BANOPHEN- diphenhydramine hcl capsule
Bryant Ranch Prepack

0836-Major(100C/1000C)

Active Ingredient (in each banded capsule)

Diphenhydramine Hydrochloride 50 mg

Purpose

Antihistamine

Use

Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies

- runny nose
- sneezing
- itchy, watery eyes
- itchy throat and nose
- Temporarily relieves these symptoms due to the common cold
  - runny nose
  - sneezing

Do not use

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

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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.

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 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.

Other Information

- Store in a dry place at 15° – 30°C (59° – 86°F).
- Do not use if either capsule band or imprinted safety seal under cap is broken or missing
- Protect from moisture
- Contains lactose

Inactive Ingredients

D&C Red #28, FD&C Blue #1, FD&C Red #40, Gelatin, Lactose and Starch.

Questions?

Questions or comments?1-800-231-4670

Distributed by: MAJOR® PHARMACEUTICALS

Indianapolis, IN 46268

(800) 616-2471

www.majorpharmaceuticals.com

HOW SUPPLIED

NDC: 71335-0081-1: 15 Capsules in a BOTTLE

NDC: 71335-0081-2: 20 Capsules in a BOTTLE

NDC: 71335-0081-3: 30 Capsules in a BOTTLE

NDC: 71335-0081-4: 10 Capsules in a BOTTLE

NDC: 71335-0081-5: 6 Capsules in a BOTTLE

NDC: 71335-0081-6: 100 Capsules in a BOTTLE

NDC: 71335-0081-7: 90 Capsules in a BOTTLE

NDC: 71335-0081-8: 60 Capsules in a BOTTLE

NDC: 71335-0081-9: 2 Capsules in a BOTTLE

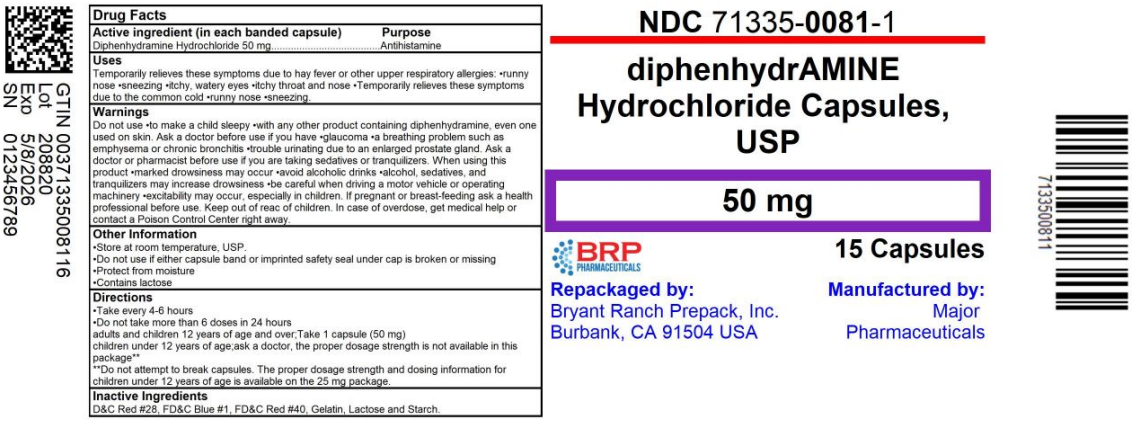
NDC: 71335-0081-0: 12 Capsules in a BOTTLE

Repackaged/Relabeled by:

Bryant Ranch Prepack, Inc.

Burbank, CA 91504

Diphenhydramine 50 mg Capsule



BANOPHEN

diphenhydramine hcl capsule

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71335-0081(NDC:0904-5307)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	50 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)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	pink	**Score**	no score
Shape	CAPSULE	**Size**	14mm
Flavor		**Imprint Code**	CPC;836
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71335-0081-1	15 in 1 BOTTLE; Type 0: Not a Combination Product	05/18/2020	
2	NDC:71335-0081-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2018	
3	NDC:71335-0081-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	08/15/2018	
4	NDC:71335-0081-4	10 in 1 BOTTLE; Type 0: Not a Combination Product	09/09/2019	
5	NDC:71335-0081-5	6 in 1 BOTTLE; Type 0: Not a Combination Product	08/07/2019	
6	NDC:71335-0081-6	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/07/2018	
7	NDC:71335-0081-7	90 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2022	
8	NDC:71335-0081-8	60 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2022	
9	NDC:71335-0081-9	2 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2022	
10	NDC:71335-0081-0	12 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	11/02/2009	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-0081) , RELABEL(71335-0081)