DIPHENHYDRAMINE HCL STRAWBERRY FLAVOR- diphenhydramine hcl solution Innovida Pharmaeutique Corporation

Active Ingredient (in each 5mL tsp)

Diphenhydramine HCl, 12.5 mg

Purpose

Antihistamine

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 you have

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- trouble urinating due to the enlarged prostate gland

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

When using this product:

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

If pregnant of 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 (1-800-222-1222) right away.

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

Temporarily relieves these symptoms due to the common cold:

- runny nose
- sneezing

Directions

- Take every 4 to 6 hours, or as directed by a doctor.
- Do not exceed recommended dose.

Adults and Children 12 years of age take 10 to 20 mL (2 to 4 tsp) not to exceed 300 and over mg in 24 hours

Children 6 to under 12 years of age Children under 6 years of age:

take 5 to 10 mL (1 to 2 tsp) not to exceed 150 mg in 24 hours consult a doctor

Other Information

- Store at controlled room temperature 15º 30ºC (59º 86ºF).
- Do not refrigerate. Avoid excessive heat or humidity.
- Protect from light.

Inactive Ingredients

Citric acid, FD&C red #40, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, sorbitol, strawberry flavor, sucralose.

Questions or Comments? 1-888-462-4166

TAMPER EVIDENT: DO NOT USE IF SEAL UNDER CAP IS TORN, BROKEN OR MISSING.

Product label

Drug Facts

Active Ingredient (in each 5 mL tsp) Purpose Diphenhydramine HCI, 12.5 mg.. .Antihistamine

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

Temporarily relieves these symptoms due to the common cold:

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

Ask a dootor before use if you have

- a breathing problem such as emphysema or chronic bronchitis

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

When using this product-

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

If pregnant of 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 (1-800-222-1222) right away. NDC: 71800-058-19



Diphenhydramine HCI

12.5 mg/5 mL

ANTIHISTAMINE



4.02 fl oz (119 mL)

Drug Facts (continued)

- Take every 4 to 6 hours, or as directed by a doctor.
- Do not exceed recommended dose.

Adults and Children	take 10 to 20 mL (2 to 4 tsp) not
12 years of age and over:	to exceed 300 mg in 24 hours
Children 6 to under	take 5 to 10 mL (1 to 2 tsp) not
12 years of age:	to exceed 150 mg in 24 hours
Children under 6 years of age:	consult a doctor

Other Information

- Store at controlled room temperature 15° 30°C (59° 86°F).
 Do not refrigerate. Avoid excessive heat or humidity.
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Inactive Ingredients

Citric acid, FD&C red #40, glycerin, methylparaben, propylene glycol, propylparab purified water, sodium citrate, sorbitol, strawberry flavor, sucralose

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HYDROCHLORIDE

Tel.: 888-462-4166

in 5 mL

DIPHENHYDRAMINE HCL STRAWBERRY FLAVOR

diphenhydramine hcl solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71800-058

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength **DIPHENHYDRAMINE HYDROCHLORIDE** (UNII: TC2D6|AD40) **DIPHENHYDRAMINE** 12.5 mg (DIPHENHYDRAMINE - UNII:8GTS82S83M)

Inactive Ingredients Ingredient Name Strength CITRIC ACID (UNII: 2968PHW8QP) FD&C RED NO. 40 (UNII: WZB9127XOA) **GLYCERIN** (UNII: PDC6A3C0OX) METHYLPARABEN (UNII: A2I8C7HI9T) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) PROPYLPARABEN (UNII: Z8IX2SC1OH) WATER (UNII: 059QF0KO0R) SODIUM CITRATE (UNII: 1Q73Q2JULR) SORBITOL (UNII: 506T60A25R) SUCRALOSE (UNII: 96K6UQ3ZD4) STRAWBERRY (UNII: 4J2TY8Y81V)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		119 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/10/2025	
Marketing Information				
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ОТ	C Monograph Dru	g M012	06/10/2025	

Labeler - Innovida Pharmaeutique Corporation (080892908)

Revised: 6/2025 Innovida Pharmaeutique Corporation