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

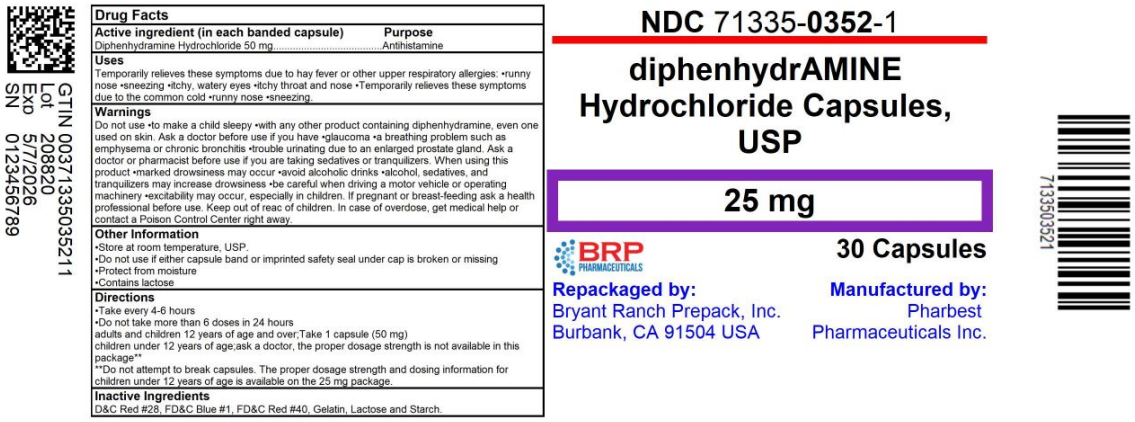
HOW SUPPLIED

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 HCl Capsules 25 mg



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**
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