

**ALLERGY RELIEF CHILDRENS- diphenhydramine hcl solution
WALMART INC.**

Equate 44-018 RESERVED 79903-368

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
 - sneezing
 - itchy, watery eyes
 - 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

- glaucoma
- a breathing problem such as chronic bronchitis

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**
- do not take more than 6 doses in 24 hours
- mL = milliliter
- only use the dose cup provided
- find right dose on chart below
- take every 4 to 6 hours, or as directed by a doctor

| 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?

1-888-287-1915

Principal Display Panel

equate™

NDC 79903-368-04

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

children's
**ALLERGY
RELIEF**

Diphenhydramine HCl,
12.5 mg per 5 mL
Oral Solution

Antihistamine

Sugar-Free

Dye-Free

6-11 YEARS

Relieves:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Throat

or Nose

**4-6
HOURS/
DOSE**

Clear

Bubble Gum

Flavored

Dosage Cup

Included

4 FL OZ (118 mL)

**TAMPER EVIDENT: DO NOT USE IF PRINTED
NECK WRAP IS BROKEN OR MISSING**

DISTRIBUTED BY: Walmart Inc.

Bentonville, AR 72716

PRODUCT OF CHINA

*This product is not manufactured or distributed
by Kenvue Inc., owner of the registered trademark
Children's Benadryl® Dye-Free Allergy.

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Equate 44-018

ALLERGY RELIEF CHILDRENS

diphenhydramine hcl solution

| Product Information | | | | |
|---|--|--|----------------------|--------------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:79903-368 | |
| 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:79903-368-04 | 1 in 1 CARTON | 06/19/2025 | |
| 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 | | 06/19/2025 | |

Labeler - WALMART INC. (051957769)

| Establishment | | | |
|-------------------------|---------|-----------|--|
| Name | Address | ID/FEI | Business Operations |
| LNK International, Inc. | | 967626305 | manufacture(79903-368) , pack(79903-368) |

Revised: 6/2025

WALMART INC.