ALLERGY RELIEF- diphenhydramine hydrochloride tablet, coated
Chain Drug Marketing Association

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

1090 - QCH - 2014-1028

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 with any other product containing diphenhydramine, even one used on skin

Ask a doctor before use if you have

- glaucoma
- trouble urinating due to an enlarged prostate gland
- 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
- alcohol, sedatives, and tranquilizers may increase drowsiness
- excitability may occur, especially in children
- 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 overdose, get medical help or contact a Poison Control Center right away.

Directions

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

adults and children 12 years of age and over	25 mg to 50 mg (1 to 2 tablets)
children 6 to under 12 years of age	12.5 mg * to 25 mg (1 tablet)
children under 6 years of age	ask a doctor

* 12.5 mg dosage strength is not available in this package. Do not attempt to break tablets.

Other information

- each tablet contains: **calcium 45 mg**
- store at room temperature 15°-30°C (59°-86°F)
- retain carton for complete product information

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, D&C red #27, dibasic calcium phosphate, FD&C yellow #6, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, titanium dioxide

PRINCIPAL DISPLAY PANEL

NDC 63868-492-24

QUALITY CHOICE

†Compare to Active Ingredient in BENADRYL® Allergy

Allergy Relief

Antihistamine

Diphenhydramine HCl, 25 mg

For Relief of:

Sneezing

Itchy, Watery Eyes

Runny Nose

Itchy Throat

24 Tablets


NC
NC
NC
Drug Facts
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 ■ 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 with any other product containing diphenhydramine, even one used on skin
Ask a doctor before use if you have ■ glaucoma ■ trouble urinating due to an enlarged prostate gland
■ 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
■ alcohol, sedatives, and tranquilizers may increase drowsiness ■ excitability may occur, especially in children
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QUALITY CHOICE
Allergy Relief
Diphenhydramine HCl, 25 mg | Antihistamine
NDC 63868-492-24
QUALITY CHOICE
†Compare to Active Ingredient in BENADRYL® Allergy
Allergy Relief
Antihistamine
Diphenhydramine HCl, 25 mg
For Relief of:
Sneezing
Itchy, Watery Eyes
Distributed by C.D.M.A., Inc.©
43157 W. Nine Mile
Novi, MI 48376-0995
www.qualitychoice.com
Questions: 248-449-9300
F10900100H_R0
100% SATISFACTION GUARANTEED
BLISTER UNITS
OR BROKEN
6 35515 94576 0

ALLERGY RELIEF

diphenhydramine hydrochloride tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-492
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)	
CROSCARMELLOSE 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: 3WQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

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:63868-492-24	2 in 1 CARTON	06/13/2014	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:63868-492-10	1 in 1 CARTON	06/13/2014	
2		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:63868-492-03	1 in 1 CARTON	06/22/2014	
3		300 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 final	part341	06/13/2014	

Labeler - Chain Drug Marketing Association (011920774)

Revised: 5/2023 Chain Drug Marketing Association