GOOD NEIGHBOR PHARMACY CHILDRENS ALLERGY- diphenhydramine hydrochloride solution Proficient Rx LP

Children's Allergy 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
- itching of the nose or throat
- runny nose
- itchy, watery eyes

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

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

- find right dose on chart below
- mL = milliliter
- 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

Attention: use only enclosed dosing cup specifically designed for use with this product. Do not use any other dosing device.

Other information

- each 5 mL (1 tsp) contains: sodium 15 mg
- store at 20-25°C (68-77°F). Protect from light. Store in outer carton until contents used.
- do not use if printed neckband is broken or missing

Inactive ingredients

anhydrous citric acid, D&C red #33, FD&C red #40, flavor, glycerin, high fructose corn syrup, poloxamer 407, purified water, sodium benzoate, sodium chloride, sodium citrate, sorbitol solution

Questions or comments?

1-800-719-9260

Principal Display Panel

Compare to Children's Benadryl® Allergy active ingredient

children's

Allergy

Antihistamine

12.5 mg/5 mL

diphenhydramine HCl

oral solution

For Allergy Relief

Runny Nose

Sneezing

Itchy, Watery Eyes

Itchy Throat

4-6 Hours/Dose

Alcohol Free

Cherry Flavored

4 fl oz (118 mL)

Relabeled by:

Proficient Rx LP

Thousand Oaks, CA 91320



Scan Here



NDC 71205-259-04

Relabeled By: Proficient Rx LP Thousand Oaks, CA 91320

Diphenhydramine 12.5mg/ 5mL

Lot #:00000

NDC 71205-259-04

4 fl oz (118 mL) Oral Solution SN# MASTER Exp:00/00/00

Diphenhydramine 12.5mg/ 5mL 4 fl oz (118 mL) Oral Solution Lot #:00000 SN# MASTER NDC 71205-259-04 Exp:00/00/00

Diphenhydramine 12.5mg/ 5mL 4 fl oz (118 mL) Oral Solution Lot #:00000 SN# MASTER NDC 71205-259-04 Exp:00/00/00



GTIN: 00371205259044 SN# MASTER Exp. 00/00/00 Lot #:00000

Diphenhydramine 12.5mg/ 5mL

4 fl oz (118 mL) Oral Solution

Each 5ml (1tsp) contains: Diphenhydramine HCI 12.5mg Antihistamine

See Box

Product ID: SD025904

Dist. By: AmerisourceBergen 1300 Morris Drive Chesterbrook, PA 19087 Store at 20°-25°C (68°-77°F)

Keep medication out of the reach of children

GOOD NEIGHBOR PHARMACY CHILDRENS ALLERGY

diphenhydramine hydrochloride solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71205-259(NDC:24385-379)

ORAL Route of Administration

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)		
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)		
POLOXAMER 407 (UNII: TUF2IVW3M2)		
WATER (UNII: 059QF0KO0R)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)		
SORBITOL (UNII: 506T60A25R)		

Product Characteristics			
Color	RED (Bluish-Red)	Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71205-259- 04	1 in 1 CARTON	05/01/2019	
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	08/15/1989	

Labeler - Proficient Rx LP (079196022)

Establishment			
Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(71205-259), RELABEL(71205-259)

Revised: 1/2024 Proficient Rx LP