DIPHENHYDRAMINE HYDROCHLORIDE- diphenhydramine hydrochloride capsule Walgreen Company
0835K-Walgreen
Drug Facts
Active ingredient (in each capsule) Diphenhydramine HCl 25 mg
<b>Purpose</b> Antihistamine
Uses
<ul> <li>□ temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:</li> <li>□ runny nose</li> <li>□ sneezing</li> <li>□ itchy, watery eyes</li> <li>□ itching of the nose or throat</li> <li>□ temporarily relieves these symptoms due to the common cold:</li> <li>□ runny nose</li> <li>□ sneezing</li> </ul>
Do not use
☐ to make a child sleepy
$\hfill \square$ 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
<ul><li>□ avoid alcoholic drinks</li><li>□ alcohol, sedatives and tranquilizers may increase drowsiness</li></ul>
use caution 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.
In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

### **Directions**

$\ \square$ take every 4 to 6 hours, or as directed by a doctor	
do not take more than 6 doses in 24 hours	
adults and children 12 years of age and over	1 to 2 capsules
children 6 to under 12 years of age	1 capsule
children under 6 years of age	do not use

### Other information

☐ store in a dry place at 15° – 30°C (59° – 86°F)
☐ protect from moisture
$\hfill \square$ do not use if imprinted safety seal under cap is broken or missing or if red band around capsule is broken or missing

corn starch, D&C red #28, FD&C blue #1, FD&C red #40, gelatin, lactose monohydrate, magnesium stearate, sodium lauryl sulfate

## Questions or comments?

1-800-925-4733

Walgreens Pharmacist Recommended.

Our pharmacists recommend the Walgreens brand. We invite you to compare to national brands.

††This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Benadryl®.

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PRODUCT OF CHINA. MANUFACTURED AND PACKAGED IN THE USA USING DOMESTIC AND IMPORTED INGREDIENTS.

NDC 0363-8350-02

Compare to the active ingredient in Benadryl® Allergy††

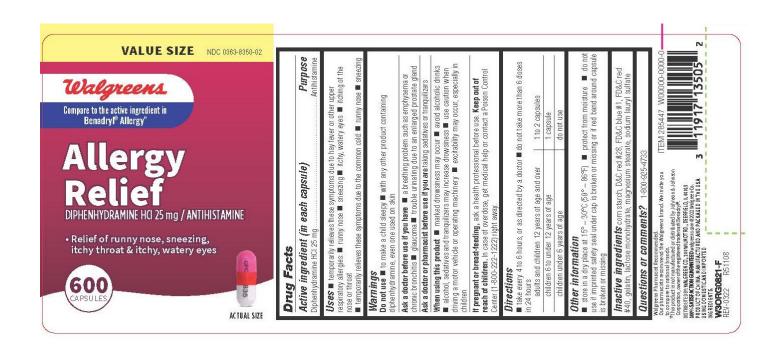
Allergy

Relief

DIPHENHYDRAMINE HCl 25 mg / ANTIHISTAMINE

• Relief of runny nose, sneezing, itchy throat & itchy, watery eyes

600 Capsules



## DIPHENHYDRAMINE HYDROCHLORIDE

diphenhydramine hydrochloride capsule

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-8350		
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			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
GELATIN (UNII: 2G86QN327L)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
STARCH, CORN (UNII: O8232NY3SJ)				
<b>D&amp;C RED NO. 28</b> (UNII: 767IP0Y5NH)				

Product Characteristics				
Color	pink ( HALF PINK AND HALF CLEAR WITH WHITE POWDER INSIDE AND SEALED WITH RED BAND)	Score	no score	
Shape	CAPSULE	Size	14mm	
Elaver		Imprint	CDC:035	

Fiavor	Code	CPC,033
Contains		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-8350- 06	365 in 1 BOTTLE; Type 0: Not a Combination Product	08/12/2022	
2	NDC:0363-8350- 02	600 in 1 BOTTLE; Type 0: Not a Combination Product	10/17/2022	
3	NDC:0363-8350- 01	1 in 1 CARTON	02/02/2024	
3		100 in 1 BOTTLE; 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	08/12/2022	

# Labeler - Walgreen Company (008965063)

Revised: 2/2024 Walgreen Company