

DIPHENHYDRAMINE HYDROCHLORIDE- diphenhydramine hydrochloride capsule
BANOPHEN- diphenhydramine hcl capsule
Major Pharmaceuticals

0835&0836(box unit)-Major

Active Ingredient (in each banded capsule)

Diphenhydramine HCl... 25 mg

Diphenhydramine HCl... 50 mg

Purpose

Antihistamine

Use

25 MG

- 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

50 MG

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

WARNINGS

Do not use

25 MG

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

50 MG

- to make a child sleepy
- with any other product containing diphenhydramine, including one applied topically

Ask a doctor before use if you have

25 MG

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- trouble urinating due to an enlarged prostate gland

50 MG

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the 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

25 MG

adults and children 12 years of age and over	1 to 2 capsules
children 6 years to under 12 years of age	1 capsule
children under 6 years of age	do not use this product in children under 6

|years of age

50 MG

adults and children 12 years of age and over	1 capsule
children 6 years to 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 20°C - 25°C (68°F - 77°F); excursions permitted to 15° - 30°C (59° - 86°F) [See USP Controlled Room Temperature]
- 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,

Livonia, MI 48152

Major-24BB

DO NOT USE IF CARTON IS OPENED OR IF A BLISTER UNIT IS TORN, BROKEN, OR SHOWS ANY SIGNS OF TAMPERING.	
Drug Facts	
Active ingredient (in each banded capsule) Diphenhydramine Hydrochloride 25 mg	Purpose Antihistamine
Uses ■ temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: ■ runny nose ■ sneezing ■ itchy, watery eyes ■ itching of the nose or throat ■ 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 one used on skin	
Ask a doctor before use if you have ■ a breathing problem such as emphysema or chronic bronchitis ■ glaucoma ■ trouble urinating due to an enlarged prostate gland	
Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers	

BAN
Compare to the active ingredient of BENADRI

OPHEN™
24 CAPSULES

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 to 6 hours ■ do not take more than 6 doses in 24 hours

adults and children 12 years of age and over	1 to 2 capsules
children 6 to under 12 years of age	1 capsule
children under 6 years of age	do not use this product in children under 6 years of age

Other information ■ store at room temperature, USP ■ protect from moisture ■ contains lactose

Inactive ingredients D&C red #28, FD&C blue #1, FD&C red #40, gelatin, lactose and starch

Questions or comments? (800) 616-2471

*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., owner of the registered trademark Benadryl®

Made in the U.S.A. 50844 M-29 Rev. 10/16 4568-10-16
Distributed by: **MAJOR® PHARMACEUTICALS**
17177 N Laurel Park Drive, Suite 233, Livonia, MI 48152



BANOPHEN™
Compare to the active ingredient of BENADRYL®
24 CAPSULES

ANTIHISTAMINE
For the temporary relief from symptoms of:
• Upper Respiratory Allergies • Hay Fever



Compare to the active ingredient of BENADRYL®*
Each Capsule Individually Banded For Your Protection
24 CAPSULES

BANOPHEN™
Complete Allergy Medication
Diphenhydramine HCl 25 mg

MAJOR®
NDC 0904-2035-24

BANOPHEN™
Compare to the active ingredient of BENADRYL®
24 CAPSULES

BANOPHEN™
Compare to the active ingredient of BENADRYL®*
24 CAPSULES



NDC 0904-5306-80

BANOPHEN™

Complete Allergy Medication
Diphenhydramine HCl 25 mg

ANTIHISTAMINE

For the temporary relief of the symptoms of:
• Upper Respiratory Allergies • Hay Fever



Compare to the active ingredient of **BENADRYL®***

1000 CAPSULES

Each Capsule Individually Banded For Your Protection

THIS LARGE PACKAGE SIZE FOR DISPENSING PURPOSE ONLY

Drug Facts

Active ingredient (in each banded capsule)
Diphenhydramine HCl 25 mg

Purpose
Antihistamine

Uses

Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: ■ runny nose ■ sneezing ■ itchy, watery eyes ■ itching of the nose or throat ■ 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 one used on skin

