ALLERGY RELIEF- diphenhydramine hydrochloride capsule
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.

QCH - 1113 - 2019-1004

Drug Facts

Active ingredient (in each capsule)

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 capsules)
children 6 to under 12 years of age	12.5 mg * to 25 mg (1 capsule)
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 capsules.

Other information

- store at room temperature 15°-30°C (59°-86°F)
- protect from moisture
- retain carton for complete product information

Inactive ingredients

benzyl alcohol, butylparaben, D&C red #28, edible black ink, FD&C blue #1, FD&C red #40, gelatin, lactose, magnesium stearate, methylparaben, polysorbate 80, propylparaben, sodium lauryl sulfate

PRINCIPAL DISPLAY PANEL

NDC 63868-087-24

QUALITY CHOICE

†Compare to BENADRYL® Allergy Kapseals active ingredient

Allergy Relief

Antihistamine

Diphenhydramine HCl

For Relief of:

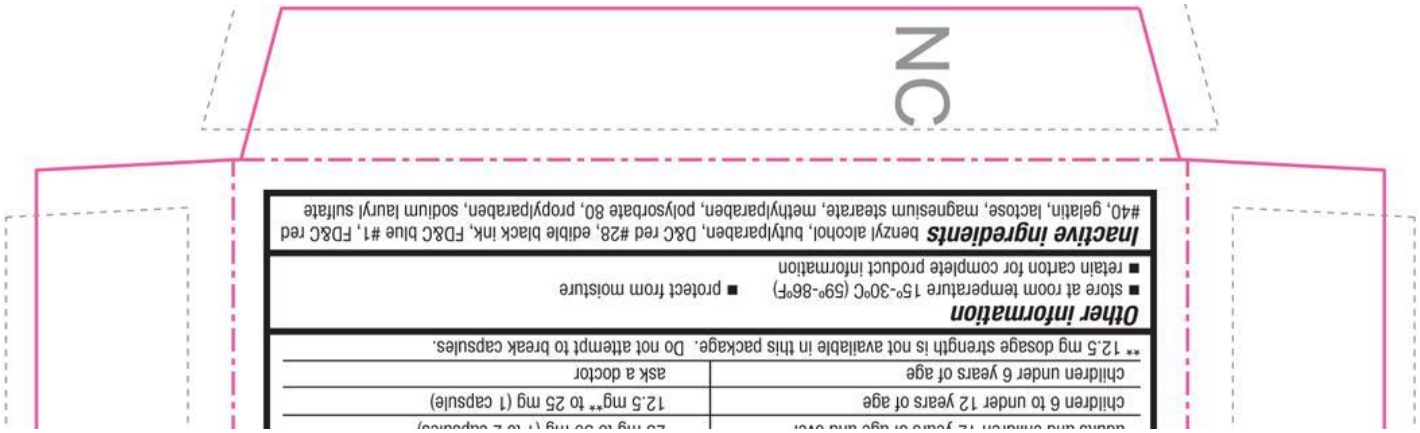
Sneezing

Itchy, Watery Eyes

Runny Nose

Itchy Throat

24 Capsules



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24 Capsules

©Distributed by C.D.M.A., Inc.
43157 W. Nine Mile
Novi, MI 48376-0995
www.qualitychoice.com
Questions: 248-449-9300

SATISFACTION GUARANTEED

F111301QCH_R2

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

DO NOT USE IF BLISTER UNITS ARE TORN OR BROKEN OR BAND AROUND ANY CAPSULE IS BROKEN OR MISSING

6 35515 90180 3

INK AND COATING FREE FOR LOT AND EXPIRATION STAMPING

QUALITY CHOICE

Allergy Relief

Antihistamine

ALLERGY RELIEF

diphenhydramine hydrochloride capsule

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-087
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**
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Product Characteristics			
Color	pink, white	**Score**	no score
Shape	OVAL	**Size**	14mm
Flavor		**Imprint Code**	A;20
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63868-087-24	2 in 1 CARTON	11/10/2008	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:63868-087-01	1 in 1 CARTON	11/10/2008	
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 final	part341	11/10/2008	

Labeler - Chain Drug Marketing Association (011920774)

Revised: 10/2019 Chain Drug Marketing Association