

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

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 right away.

Directions

- **do not take more than directed**
- find right dose on chart below
- mL = milliliter
- only use the dose cup provided
- 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 2 to 5 years	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 per 5 mL oral solution/antihistamine

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 Kenvue Inc., owner of the registered trademark Children's Benadryl® Dye-Free Allergy.

50844 REV0123A01836

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Drug Facts KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

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Drug Facts (continued)

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B-1920-018-36
REV0123A01836

dye-free
children's allergy relief
diphenhydramine HCl, 12.5 mg per 5 mL oral solution/antihistamine



NDC 11673-918-36

Compare to active ingredient in **Children's Benadryl® Dye-Free Allergy***

dye-free children's allergy relief

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alcohol and sugar free

for relief of:

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





4 FL OZ (118 mL)

dye-free
children's allergy relief
diphenhydramine HCl, 12.5 mg per 5 mL oral solution/antihistamine



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C-000703-01-049



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No print/No varnish
Lot & Exp date

Target 44-018

CHILDRENS ALLERGY RELIEF DYE-FREE			
diphenhydramine hcl solution			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-918

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 in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	BUBBLE GUM	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-918-36	1 in 1 CARTON	05/20/2018	
1		118 mL 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	05/20/2018	

Labeler - Target Corporation (006961700)

Establishment

Name	Address	ID/FEI	Business Operations
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Revised: 7/2024

Target Corporation