ALLERGY RELIEF- diphenhydramine hydrochloride capsule
Chain Drug Marketing Association

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:

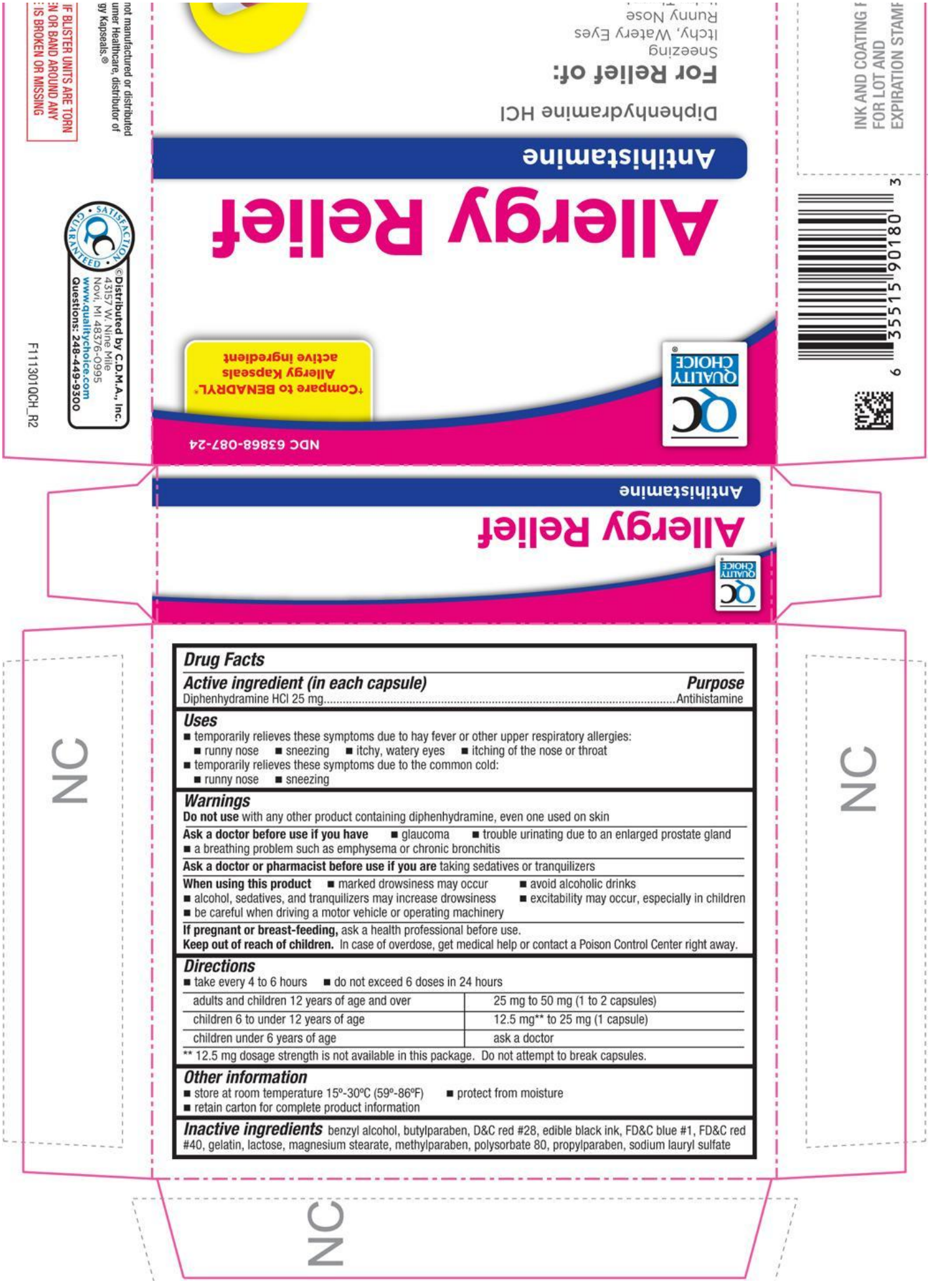
Sneezing

Itchy, Watery Eyes

Runny Nose

Itchy Throat

24 Capsules


Allergy Relief
Antihistamine
Diphenhydramine HCl
For Relief of:
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NDC 63868-087-24
QUALITY CHOICE
Allergy Relief
Antihistamine
Drug Facts
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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
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Ask a doctor before use if you have ■ glaucoma ■ trouble urinating due to an enlarged prostate gland
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NC
NC
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IS BROKEN OR MISSING
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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	04/30/2027
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 Drug	M012	11/10/2008	

Labeler - Chain Drug Marketing Association (011920774)

Revised: 10/2024 Chain Drug Marketing Association