DYE FREE ALLERGY CHILDRENS- 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

- with any other product containing diphenhydramine, even one used on skin
- to make a child sleepy

Ask a doctor before use if the child has

- a breathing problem such as chronic bronchitis
- glaucoma
- 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 8 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, carboxymethycellulose sodium, flavor, glycerin, purified water, saccharin sodium, sodium benzoate, sodium citrate, sorbitol

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

Dye-Free

Children's

Allergy

Antihistamine

Diphenhydramine Hydrochloride 12.5mg / 5 mL Oral Solution

Liquid Medication

Relieves

Runny Nose | Sneezing |

Itchy, Watery Eyes | Itchy Throat

For Ages 6 to 11 Years

Alcohol-Free

Dye-Free

Bubble Gum Flavored

FL OZ (mL)

*This product is not manufactured or distributed by McNeil Consumer Healthcare, distributors of 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

Package Label

Drug Facts (continued) Inactive ingredients carboxymethylcellulose sodium, citric acid, flavor, glycerin, purified water, saccharin sodium, sodium benzoate, sodium citrate, Questions or comments? Call 1-248-449-9300 Monday-Friday 9AM-5PM EST *This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Benadry!® 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, M 48375 www.qualitychoice.com Questions: 248-449-9300



Drug Facts

Active ingredient

Purpose .Antihistamin

(in each 5 mL) Diphenhydramine HCl 12.5 mg.

- 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

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 to make a child sleepy
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QUALITY CHOICE Dye-Free Children's Allergy

DYE FREE ALLERGY CHILDRENS

diphenhydramine hydrochloride liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63868-128

ORAL Route of Administration

Active Ingredient/Active Moiety

Basis of Strength Ingredient Name Strength **DIPHENHYDRAMINE HYDROCHLORIDE** (UNII: TC2D6JAD40) **DIPHENHYDRAMINE** 12.5 mg **HYDROCHLORIDE** (DIPHENHYDRAMINE - UNII:8GTS82S83M) in 5 mL

Inactive Ingredients			
Ingredient Name	Strength		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)			
GLYCERIN (UNII: PDC6A3C0OX)			
WATER (UNII: 059QF0KO0R)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)			
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
SORBITOL (UNII: 506T60A25R)			

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	BUBBLE GUM	Imprint Code	
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:63868- 128-04	1 in 1 BOX	02/29/2016		
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	02/29/2016	

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

Revised: 4/2024 QUALITY CHOICE (Chain Drug Marketing Association)