

**DIPHENHYDRAMINE HYDROCHLORIDE- diphenhydramine
hydrochloride capsule
Major Pharmaceuticals**

0835K- Major

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
- 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
- glaucoma
- 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 doses in 24 hours

adults and children 12 years of age and over	1 to 2 capsules
children 6 to under 12 years of age	1 capsule
children under 6 years of age	do not use this product in children under 6 years of age

Other information

□ store in a dry place at 15° - 30°C (59° - 86°F)

corn starch, D&C red #28, FD&C blue #1, FD&C red #40, gelatin, lactose monohydrate, magnesium stearate, sodium lauryl sulfate

Questions or comments?

1-800-616-2471

Distributed by: MAJOR® PHARMACEUTICALS, Indianapolis, IN 46268

*This product is not manufactured or distributed by Kenvue Inc., owner of the registered trademark Benadryl®.

RETAIN CARTON FOR COMPLETE PRODUCT INFORMATION

DO NOT USE IF CARTON IS OPENED OR IF A BLISTER UNIT IS TORN, BROKEN, OR SHOWS ANY SIGNS OF TAMPERING.

DO NOT USE IF RED CAPSULE BAND IS BROKEN OR MISSING.

Major

BENOPHEN

NDC 0904-7237-24

Diphenhydramin HCl 25 mg

Antihistamine

For the temporary relief of the symptoms of:

- Upper Respiratory Allergies
- Hay Fever

Compare to the active ingredient in BENADRYL® Allergy*

24 CAPSULES

EACH CAPSULE INDIVIDUALLY
BANDED FOR YOUR PROTECTION

MAJOR[®]

NDC 0904-7237-24

compare to the active ingredient of Benadryl[®] Allergy*

banophen[™]

diphenhydramine HCl 25 mg
antihistamine

for the temporary relief from symptoms of:

- upper respiratory allergies
- hay fever

24 capsules



each capsule individually banded for your protection

R53756

MAJOR[®]

NDC 0904-7237-24

banophen[™]

diphenhydramine HCl 25 mg

NDC 0904-7237-24



Rev. 03/25 M-29 Re-order No. 701309

INDIANAPOLIS, IN 46268
MAJOR[®] PHARMACEUTICALS
(800) 616-2471
www.major-rugby.com

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Questions or comments? 1-800-231-4670

Inactive ingredients corn starch, D&C red #28, FD&C blue #1, FD&C red #40, gelatin, lactose monohydrate, magnesium stearate, sodium lauryl sulfate

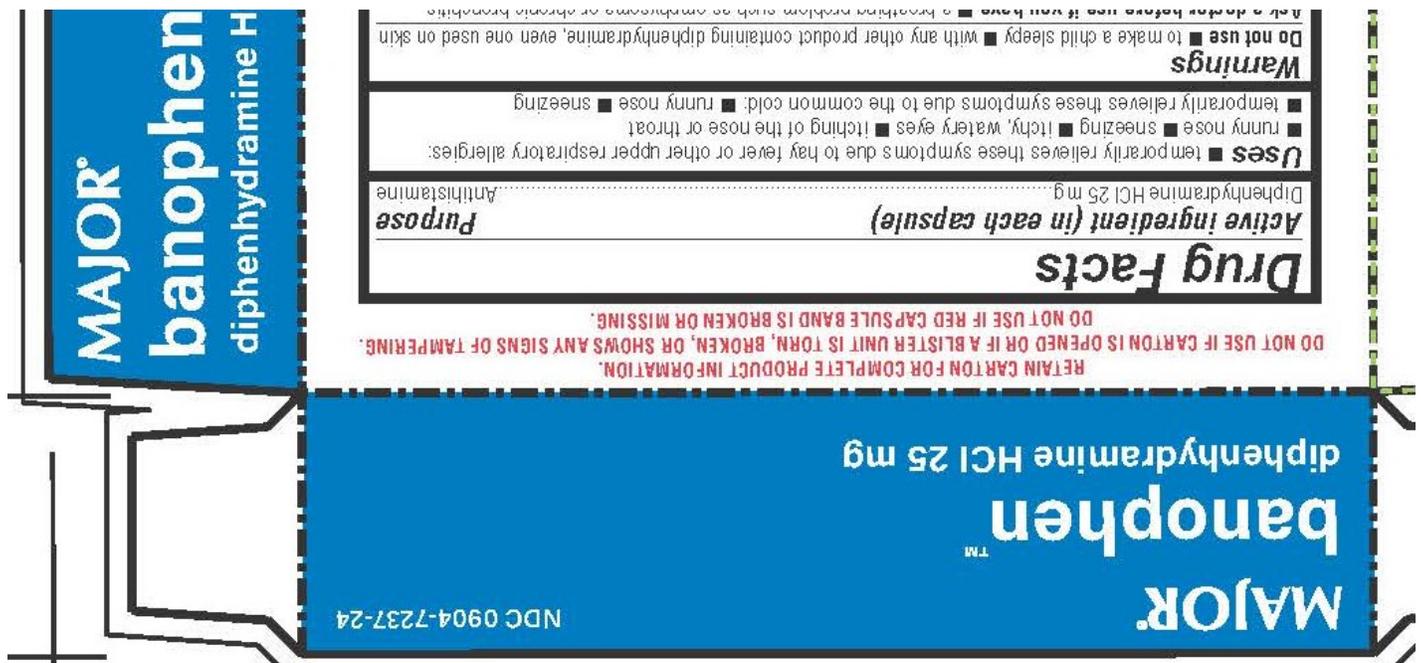
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■ glaucoma ■ trouble urinating due to an enlarged prostate gland
When using this product ■ marked drowsiness may occur ■ avoid alcoholic drinks ■ alcohol, sedatives and tranquilizers may increase drowsiness ■ be careful when driving a motor vehicle 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).

™

Cl 25 mg



DIPHENHYDRAMINE HYDROCHLORIDE

diphenhydramine hydrochloride capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-7237
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
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	pink (Half pink and half clear with white powder inside and sealed with red band)	Score	no score
Shape	CAPSULE	Size	14mm

Flavor		Imprint Code	CPC;835	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-7237-80	1000 in 1 BOTTLE; Type 0: Not a Combination Product	04/14/2022	04/30/2028
2	NDC:0904-7237-60	100 in 1 BOTTLE; Type 0: Not a Combination Product	04/14/2022	02/29/2028
3	NDC:0904-7237-24	2 in 1 CARTON	04/14/2022	01/31/2028
3		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:0904-7237-61	10 in 1 BOX, UNIT-DOSE	04/14/2022	02/29/2028
4		10 in 1 BLISTER PACK; 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	04/14/2022	04/30/2028	

Labeler - Major Pharmaceuticals (191427277)

Revised: 2/2026

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