# ALLERGY RELIEF- diphenhydramine hcl tablet, coated NORTHEAST PHARMA

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hpc 681 (683)

### Active ingredient (in each tablet)

Diphenhydramine HCl 25mg

### **Purpose**

**Antihistamine** 

#### Uses

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

## 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

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

## Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers

## When using this product

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

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 right away.

#### **Directions**

- do not exceed recommended dose
- adults and children 12 years and over, take 1 to 2 tablets every 4-6 hours, as needed; not more than 12 tablets in 24 hours, or as directed by a doctor
- children under 12 years: consult a doctor

#### Other information

- each tablet contains: calcium 30 mg
- store at 20-25°C (68-77°F); excursions permitted between 15°C 30°C (59°F 86°F)
- protect from moisture
- **Tamper Evident:** Do not use if imprinted seal under cap is missing or broken.

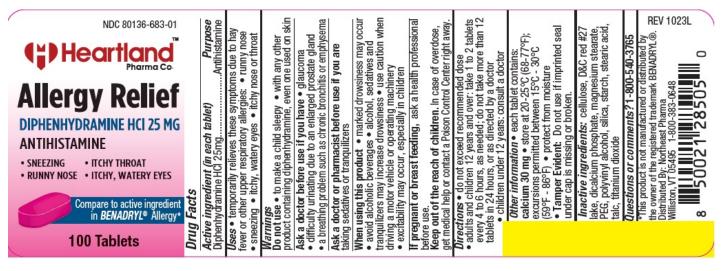
#### **Inactive ingredients**

cellulose, D&C red #27 lake, dicalcium phosphate, magnesium stearate, PEG, polyvinyl alcohol, silica, starch, stearic acid, talc, titanium dioxide

#### Questions or comments?

1-800-540-3765

## package label



# ALLERGY RELIEF

diphenhydramine hcl tablet, coated

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80136-683		
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		
STEARIC ACID (UNII: 4ELV7Z65AP)			
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)			
STARCH, CORN (UNII: O8232NY3SJ)			
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
<b>D&amp;C RED NO. 27</b> (UNII: 2LRS185U6K)			
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3M)Q0SDW1A)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
TALC (UNII: 7SEV7J4R1U)			

Product Characteristics				
Color	pink	Score	no score	
Shape	OVAL	Size	11mm	
Flavor		Imprint Code	44;329	
Contains				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:80136-683- 01	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2023		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/01/2023	

## Labeler - NORTHEAST PHARMA (081232935)

## Registrant - Geri-Care Pharmaceutical Corp (611196254)

Revised: 11/2023 NORTHEAST PHARMA