Ask a doctor before use if you have ■ a breathing problem such as emphysema or chronic bronchitis ■ glaucoma ■ 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 to 6 hours ■ do not take more than 6 doses in 24 hours
adults and children 12 years of age and over 1 to 2 capsules
children 6 to under 12 years of age 1 capsule

do not use this product in children under 6 years of age
children under 6 years of age

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 or comments? (800) 616-2471

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17171 Union Road, Suite 220,
Livonia, MI 48152

Re-Order No. 00829
M-29 Rev. 10/16
4570-10-16
Lot # & Exp. Date:
3 09045 30680 2



R51834

100 CAPSULES
ANTIHISTAMINE
50 mg
DIPHENHYDRAMINE HCl
Unit Dose
MAJOR®
NDC 0904-2056-61



NDC 0904-2056-61
MAJOR®
Unit Dose
DIPHENHYDRAMINE HCl
50 mg
ANTIHISTAMINE
100 CAPSULES

Unit Dose
DIPHENHYDRAMINE HCl
ANTIHISTAMINE
DO NOT USE IF CARTON IS OPENED OR IF A BLISTER UNIT IS TORN, BROKEN, OR SHOWS ANY SIGNS OF TAMPERING. DO NOT USE IF RED CAPSULE BAND IS BROKEN OR MISSING.

Drug Facts

Active ingredient (in each capsule)	Purpose
Diphenhydramine Hydrochloride 50 mg	Antihistamine

Uses
Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: ■ sneezing ■ runny nose ■ itchy, watery eyes ■ itchy throat and nose

Warnings
Do not use ■ to make a child sleepy ■ with any other products 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 ■ difficulty in urination due to enlargement of the 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 ■ use caution when driving a motor vehicle or operating machinery ■ excitability may occur, especially in children

NDC 0904-2056-61
MAJOR®
Unit Dose
DIPHENHYDRAMINE HCl
50 mg
ANTIHISTAMINE
100 CAPSULES

Drug Facts (continued)

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 (1-800-222-1222) right away.

Directions
Take every 4 to 6 hours ■ do not take more than 6 doses in 24 hours
adults and children 12 years of age and over 1 capsule
child or 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) ■ protect from moisture ■ contains lactose

Inactive ingredients D&C red #28, FD&C blue #1, FD&C red #40, gelatin, lactose and starch

Questions or comments? 1-800-231-4670

Keep this carton for complete product information
Distributed by: MAJOR® PHARMACEUTICALS
Indianapolis, IN 46268
(800) 616-2471
www.majorpharmaceuticals.com

M-29 Rev. 03/23 Re-Order No. 301575



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DIPHENHYDRAMINE HYDROCHLORIDE

diphenhydramine hydrochloride capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-5306
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
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	pink (half pink and half clear with white powder inside)	Score	no score
Shape	CAPSULE	Size	14mm
Flavor		Imprint Code	CPC;835
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-5306-60	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/02/2009	07/25/2025
2	NDC:0904-5306-80	1000 in 1 BOTTLE; Type 0: Not a Combination Product	01/02/2009	06/30/2025
3	NDC:0904-5306-61	10 in 1 BOX	01/02/2009	07/31/2025
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:0904-5306-24	2 in 1 CARTON	03/15/2019	04/30/2025
4		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/02/2009	07/31/2025

DIPHENHYDRAMINE HYDROCHLORIDE

diphenhydramine hydrochloride capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-2056
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 (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:0904-2056-61	10 in 1 BOX	01/02/2009	12/31/2027
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/02/2009	12/31/2027

BANOPHEN

diphenhydramine hcl capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-2035
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
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	pink (half pink and half clear with white powder inside)	Score	no score
Shape	CAPSULE	Size	14mm
Flavor		Imprint Code	CPC;835
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-2035-24	2 in 1 CARTON	01/02/2009	08/31/2021
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/02/2009	08/31/2021

Labeler - Major Pharmaceuticals (191427277)

