

ALLERGY RELIEF- diphenhydramine hydrochloride tablet
Cabinet Health, Inc

DIPHENHYDRAMINE HCl Tablets, USP 25mg

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 you are taking 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;

ask a health professional before use

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

1-908-242-6108 (Mon-Fri 8AM-5PM EST)

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.

82725-1001-1



Compare to BENADRYL®
Allergy ULTRATAB® Tablets
active ingredient†

ALLERGY RELIEF

ANTIHISTAMINE
DIPHENHYDRAMINE HCl 25 mg

For the temporary relief of:
• Sneezing • Itchy Throat
• Runny Nose • Itchy, Watery Eyes



600 TABLETS

Drug Facts

Active ingredient (in each tablet)	Purpose
Diphenhydramine HCl 25mg.....	Antihistamine

Uses

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

- trouble urinating due to an enlarged prostate gland
- glaucoma

Ask your doctor or pharmacist before use if you are

- a breathing problem such as emphysema or chronic bronchitis
- taking the blood thinning drug warfarin

Drug Facts (continued)

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

When using this product

- drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness. avoid alcoholic beverages
- 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

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 over	1 to 2 tablets
children 6 to under 12 years	1 tablets
children under 6 years	do not use

Drug Facts (continued)

Other information

- protect from moisture
- store at 25°C (77°F) excursions permitted between 15°-30°C (59°-86°F)

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, dicalcium phosphate, d&c red 27, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, titanium dioxide, talc

Questions or Comments? contact 1-908-242-6108 Mon-Fri 8:00 AM EST to 5:00 PM PST

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

IMPORTANT: READ THE DIRECTIONS AND WARNINGS BEFORE USE

Distributed By: Cabinet Health P.B.C
Brooklyn, NY 11222 ITEM# 10001-01 Lot.

EXP.



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ALLERGY RELIEF

diphenhydramine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82725-1001
Route of Administration	ORAL		

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

Inactive Ingredients		
Ingredient Name		Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)		
D&C RED NO. 27 (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: 6OZP39ZG8H)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

Product Characteristics			
Color	pink	Score	no score
Shape	CAPSULE	Size	11mm
Flavor		Imprint Code	N02
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82725-1001-1	600 in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	11/01/2024	

Labeler - Cabinet Health, Inc (117102391)

Revised: 8/2025

Cabinet Health, Inc