

**DIPHENHYDRAMINE HCL- diphenhydramine hcl capsule
Bryant Ranch Prepack**

Drug Facts

Active ingredient (in each capsule)

Diphenhydramine HCL 25 mg

Purpose

Antihistamine

Uses:

Temporarily relieves these symptoms associated with the common cold, hay fever, or other respiratory allergies.

- sneezing
- nasal congestion
- runny nose
- itchy, watery eyes

Warnings:

Do not use

- With any other product containing Diphenhydramine HCL, including one applied topically.

Ask a doctor or pharmacist before use

If you have

- trouble urinating due to enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- if you are taking sedatives or tranquilizers

When using this product

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

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.

Directions:

- Take every 4-6 hours
- Do not take more than 6 doses in 24 hours.

Adults and children 12 years or over	1 to 2 capsule
Children 6 to under 12 years	1 capsule
Children under 6 years	ask a doctor

Other information:

- Store at room temperature 15-30 degrees C (59-86 degrees F)
- Protect from excessive moisture

Inactive ingredients: Black Iron Oxide, D & C Red #28, FD & C Blue #1, FD & C Red #40, Gelatin, Lactose Monohydrate, Magnesium Stearate, Silicon Dioxide, Sodium Lauryl Sulfate

HOW SUPPLIED

Diphenhydramine HCl 25 mg

- NDC: 71335-0352-1: 30 Capsules in a BOTTLE
- NDC: 71335-0352-2: 20 Capsules in a BOTTLE
- NDC: 71335-0352-3: 42 Capsules in a BOTTLE
- NDC: 71335-0352-4: 24 Capsules in a BOTTLE
- NDC: 71335-0352-5: 15 Capsules in a BOTTLE
- NDC: 71335-0352-6: 60 Capsules in a BOTTLE
- NDC: 71335-0352-7: 10 Capsules in a BOTTLE
- NDC: 71335-0352-8: 6 Capsules in a BOTTLE
- NDC: 71335-0352-9: 90 Capsules in a BOTTLE
- NDC: 71335-0352-0: 100 Capsules in a BOTTLE

Repackaged/Relabeled by:
Bryant Ranch Prepack, Inc.
Burbank, CA 91504

Diphenhydramine HCl Capsules 25 mg



GTIN 00371335035211
 Lot 208620
 Exp 3/28/2027
 SN 0123456789



Package Insert

Drug Facts	
Active ingredient (in each banded capsule) Diphenhydramine Hydrochloride 25 mg	Purpose Antihistamine
Uses Scan Package Insert QR Code for additional information.	
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 -a breathing problem such as emphysema or chronic bronchitis -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.	
Other Information <ul style="list-style-type: none"> •Store at room temperature, USP. •Do not use if either capsule band or imprinted safety seal under cap is broken or missing •Protect from moisture •Contains lactose 	
Directions <ul style="list-style-type: none"> •Take every 4-6 hours •Do not take more than 6 doses in 24 hours adults and children 12 years of age and over;Take 1 capsule (25 mg) children under 12 years of age;ask a doctor, the proper dosage strength is not available in this package** **Do not attempt to break capsules. The proper dosage strength and dosing information for children under 12 years of age is available on the 25 mg package.	
Inactive Ingredients D&C Red #28, FD&C Blue #1, FD&C Red #40, Gelatin, Lactose and Starch.	

NDC 71335-0352-1

diphenhydRAMINE
Hydrochloride Capsules,
USP

25 mg

30 Capsules



Repackaged by:
 Bryant Ranch Prepack, Inc.
 Burbank, CA 91504 USA

Manufactured by:
 SDA Laboratories Inc.



7133503521

DIPHENHYDRAMINE HCL

diphenhydramine hcl capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71335-0352(NDC:66424-020)
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
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
D&C RED NO. 28 (UNII: 7671P0Y5NH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Product Characteristics

Color	pink	Score	no score
Shape	CAPSULE	Size	14mm

Flavor		Imprint Code	PH014	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71335-0352-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/19/2018	
2	NDC:71335-0352-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	11/21/2018	
3	NDC:71335-0352-3	42 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	
4	NDC:71335-0352-4	24 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	
5	NDC:71335-0352-5	15 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	
6	NDC:71335-0352-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	
7	NDC:71335-0352-7	10 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	
8	NDC:71335-0352-8	6 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	
9	NDC:71335-0352-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	
10	NDC:71335-0352-0	100 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M012	01/27/2010		

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-0352) , RELABEL(71335-0352)

Revised: 3/2025

Bryant Ranch Prepack