## ALLERGY RELIEF- diphenhydramine hydrochloride tablet HealthLife of USA

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#### **DIPHENHYDRAMINE HCI 25mg, USP**

#### **Active Ingredient**

(in each tablet)

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
- itchy nose or throat
- temporarily relieves these symptoms of the common cold:
- runny nose
- sneezing

#### **Warnings**

#### Do not use

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

## Ask a doctor before use if you have

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

## Ask a doctor or pharmacist before use if youaretaking sedatives or tranquilizers

## When using this product

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

#### If pregnant or breast-feeding;

#### Keep out of reach of children.

In case of accidental overdose, contact a doctor or Poison Control Center immediately. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Do not exceed recommended dosage.

#### **Directions**

- take every 4 to 6 hours, not more than 6 doses in 24 hours
- Adults and children 12 years of age and older:1 or 2 tablets
- children 6 to under 12 years of age:1 tablet
- children 4 to under 6 years of age:do not use unless directed by a doctor
- children under 4 years of age:do not use

#### Other Information

- each tablet contains : calcium 20 mg
- store at controlled room temperature 20°-25°C (68°-77°F).
- read all product information before using.
- TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

#### **Inactive Ingredients**

Colloidal silicon Dioxide, Croscarmellose Sodium, Dicalcium Phosphate, D & C Red, Magnesium stearate, Microcrystalline cellulose, Polyvinyl alcohol, Titanium dioxide, Talc

#### **Questions or Comments**

1-844-832-1138 (Mon-Fri 9AM-5PM EST) or www.healthlifeofusa.com

## PACKAGE LABEL.PRINCIPAL DISPLAY PANEL DIPHENHYDRAMINE HYDROCHLORIDE TABLET, USP 25 MG

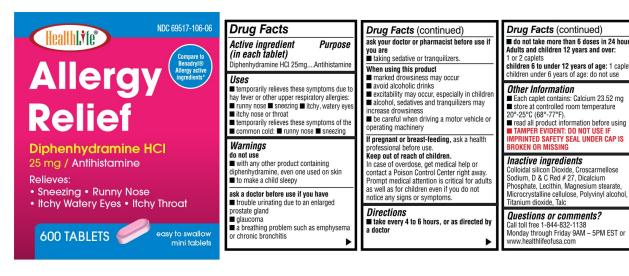
#### **ANTIHISTAMINE**

\* This product is not manufactured or distributed by McNeil-Consumer Healthcare, owner of the registered trademark Benadryl Allergy.

69517-106-24 24 Caplets

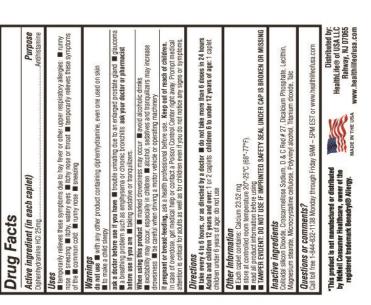


69517-106-06 600Caplets



#### 69517-106-10 1000 Caplets







HealthLife® of USA LLC Rahway, NJ 07065 www.healthlifeofusa.com

> Lot No.: Exp. Date:

#### **ALLERGY RELIEF**

diphenhydramine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69517-106
Route of Administration	ORAL		

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

Inactive Ingredients	
Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
<b>D&amp;C RED NO. 27</b> (UNII: 2LRS185U6K)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	pink	Score	no score
Shape	CAPSULE	Size	11mm
Flavor		Imprint Code	EL
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69517-106- 24	24 in 1 BOTTLE	01/01/2016	
1		1 in 1 CARTON; Type 0: Not a Combination Product		
2	NDC:69517-106- 30	1 in 1 CARTON	06/05/2017	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:69517-106- 10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	05/12/2012	
4	NDC:69517-106- 06	600 in 1 BOTTLE; Type 0: Not a Combination Product	05/12/2012	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	05/12/2012	

## Labeler - HealthLife of USA (079656178)

Revised: 10/2023 HealthLife of USA