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

- aAvoid 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 HCI Capsules 25 mg



N 00371335035211 208820 5/7/2026 0123456789

Drug Facts
Active ingredient (in each banded capsule)
Diphenhydramine Hydrochloride 50 mg.
Uses Purpose

Diphenhydramine Hydrochloide 50 mg. Antihistamine

Uses

Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: runny nose -sneezing -ktohy, watery eyes -ktohy throat and nose -Temporarily relieves these symptoms due to the common cold -runny nose -sneezing.

Warnings

Do not use -to make a child sleepy -with any other product containing diphenhydramine, even on used on skin -laks a doctor before use if you have -glaucoma -a herathing problem such as emphysema or chronic bronchita -trouble urnating due to an enlarged protestle gland. Ask a emphysema or chronic bronchita -trouble urnating due to an enlarged protestle gland. Ask as emphysema or chronic bronchita -trouble urnating due to an enlarged protestle gland. Ask as emphysema or chronic bronchita -trouble urnating due to an enlarged protestle gland. Ask as emphysema or chronic product -marked drownieses and yoccur avoid abcholic drinks -alcohol, sedatives, and tranquilizers may increase drownieses -be careful when driving a motor vehicle or operating machinery -excitability may occur, especially in driften. If pregnant or breast-feeding ask a health professional before use. Keep out of reac of children. In case of overdose, get medical help or contact a Poisson Control Center (right away.

Other Information

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

Contains actobe

Directions

-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 (50 mg)
children under 12 years of age, ask a doctor, the proper dosage strength is not available in this nackane."

package**
"The 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

BRP

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

30 Capsules

Manufactured by: Pharmaceuticals Inc.



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** 25 mg (DIPHENHYDRAMINE - UNII:8GTS82S83M)

Insetive Insurediente

HYDROCHLORIDE

Inactive Ingredients		
Ingredient Name	Strength	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)		
D&C RED NO. 28 (UNII: 767IP0Y5NH)		
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: 4/2024 Bryant Ranch Prepack