

ALLERGY RELIEF- diphenhydramine hcl capsule
Chain Drug Consortium

Premier Value 44-190

Active ingredient (in each banded capsule)

Diphenhydramine HCl 25 mg

Purpose

Antihistamine

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
 - runny nose
 - itchy, watery eyes
 - sneezing
 - 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

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- difficulty in urination due to enlargement of the 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 beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- use caution 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.

Directions

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

adults and children 12 years and over	1 to 2 capsules
children 6 to under 12 years	1 capsule
children under 6 years	do not use

Other information

- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- protect from moisture
- see end flap for expiration date and lot number

Inactive ingredients

butylparaben, corn starch, D&C red #28, edible ink, FD&C blue #1, FD&C red #40, gelatin, lactose anhydrous, magnesium stearate, methylparaben, polysorbate 80, propylparaben, silicon dioxide

Questions or comments?

1-800-426-9391

Principal Display Panel

**Premier
Value®**

*COMPARE TO THE ACTIVE INGREDIENT
IN BENADRYL®

Allergy Relief

Diphenhydramine HCl, 25 mg

ANTIHISTAMINE

Allergy relief for:

- sneezing

- runny nose
- itchy, watery eyes
- itchy throat

actual size

100 Capsules

PV

INDEPENDENTLY TESTED

SATISFACTION GUARANTEED

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL
UNDER CAP IS BROKEN OR MISSING OR IF RED BAND
AROUND CAPSULE IS BROKEN OR MISSING**

*This product is not manufactured or distributed
by Johnson & Johnson Corporation, owner of the
registered trademark Benadryl®.

50844 REV1117B19012

Distributed By:

Pharmacy Value Alliance, LLC

**407 East Lancaster Avenue,
Wayne, PA 19087**

**If for any reason you are
not satisfied with this product,
please return it to the store
where purchased for a full refund.**



Premier Value 44-190

ALLERGY RELIEF

diphenhydramine hcl capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-640
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		
BUTYLPARABEN (UNII: 3QPI1U3FV8)				
STARCH, CORN (UNII: O8232NY3SJ)				
D&C RED NO. 28 (UNII: 767IP0Y5NH)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)				
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
METHYLPARABEN (UNII: A2I8C7HI9T)				
POLYSORBATE 80 (UNII: 6OZP39ZG8H)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
Product Characteristics				
Color	pink, white	Score	no score	
Shape	CAPSULE	Size	14mm	
Flavor		Imprint Code	44;107	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-640-24	2 in 1 CARTON	03/15/1990	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:68016-640-10	1 in 1 CARTON	03/15/1990	
2		100 in 1 BOTTLE; 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	03/15/1990	

Labeler - Chain Drug Consortium (101668460)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(68016-640)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	pack(68016-640)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(68016-640)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(68016-640)

Revised: 9/2023

Chain Drug Consortium