

ALLERGY RELIEF- diphenhydramine hydrochloride tablet, coated
VALU MERCHANDISERS COMPANY

1090-BST-2021-1116

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
 - sneezing
 - itchy, watery eyes
 - 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

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

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 (1-800-222-1222).

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 tablet
children under 6 years	do not use

Other information

- **each tablet contains:** calcium 45 mg
- store between 20-25°C (68-77°F)
- retain carton for complete product information

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, D&C red #27 aluminum lake, dibasic calcium phosphate, FD&C yellow #6 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide

PRINCIPAL DISPLAY PANEL

COMPARE TO THE ACTIVE INGREDIENT IN BENADRYL® ALLERGY ULTRATABS®†

Best Choice®

Antihistamine

Allergy Relief

Actual Size

Diphenhydramine HCl

For Allergy Relief

Sneezing, Itchy, Watery Eyes, Runny Nose, Itchy Throat

24 TABLETS

25 MG EACH



BEST CHOICE



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F109001BST_R1

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†This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Benadryl® Allergy Ultratabs®

DO NOT USE IF BLISTER UNITS ARE TORN OR BROKEN

PROUDLY DISTRIBUTED BY:
 VALU MERCHANTS, CO.
 5000 KANSAS AVE
 KANSAS CITY, KS 66106

Best Choice.
100% Guaranteed

www.bestchoicebrand.com
 Report serious side effects to:
 881 Main Street, Lumberton, NJ 08048

Antihistamine Allergy Relief



COMPARE TO THE ACTIVE INGREDIENT IN BENADRYL® ALLERGY ULTRATABS†

Antihistamine Allergy Relief



Diphenhydramine HCl

For Allergy Relief
 Sneezing, Itchy, Watery Eyes,
 Runny Nose, Itchy Throat

24 TABLETS
 25 MG EACH

NC

NC



ALLERGY RELIEF

diphenhydramine hydrochloride tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63941-090
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
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	pink	Score	no score
Shape	OVAL	Size	11mm
Flavor		Imprint Code	25;052
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63941-090-01	2 in 1 CARTON	05/19/2017	01/31/2028
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		

2	NDC:63941-090-03	1 in 1 CARTON	02/27/2017	11/30/2027
2		100 in 1 BOTTLE, PLASTIC; 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	02/27/2017	01/31/2028

Labeler - VALU MERCHANDISERS COMPANY (868703513)

Revised: 12/2025

VALU MERCHANDISERS COMPANY