

**DIPHENHYDRAMINE HYDROCHLORIDE- diphenhydramine
hydrochloride capsule
PD-Rx Pharmaceuticals, Inc.**

Drug Facts

Active ingredient (in each capsule)

Diphenhydramine HCl 25 mg

Purpose

Antihistamine

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

Ask a doctor before use if you have

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

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

- take every 4 to 6 hours, or as directed by a doctor

□ do not take more than 6 doses in 24 hours

adults and children 12 years of age and over	1 to 2 capsules
children 6 to under 12 years of age	1 capsule
children under 6 years of age	do not use this product in children under 6 years of age

Other information

□ store in a dry place at 15° - 30°C (59° - 86°F)

corn starch, D&C red #28, FD&C blue #1, FD&C red #40, gelatin, lactose monohydrate, magnesium stearate, sodium lauryl sulfate

Questions or comments?

1-800-616-2471

TAMPER EVIDENT: DO NOT USE IF SAFTY SEAL IS BROKEN OF MISSING FROM BOTTLE
DO NOT USE IF RED CAPSULE BAND IS BROKEN OR MISSING.

Diphenhydramin HCl 25 mg

Antihistamine

EACH CAPSULE INDIVIDUALLY
BANDED FOR YOUR PROTECTION

Drug Facts	
Active Ingredient (in each capsule)	Purpose
Diphenhydramine HCl 25 mg	Antihistamine
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	
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 • a breathing problem such as emphysema or chronic bronchitis • glaucoma • trouble urinating due to an enlarged prostate gland	
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	

GTIN: 00372789317304
SNO: D23A43000005
EXP: 12/2025
LOT: D23A43

NDC 72789-317-30



Diphenhydramine
HCL 25 mg

Marketed and Packaged By
PD-Rx Pharmaceuticals, Inc
Oklahoma City, OK 73127
1-405-942-3040 v.4.4.0

30 Capsules
TAMPER EVIDENT: DO NOT USE IF
SAFETY SEAL IS BROKEN OR MISSING FROM BOTTLE.

Drug Facts (continued)	
If pregnant or breastfeeding , 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: • take every 4 to 6 hours, or as directed by a doctor • do not take more than 6 doses in 24 hours	
Adults and children 12 years of age and over	1 to 2 capsules
children 6 to under 12 years of age	1 capsule
children under 6 years of age	do not use this product in children under 6 years of age
Other information • store in a dry place at 15° - 30°C (59° - 86°F)	
Inactive Ingredients corn starch, D&C red # 28, FD&C blue # 1, FD&C red # 40, gelatin, lactose monohydrate, magnesium stearate, sodium lauryl sulfate	
Questions or comments? 1-800-616-2471	

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DIPHENHYDRAMINE HYDROCHLORIDE

diphenhydramine hydrochloride capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72789-317(NDC:0904-7237)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	pink (Half pink and half clear with white powder inside and sealed with red band)	Score	no score
Shape	CAPSULE	Size	14mm
Flavor		Imprint Code	CPC;835
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72789-317-10	10 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2023	02/29/2028
2	NDC:72789-317-12	12 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2023	02/29/2028
3	NDC:72789-317-15	15 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2023	02/29/2028
4	NDC:72789-317-20	20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2023	02/29/2028
5	NDC:72789-317-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2023	02/29/2028

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	04/14/2022	02/29/2028

Labeler - PD-Rx Pharmaceuticals, Inc. (156893695)

Registrant - PD-Rx Pharmaceuticals, Inc. (156893695)

Establishment

Name	Address	ID/FEI	Business Operations
PD-Rx Pharmaceuticals, Inc.		156893695	repack(72789-317)

Revised: 2/2026

PD-Rx Pharmaceuticals, Inc.