

**ALLERGY RELIEF- diphenhydramine hcl tablet, film coated  
Walgreen Company**

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**Walgreens 44-329 Allergy Relief**

***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**

- 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**

- **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
- 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**

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**

**Walgreens**

• **WALGREENS** •  
**PHARMACIST RECOMMENDED†**

Compare to the active ingredient in Benadryl® Allergy ULTRATABS®††

**Allergy Relief**

DIPHENHYDRAMINE HCl 25 mg / ANTIHISTAMINE

- Relief of runny nose, sneezing, itchy throat & itchy, watery eyes

**100**  
COATED  
MINI TABS

ACTUAL SIZE

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

†Our pharmacists recommend the Walgreens brand.  
We invite you to compare to national brands.

††This product is not manufactured or distributed  
by Kenvue Inc., owner of the registered trademarks  
Benadryl® and ULTRATABS®.

50844 REV0721A32912

NDC 0363-3291-12

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**DEERFIELD, IL 60015**  
**100% SATISFACTION GUARANTEED**  
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**ALLERGY RELIEF**

diphenhydramine hcl tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0363-3291
<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

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-3291-08	2 in 1 CARTON	09/27/2021	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0363-3291-22	4 in 1 CARTON	09/27/2021	
2		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:0363-3291-07	3 in 1 CARTON	12/30/2024	
3		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:0363-3291-12	1 in 1 CARTON	09/27/2021	
4		100 in 1 BOTTLE, PLASTIC; 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	09/27/2021	

**Labeler** - Walgreen Company (008965063)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(0363-3291)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(0363-3291) , pack(0363-3291)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(0363-3291)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(0363-3291)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(0363-3291)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117597853	pack(0363-3291)

Revised: 2/2026

Walgreen Company