

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

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

PRINCIPAL DISPLAY PANEL

Allergy Relief
Diphenhydramine HCl 25 mg

Antihistamine

Sneezing, Runny Nose, Itchy Throat,

Itchy, Watery Eyes

36 tablets

VALUHEALTH
BY SPIRIT

allergy relief

Diphenhydramine HCl 25 mg - Antihistamine

Relief of:
sneezing, runny nose, Itchy throat, Itchy, watery eyes

36 tablets

DO NOT USE IF SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

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

Directions ■ 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 tablets	
children 6 to under 12 years of age	1 tablet	
children under 6 years of age	do not use	

Other information
 ■ each tablet contains calcium 24 mg
 ■ store between 20-25° C (68-77° F). Protect from light.

Questions or comments? 1-888-333-9792

Distributed By: Spirit Pharmaceuticals, LLC
Ronkonkoma, NY 11779 REV 05/23

88905-5 Made in India

LOT:
EXP:

VALUHEALTH
BY SPIRIT

allergy relief

Diphenhydramine HCl 25 mg - Antihistamine

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

36 tablets

Actual Size

88905-5



8 50026 88905 5

LOT:
EXP:

Compare to BENADRYL® Allergy ULTRATAB® Tablets active ingredient!

Drug Facts (continued)

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

†This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Benadryl®.

TAMPER EVIDENT - DO NOT USE IF CARTON IS OPENED OR SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

Drug Facts

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Drug Facts (continued)

■ 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 (1-800-222-1222).

Directions
 ■ 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 tablets	
children 6 to under 12 years of age	1 tablet	
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ALLERGY RELIEF

diphenhydramine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68210-4127
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: 3WJQ05DW1A)	
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:68210-4127-1	1 in 1 CARTON	04/27/2021	
1		36 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	04/27/2021	

Labeler - Spirit Pharmaceutical LLC (179621011)

Revised: 12/2023

Spirit Pharmaceutical LLC