tablets





#### Packaging: 400 tablets

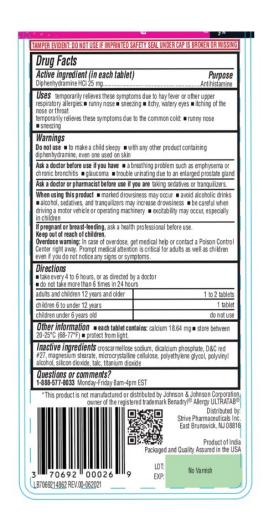


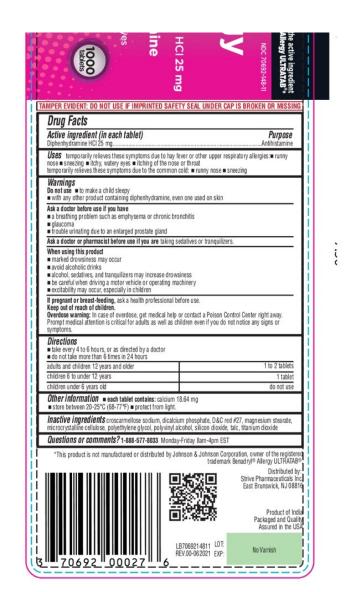
#### Packaging: 600 tablets



#### Packaging: 1000 tablets







### DIPHENHYDRAMINE HYDROCHLORIDE 25MG

diphenhydramine hydrochloride 25mg tablet

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:70692-148

Route of Administration ORAL

## **Active Ingredient/Active Moiety**

Ingredient Name

Basis of Strength

DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40)
(DIPHENHYDRAMINE - UNII:8GTS82S83M)

DIPHENHYDRAMINE HYDROCHLORIDE

25 mg

DITTENTI DIVAMINE ONII.001302303M

Inactive Ingredients		
Strength		

TALC (UNII: 7SEV7J4R1U)

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
<b>D&amp;C RED NO. 27</b> (UNII: 2LRS185U6K)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	

Product Characteristics			
Color	pink	Score	no score
Shape	CAPSULE	Size	12mm
Flavor		Imprint Code	DP
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:70692-148- 01	100 in 1 PACKAGE; Type 0: Not a Combination Product	12/28/2018		
2	NDC:70692-148- 46	1 in 1 CARTON	04/06/2023		
2		72 in 1 BOTTLE; Type 0: Not a Combination Product			
3	NDC:70692-148- 40	400 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2021		
4	NDC:70692-148- 62	600 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2021		
5	NDC:70692-148- 11	1000 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2021		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M012	12/20/2018		

## Labeler - Strive Pharmaceuticals Inc (080028013)

Revised: 12/2024 Strive Pharmaceuticals Inc