

**ALLERGY RELIEF- diphenhydramine hcl tablet, film coated**  
**Rite Aid Corporation**

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**Rite Aid 44-329**

***Active ingredient (in each tablet)***

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**

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- 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**

- **do not take more than directed**
- take every 4 to 6 hours, or as directed by a doctor
- do not take more than 6 times in 24 hours

adults and children 12 years and over	1 to 2 tablets
children 6 to under 12 years	1 tablet
children under 6 years	do not use

**Other information**

- **each tablet contains:** calcium 30 mg
- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- protect from moisture
- store at 25°C (77°F); excursions permitted between 15°C-30°C (59°F-86°F)
- see end flap for expiration date and lot number

**Inactive ingredients**

corn starch, D&C red #27 aluminum lake, dibasic calcium phosphate dihydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, silicon dioxide, stearic acid, talc, titanium dioxide

**Questions or comments?**

**1-800-426-9391**

**Principal Display Panel**

Compare to the active ingredient in  
**Benadryl® Allergy ULTRATAB® Tablets\***

NDC 11822-0329-8

**ALLERGY RELIEF**  
DIPHENHYDRAMINE HCl 25 mg

**ANTIHISTAMINE**

RELIEF OF

Sneezing • Runny nose  
Itchy, watery eyes  
Itchy throat

ACTUAL SIZE

**24**

MINITABS

25 mg EACH

**TAMPER EVIDENT: DO NOT USE IF PACKAGE IS  
OPENED OR IF BLISTER UNIT IS TORN, BROKEN  
OR SHOWS ANY SIGNS OF TAMPERING**

\*This product is not  
manufactured or distributed  
by Johnson & Johnson  
Corporation, owner of the  
registered trademark  
Benadryl® Allergy  
ULTRATAB® Tablets.  
50844 REV0721H32908

**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 11822-0329-8

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# ALLERGY RELIEF

DIPHENHYDRAMINE HCl 25 mg

## ANTIHISTAMINE

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

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MINITABS  
25 mg EACH

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50844 REV0721H32908

B-1702-329-08-R3  
REV0721H32908

**Warnings**

- runny nose ■ itchy, watery eyes
- 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.

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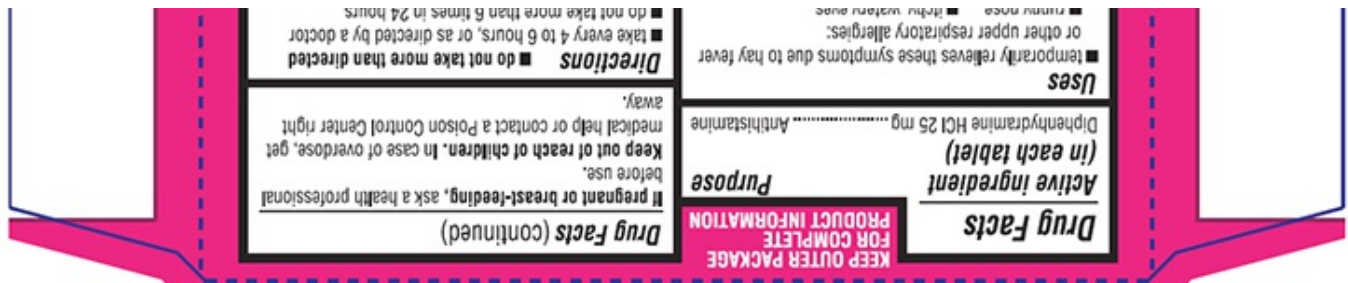
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**Rite Aid 44-329**

**ALLERGY RELIEF**

diphenhydramine hcl tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11822-0329
<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>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>D&amp;C RED NO. 27 ALUMINUM LAKE</b> (UNII: ZK64F7XSTX)	
<b>DIBASIC CALCIUM PHOSPHATE DIHYDRATE</b> (UNII: O7TSZ97GEP)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POLYVINYL ALCOHOL, UNSPECIFIED</b> (UNII: 532B59J990)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

**Product Characteristics**

<b>Color</b>	pink	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	44;329
<b>Contains</b>			

**Packaging**

		<b>Marketing Start</b>	<b>Marketing End</b>
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-0329-8	2 in 1 CARTON	03/02/1990	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11822-0329-4	4 in 1 CARTON	03/02/1990	
2		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:11822-0329-2	1 in 1 CARTON	03/02/1990	
3		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:11822-0329-6	1 in 1 CARTON	03/02/1990	
4		200 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
5	NDC:11822-0329-1	365 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/02/1990	
6	NDC:11822-0329-5	1 in 1 CARTON	03/02/1990	10/08/2021
6		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
7	NDC:11822-0329-7	3 in 1 CARTON	03/02/1990	09/09/2017
7		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
8	NDC:11822-0329-3	1 in 1 PACKAGE	03/02/1990	10/08/2019
8		10 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
9	NDC:11822-0329-9	1 in 1 CARTON	03/02/1990	10/16/2021
9		2 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	03/02/1990	

**Labeler** - Rite Aid Corporation (014578892)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(11822-0329)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(11822-0329) , pack(11822-0329)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(11822-0329)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(11822-0329)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(11822-0329)

Revised: 6/2024

Rite Aid Corporation