

CABINET ALLERGY RELIEF- diphenhydramine hydrochloride tablet, film coated
Spirit Pharmaceutical LLC

Cabinet Allergy Relief

Drug Facts

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
 - itchy, watery eyes
 - sneezing
 - 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
- trouble urinating due to an enlarged prostate gland
- glaucoma

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. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- take every 4 to 6 hours
- do not take more than 6 doses in 24 hours

adults and children 12 years of age and over	1 to 2 tablets
children 6 to under 12 years of age	1 tablet
children under 6 years of age	do not use this product in children under 6 years of age

Other information

- store at controlled room temperature 15°-30° C (59°-86° F)
- protect from moisture and light
- see end flap for expiration date and lot number
- **each tablet contains** calcium 24 mg

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, dicalcium phosphate, D&C red # 27 Al lake, lecithin, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, talc, titanium dioxide

Questions or comments?

1-888-333-9792

Distributed by:
Cabinet Health, Inc.
wearecabinet.com
Made in India

PRINCIPAL DISPLAY PANEL - 25 mg Tablet Bottle Carton

Compare to the active ingredient
in Benadryl® Allergy*

CABINET:

Allergy Relief

Diphenhydramine HCl 25mg

Warnings
Do not use
• with any other product containing diphenhydramine, even one used on skin

Directions
• take every 4 to 6 hours
• do not take more than 6 doses in 24 hours

adults & children 12 years & over	1 to 2 tablets
children 6 to under 12 years	1 tablet

• do not use this product in children under 6 years of age

Other information
contains salicin 24 mg
• store between 15-30°C
• protect from moisture and light
• read and keep container for complete instructions and product information

Questions or comments?
1-800-845-8454

*This product is not manufactured or distributed by Mergal Consumer Products, Inc. For more information, please contact Mergal Consumer Products, Inc. at 1-800-845-8454.

Distributed by: Cabinet Health, Inc.
www.cabinet.com Made in India

CABINET ALLERGY RELIEF

diphenhydramine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68210-1280
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)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	pink	Score	no score
Shape	OVAL	Size	11mm

Flavor		Imprint Code	S4	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210-1280-1	1 in 1 CARTON	10/16/2019	
1		25 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:68210-1280-0	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/16/2019	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	10/16/2019		

Labeler - Spirit Pharmaceutical LLC (179621011)

Revised: 12/2024

Spirit Pharmaceutical LLC