

**CHILDRENS ALLERGY- diphenhydramine hydrochloride liquid
QUALITY CHOICE (Chain Drug Marketing Association)**

Drug Facts

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:
- sneezing
- runny nose
- itchy, watery eyes
- itching of the nose or throat

Warnings

Do not use

- to make a child sleep
- 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
- a sodium-restricted diet

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

- do not take more than 6 doses in 24 hours
- take every 4 to 6 hours, or as directed by a doctor
- measure only with dosing cup provided. Do not use any other dosing device.
- mL = milliliter
- keep dosing cup with product
- find the right dose on the chart below

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

Other information

- each 5 mL contains: **sodium 6 mg**
- store between 20-25°C (68-77°F). Do not refrigerate.
- Protect from light. Store in outer carton until contents are used

Inactive ingredients

anhydrous citric acid, D&C red #33, FD&C red #40, flavor, glycerin, poloxamer 407, purified water, sodium benzoate, sodium chloride, sodium citrate, sucrose

Questions or comments?

call **1-248-449-9300 Monday-Friday 9AM-5PM EST**

Principal Display Panel

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

Children's Allergy

Oral Solution

Diphenhydramine HCl 12.5 mg

Antihistamine

For Ages 6 to 11 Years

Alcohol Free

Bubblegum Flavor

FL OZ (mL)

*This product is not manufactured or distributed by McNeil Consumer Healthcare distributor Benadryl® Allergy.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF PRINTED SAFETY SEAL AROUND BOTTLE OR UNDER CAP IS BROKEN OR MISSING.

Distributed by C.D.M.A., Inc.©

43157 W. Nine Mile

Novi, MI 48376-0995

www.qualitychoice.com

Questions: 248-449-9300

Package Label

Drug Facts (continued)
Inactive ingredients citric acid, D&C red #33, FD&C red #40, flavor, glycerin, poloxamer 407, purified water, sodium benzoate, sodium chloride, sodium citrate, sucrose
Questions or comments?
 Call 1-248-449-9300 Monday-Friday 9AM-5PM EST

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NDC 63868-370-04

QC QUALITY CHOICE

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

Children's Allergy

Oral Solution

Diphenhydramine HCl 12.5 mg
Antihistamine

For Ages 6 to 11 Years
Alcohol Free

Bubble Gum Flavor

4 FL OZ (118 mL)

NDC 63868-370-04

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Warnings	
<p>Do not use</p> <ul style="list-style-type: none"> ■ with any other product containing diphenhydramine, even one used on the skin ■ to make a child sleepy 	
<p>Ask a doctor before use if the child has</p> <ul style="list-style-type: none"> ■ a breathing problem such as chronic bronchitis ■ glaucoma ■ a sodium-restricted diet 	
<p>Ask a doctor or pharmacist before use if the child is taking sedatives or tranquilizers.</p>	
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PLD-F240C FC004037

Lot No.:

Exp. Date:

QUALITY CHOICE Children's Allergy

CHILDRENS ALLERGY

diphenhydramine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-370
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)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color	pink	Score	
Shape		Size	
Flavor	BUBBLE GUM	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-370-04	1 in 1 BOX	05/30/2014	
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 Drug	M012	05/30/2014	

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Revised: 4/2024

QUALITY CHOICE (Chain Drug Marketing Association)