BANOPHEN- diphenhydramine hcl capsule Preferred Pharmaceuticals Inc.

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
 - o runny nose
 - o sneezing

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.

Directions

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

adults and children 12 years of age and	Take 1 capsule (50 mg)	
over		
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 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

Inactive Ingredients

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

Questions?

Questions or comments? (800) 616-2471

Distributed by

MAJOR® PHARMACEUTICALS 17177 N Laurel Park Drive, Suite 233, Repackaged By: Preferred Pharmaceuticals Inc.

NDC 68788-7830

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

Diphenhydramine HCL Capsules,

USP 50mg Generic for Benadryl

Each capsule contains: Diphenhydramine HCL

Pkg Size: Exp Date: ##/##/####

Mfg: Major Pharm.; Livonia, MI Prod#:

Frod#:

Warning

Store at room temperature2 0° to 25°C (68° to 77°F).

Protect from moisture. Do not use with any other
moisture. Do not use with any other
moisture of the protection of the protection of the moisture of th



May cause drowsiness. Do not drink alcohol while taking this Directions English

CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed.

> alcoholicas mientras coma esta medicina.

oma

No tome bebidas



######## EXP ##/##/##

Instrucciones Espanol:

Puede causar somnolencia. Prod# (NDC):

Qty: Ins:

Diphenhydramine HCL Capsules, USP 50mg Qty: Insurance NDC: Lot: Bat:

Diphenhydramine HCL Capsules, USP 50mg

Diphenhydramine HCL Capsules, Qty: Ins: Lot: Bat: Prod# (NDC):

Diphenhydramine HCL Capsules, USP 50mg Qty: Ins: Lot: Bat: Prod# (NDC):

Patient

BANOPHEN

diphenhydramine hcl capsule

Product Information

Product Type HUMAN OTC DRUG NDC:68788-7830(NDC:0904-5307) **Item Code (Source) Route of Administration ORAL**

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength **DIPHENHYDRAMINE HYDROCHLORIDE** (UNII: TC2D6|AD40) **DIPHENHYDRAMINE** 50 mg (DIPHENHYDRAMINE - UNII:8GTS82S83M) **HYDROCHLORIDE**

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			

I	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:68788- 7830-1	15 in 1 BOTTLE; Type 0: Not a Combination Product	01/08/2021		
	NDC:68788- 7830-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/08/2021		

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	341	01/08/2021	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment				
Name	Address	ID/FEI	Business Operations	
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-7830)	

Revised: 3/2025 Preferred Pharmaceuticals Inc.