

SILADRYL ALLERGY MEDICINE- diphenhydramine hydrochloride liquid

Proficient Rx LP

Siladryl Allergy Liquid Medicine

Active Ingredient: Diphenhydramine HCl 12.5 mg (in each 5 mL (teaspoonful)(TSP))

Purpose: Antihistamine

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
- - o runny nose
 - o sneezing
 - o itchy, watery eyes
 - o 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 you have

- glaucoma
- trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis
- a sodium restricted diet

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers

When using this product

- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

If pregnant or breast-feeding, ask a health professional before use.

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

- repeat dose every 4 to 6 hours
- do not take more than 6 doses in any 24-hour period
- Attention: use only enclosed dosing cup specifically designed for use with this product. Do not use any other dosing device.

adults and children 12 years and over	2 to 4 teaspoonfuls (TSP)
children 6 to under 12 years	1 to 2 teaspoonfuls (TSP)
children under 6 years	DO NOT USE

Other information

Each 5 mL (1 TSP) contains: Sodium 14 mg. Store at room temperature 20°-25°C (68°-77°F).

Inactive ingredients

citric acid, D&C red no. 33, FD&C red no. 40, black cherry flavor, methylparaben, propylene glycol, propylparaben, saccharin sodium, sodium citrate, sorbitol, water.

Questions

1-844-834-0530

Manufactured by:

Silarx Pharmaceuticals, Inc.
Carmel, NY 10512

Relabeled by:

Proficient Rx LP
Thousand Oaks, CA 91320

10-1043 Rev. 04/18



Scan Here



NDC 71205-410-04

Relabeled By: Proficient Rx LP
Thousand Oaks, CA 91320



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Diphenhydramine 12.5mg/ 5mL

118 mL (4 fl oz) Oral Solution

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

See Box. Sugar Free. Alcohol Free.

Product ID: SD041004

Mfr. By: Silarx Pharmaceuticals, Inc. Carmel, NY 10512

Store at 20°-25°C (68°-77°F)

Diphenhydramine 12.5mg/ 5mL
118 mL (4 fl oz) Oral Solution
Lot #:00000 SN# MASTER
NDC 71205-410-04 Exp:00/00/00

Diphenhydramine 12.5mg/ 5mL
118 mL (4 fl oz) Oral Solution
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NDC 71205-410-04 Exp:00/00/00

Diphenhydramine 12.5mg/ 5mL
118 mL (4 fl oz) Oral Solution
Lot #:00000 SN#MASTER
NDC 71205-410-04 Exp:00/00/00



GTIN: 00371205410049
SN# MASTER
Exp. 00/00/00
Lot #:00000

Keep medication out of the reach of children

SILADRYL ALLERGY MEDICINE

diphenhydramine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71205-410(NDC:54838-135)
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)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	CHERRY (black cherry)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71205-410-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/25/2020	11/30/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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OTC Monograph Drug	M012	01/01/1997	11/30/2024
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Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(71205-410) , RELABEL(71205-410)

Revised: 1/2024

Proficient Rx LP