

**RIGHT REMEDIES ALLERGY RELIEF- diphenhydramine hydrochloride capsule**  
**Strive Pharmaceuticals Inc**

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**RIGHT REMEDIES Allergy Relief**

**Drug Facts**

**Active ingredient (in each capsule)**

Diphenhydramine HCl 25 mg

**Purpose**

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

**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 a doctor
- do not take more than 6 times in 24 hours

adults and children 12 years and older	1 to 2 capsules
children 6 to under 12 years	1 capsule
children under 6 years old	do not use

## **Other information**

- store between 20-25°C (68-77°F) • protect from light.

## **Inactive ingredients**

butylparaben, D&C red #28, edible ink, FD&C blue #1, FD&C red #40, gelatin, lactose monohydrate, magnesium stearate, methylparaben, polysorbate-80, propylparaben, purified water, sodium lauryl sulphate, starch

## **Questions or comments?**

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

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

- Sneezing
- Runny nose
- Itchy, watery eyes
- Itchy throat

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

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

Distributed by:  
Strive Pharmaceuticals Inc.  
East Brunswick, NJ 08816

Product of India  
Packaged and Quality Assured in the USA

## **Packaging**





lactose monohydrate, magnesium stearate, methylparaben, polysorbate-80, propylparaben, purified water, sodium lauryl sulphate, starch

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## RIGHT REMEDIES ALLERGY RELIEF

diphenhydramine hydrochloride capsule

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:70692-832
<b>Route of Administration</b>	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
<b>BUTYLPARABEN</b> (UNII: 3QPI1U3FV8)	
<b>D&amp;C RED NO. 28</b> (UNII: 767IP0Y5NH)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GELATIN, UNSPECIFIED</b> (UNII: 2G86QN327L)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	

### Product Characteristics

<b>Color</b>	pink	<b>Score</b>	no score
<b>Shape</b>	capsule (Cylindrical)	<b>Size</b>	14mm
<b>Flavor</b>		<b>Imprint Code</b>	P25
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70692-832-11	1000 in 1 BOTTLE; Type 0: Not a Combination Product	12/20/2021	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/20/2021	

**Labeler** - Strive Pharmaceuticals Inc (080028013)

Revised: 12/2024

Strive Pharmaceuticals Inc