

**ALLERGY RELIEF- diphenhydramine hcl capsule**  
**Rite Aid Corporation**

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**Rite Aid 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
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic beverages
- 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**

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN OR IF RED BAND AROUND CAPSULE IS BROKEN OR MISSING**
- 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**

**NDC 11822-0190-8**

Compare to the active ingredient in **Benadryl®\***

**ALLERGY RELIEF**

DIPHENHYDRAMINE HCl 25 mg  
ANTIHISTAMINE

RELIEF OF  
Sneezing • Runny nose  
Itchy, watery eyes  
Itchy throat

ACTUAL SIZE

**24** CAPSULES

EACH CAPSULE INDIVIDUALLY BANDED FOR YOUR PROTECTION

**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 Kenvue, Inc., owner  
of the registered trademark Benadryl®.  
50844 REV1117E19008

**DISTRIBUTED BY:**

RITE AID,  
200 NEWBERRY COMMONS  
ETTERS, PA 17319 [www.riteaid.com](http://www.riteaid.com)

**SATISFACTION  
GUARANTEE**

If you're not satisfied, we'll  
happily refund your money.

# ALLERGY RELIEF

DIPHENHYDRAMINE HCl 25 mg

ANTIHISTAMINE

NDC 11822-0190-8

Compare to the active ingredient in Benadryl<sup>®</sup>\*

# ALLERGY RELIEF

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RELIEF OF  
Sneezing • Runny nose  
Itchy, watery eyes  
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200 NELUBERRY COMMONS  
EITERS, PA 17319 [www.riteaid.com](http://www.riteaid.com)

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or distributed by Kenec, Inc., owner  
of the registered trademark Benadryl<sup>®</sup>.  
50844 REV117E19008

no print / no varnish area  
lot no. & exp. date

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED  
OR IF BLISTER UNIT IS TORN OR BROKEN OR IF  
RED BAND AROUND CAPSULE IS BROKEN OR MISSING



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# ALLERGY RELIEF

DIPHENHYDRAMINE HCl 25 mg

ANTIHISTAMINE

B-1702-190-08-R3  
REV117E19008

**Warnings**

- sneezing ■ itching of the nose or throat
- temporarily relieves these symptoms due to the common cold: ■ runny nose ■ sneezing
- 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 ■ alcohol, sedatives, and tranquilizers may increase drowsiness ■ avoid alcoholic beverages ■ use caution when driving a motor vehicle or operating machinery ■ excitability may occur, especially in children

**Other information**

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**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



## Rite Aid 44-190

### ALLERGY RELIEF

diphenhydramine hcl capsule

#### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11822-0190
<b>Route of Administration</b>	ORAL		

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

#### Inactive Ingredients

Ingredient Name	Strength
<b>BUTYLPARABEN</b> (UNII: 3QPI1U3FV8)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>D&amp;C RED NO. 28</b> (UNII: 767IP0Y5NH)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GELATIN, UNSPECIFIED</b> (UNII: 2G86QN327L)	
<b>ANHYDROUS LACTOSE</b> (UNII: 3SY5LH9PMK)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	

#### Product Characteristics

<b>Color</b>	pink, white	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	14mm
<b>Flavor</b>		<b>Imprint Code</b>	44;107
<b>Contains</b>			

#### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-0190-8	2 in 1 CARTON	03/15/1990	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11822-0190-1	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** - Rite Aid Corporation (014578892)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(11822-0190)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	pack(11822-0190)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(11822-0190)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(11822-0190)

Revised: 8/2025

Rite Aid Corporation