

CHILDRENS ALLERGY RELIEF- diphenhydramine hcl tablet, chewable
Rite Aid Corporation

Rite Aid 44-599

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
- 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

- 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

Age (yr)	Dose (chewable tablets)
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°C-30°C (59°F-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

Compare to the active ingredient in **Children's Benadryl® Chewables***

NDC 11822-0599-4

6 YEARS & OLDER

CHILDREN'S

ALLERGY RELIEF

DIPHENHYDRAMINE HCl 12.5 mg

CHEWABLE TABLETS

ANTIHISTAMINE

for relief of: sneezing • runny nose
itchy, watery eyes
itchy throat or nose

ACTUAL SIZE

CHERRY
FLAVOR

18
CHEWABLE TABLETS

CONTAINS ONE
ACTIVE INGREDIENT

Chew or crush tablets
completely before swallowing.

**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
REV1218E59944

DISTRIBUTED BY:
RITE AID, 30 HUNTER LANE,
CAMP HILL, PA 17011
www.riteaid.com

**SATISFACTION
GUARANTEE**

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

6 YEARS & OLDER

CHILDREN'S ALLERGY RELIEF

DIPHENHYDRAMINE HCl 12.5 mg
CHEWABLE TABLETS
ANTIHISTAMINE

Compare to the active ingredient in Children's Benadryl® Chewables*

NDC 11822-0599-4

6 YEARS & OLDER

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CHERRY FLAVOR

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No print/No varnish area
Lot no/Exp date

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flavor, hydroxypropyl cellulose, magnesium stearate, mannitol, stearic acid, sucralose, sucrose

Drug Facts (continued)

B-1702-599-44-R2
REV1218E59944

Drug Facts

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

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 ■ 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

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Drug Facts (continued)

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

■ 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

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Inactive ingredients

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CHILDRENS ALLERGY RELIEF

diphenhydramine hcl tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-0599
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	12.5 mg

Inactive Ingredients

Ingredient Name	Strength
D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)	
D&C RED NO. 30 (UNII: 2S42T2808B)	
DEXTROSE MONOHYDRATE (UNII: LX22YL083G)	
ETHYLCELLULOSE, UNSPECIFIED (UNII: 7Z8S9VYZ4B)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-0599-4	3 in 1 CARTON	04/25/2011	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11822-0599-2	1 in 1 CARTON	04/25/2011	09/16/2021
2		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	04/25/2011	

Labeler - Rite Aid Corporation (014578892)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(11822-0599) , pack(11822-0599)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(11822-0599)

Revised: 7/2023

Rite Aid Corporation