CHILDRENS ALLERGY RELIEF DYE-FREE- diphenhydramine hcl liquid Target Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Target 44-018-Liquid

Active ingredient (in each 5 mL)

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 the child has

- a breathing problem such as chronic bronchitis
- glaucoma

Ask a doctor or pharmacist before use if the child is

taking sedatives or tranquilizers.

When using this product

- marked drowsiness may occur
- sedatives and tranquilizers may increase drowsiness
- excitability may occur, especially in children

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
- mL = milliliter; FL OZ = fluid ounce
- use only enclosed dosing cup designed for use with this product. Do not use any other dosing device.

- take every 4 to 6 hours, or as directed by a doctor
- do not take more than 6 doses in 24 hours

Age (yr)	Dose (mL)
children under 2 years	do not use
children I to 5 warre	do not use unless
	directed by a doctor
children 6 to 11 years	5 mL to 10 mL

Other information

- each 5 mL contains: sodium 4 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

anhydrous citric acid, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sodium chloride, sodium citrate dihydrate, sucralose, xanthan gum

Questions or comments?

Call 1-800-910-6874

Principal Display Panel

NDC 11673-918-36

Compare to active ingredient in

Children's Benadryl® Dye-Free Allergy*

dye-free

Children's allergy relief

diphenhydramine HCl 12.5 mg antihistamine/5 mL oral solution

alcohol and sugar free

for relief of:

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

up & up

BUBBLEGUM FLAVOR AGES 6-11 YEARS

4 FL OZ (118 mL)

*This product is not manufactured or distributed by

Johnson & Johnson Corporation, owner of the registered trademark Children's Benadryl® Dye-Free Allergy. 50844 ORG121701836

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

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Target 44-018

CHILDRENS ALLERGY RELIEF DYE-FREE diphenhydramine hcl liquid Product Information Product Type HUMAN OTC DRUG Route of Administration HUMAN OTC DRUG ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DIPHENHYDRAMINE HYDRO CHLO RIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	12.5 mg in 5 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)		
GLYCERIN (UNII: PDC6 A3C0 O X)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
TRISO DIUM CITRATE DIHYDRATE (UNII: B22547B95K)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		
XANTHAN GUM (UNII: TTV12P4NEE)		

Product Characteristics			
Color	WHITE (clear-colorless)	Score	
Shape		Size	
Flavor	BUBBLE GUM	Imprint Code	
Contains			

l	Packaging			
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:11673-918-36	1 in 1 CARTON	05/20/2018	
ı	1	118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part341	05/20/2018	

Labeler - Target Corporation (006961700)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	MANUFACTURE(11673-918), PACK(11673-918)

Revised: 12/2019 Target Corporation