

**ALLERGY RELIEF CHILDRENS- diphenhydramine hcl tablet, chewable**  
**Chain Drug Consortium**

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**Premier Value 44-599-Childrens-Delisted**

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

Diphenhydramine HCl 12.5 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

***Warnings***

**Do not use**

- with any other product containing diphenhydramine, even one used on skin
- to make a child sleepy

**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
- alcohol, sedatives, and tranquilizers may increase drowsiness
- use caution when driving a motor vehicle or operating machinery
- avoid alcoholic beverages
- 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 (1-800-222-1222) right away.

### **Directions**

- find right dose on chart below
- chew or crush tablets completely before swallowing; do not swallow tablets whole
- take every 4 to 6 hours, or as directed by a doctor
- do not take more than 6 times in 24 hours

<b>Age (yr)</b>	<b>Dose (chewable tablets)</b>
children under 2 years	do not use
children 2 to 5 years	do not use unless directed by a doctor
children 6 to 11 years	1 to 2 chewable tablets (12.5 mg to 25 mg)
adults and children 12 years and over	2 to 4 chewable tablets (25 mg to 50 mg)

### **Other information**

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- avoid high humidity
- see end flap for expiration date and lot number

### **Inactive ingredients**

D&C red #27 aluminum lake, D&C red #30 aluminum lake, dextrans hydrated, ethylcellulose, flavor, hydroxypropyl cellulose, magnesium stearate, mannitol, stearic acid, sucralose, sucrose

### **Questions or comments?**

**1-800-426-9391**

### **Principal display panel**

**Premier  
Value®**

\*COMPARE TO THE ACTIVE INGREDIENT  
IN CHILDREN'S BENADRYL® CHEWABLES

**Children's  
Allergy Relief**

Diphenhydramine HCl 12.5 mg Chewable Tablets

**ANTIHISTAMINE**

Relief for:

- Sneezing
- Itchy throat
- Runny nose
- Itchy, watery eyes

**Chew or crush tablets completely  
before swallowing.**

**18 Chewable  
Tablets**

Cherry flavored

**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  
Children's Benadryl® Chewables.

50844 REV1218A59944

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.



**Children's Allergy Relief**  
Diphenhydramine HCl 12.5 mg  
**ANTIHISTAMINE**



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

B-1590-599-44-R  
REV1218A59944

This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Children's Benadryl® Chewables, 50844 REV1218A59944  
Distributed By:  
Pharmacy Value Alliance, LLC  
407 East Lancaster Avenue,  
Wayne, PA 19087

**Drug Facts (continued)**  
Flavor, hydroxypropyl cellulose, magnesium stearate, mannitol, stearic acid, sucralose, sucrose

**Drug Facts (continued)**  
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

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**Drug Facts**  
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**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

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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 ■ alcohol, sedatives, and tranquilizers may increase drowsiness ■ use caution when driving a motor vehicle or operating machinery ■ avoid alcoholic beverages ■ excitability may occur, especially in children  
If pregnant or breast-feeding, ask a health professional before use.

**Purpose**  
Antihistamine

**KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION**

# ALLERGY RELIEF CHILDRENS

diphenhydramine hcl tablet, chewable

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68016-051
<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	12.5 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>D&amp;C RED NO. 27 ALUMINUM LAKE</b> (UNII: ZK64F7XSTX)	
<b>D&amp;C RED NO. 30</b> (UNII: 2S42T2808B)	
<b>DEXTROSE MONOHYDRATE</b> (UNII: LX22YL083G)	
<b>ETHYLCELLULOSE, UNSPECIFIED</b> (UNII: 7Z8S9VYZ4B)	
<b>HYDROXYPROPYL CELLULOSE, UNSPECIFIED</b> (UNII: 9XZ8H6N6OH)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6130)	
<b>MANNITOL</b> (UNII: 3OWL53L36A)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>SUCROSE</b> (UNII: C151H8M554)	

## Product Characteristics

<b>Color</b>	pink	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	12mm
<b>Flavor</b>	CHERRY	<b>Imprint Code</b>	44;599
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-051-18	3 in 1 CARTON	04/25/2011	06/30/2026
1		6 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
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OTC Monograph Drug	M012	04/25/2011	06/30/2026
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**Labeler - Chain Drug Consortium (101668460)**

**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(68016-051) , pack(68016-051)

**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(68016-051)

Revised: 7/2025

Chain Drug Consortium