CHILDRENS ALLERGY RELIEF- diphenhydramine hcl tablet, chewable Target Corporation

Target 44-585 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

- 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 (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

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°-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, dextrates hydrated, ethylcellulose, FD&C blue #1 aluminum lake, flavor, hydroxypropyl cellulose, magnesium stearate, mannitol, stearic acid, sucralose, sucrose

Questions?

Call 1-800-910-6874

Principal display panel

NDC 11673-985-09

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

children's allergy relief diphenhydramine HCl, 12.5 mg chewable tablets antihistamine

relieves:

- sneezing
- itchy, watery eyes
- runny nose
- itchy throat or nose

chew or crush tablets completely before swallowing.

up&up™

20 CHEWABLE TABLETS

GRAPE FLAVOR

ACTUAL SIZE

CHEWABLE TABLETS

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

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Target 44-585

CHILDRENS ALLERGY RELIEF

diphenhydramine hcl tablet, chewable

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-985
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)

FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)

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	purple	Score	no score
Shape	ROUND	Size	12mm
Flavor	GRAPE	Imprint Code	44;585
Contains			

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-985- 09	4 in 1 CARTON	04/01/2019	05/24/2024
1		5 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	04/01/2019	05/24/2024	

Labeler - Target Corporation (006961700)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(11673-985)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	pack(11673-985)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(11673-985)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	pack(11673-985)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(11673-985)

Revised: 10/2023 Target Corporation