

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

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

ANTI-HISTAMINE

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

**SATISFACTION
GUARANTEE**

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

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DIPHENHYDRAMINE HCl 25 mg
ANTIHISTAMINE

NDC 11822-0329-8

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B-1702-329-08-R3
REV0721H32908

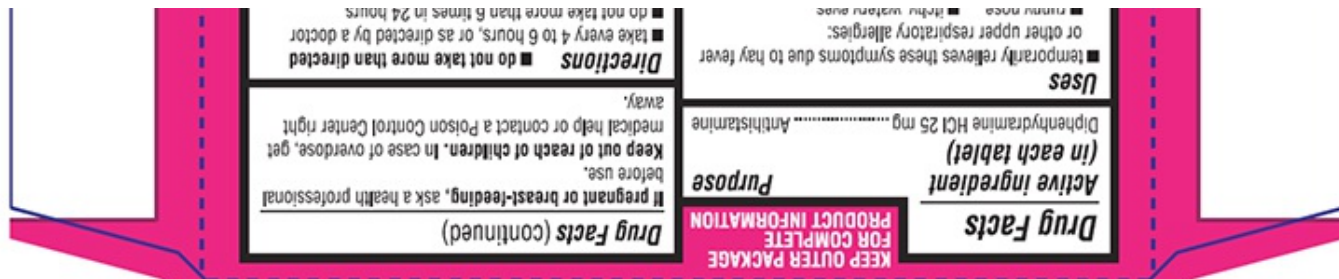
Questions or comments? 1-800-426-9391

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- Warnings**
- sneezing ■ itchy of the nose or throat ■ runny nose ■ itchy, watery eyes
 - 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 ■ 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



Rite Aid 44-329

ALLERGY RELIEF

diphenhydramine hcl tablet, film coated

Product Information

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

Product Characteristics

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

Packaging

#	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: 6/2025

Rite Aid Corporation