DIPHENHYDRAMINE HCL- diphenhydramine hcl capsule
Akron Pharma Inc.

Akron Pharma, Inc.

Active ingredient (in each banded capsule)

Diphenhydramine Hydrochloride 25 mg and 50 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

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

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- marked drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- excitability may occur, especially in children
- 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 (1-800-222-1222) right away.

Directions

- take every 4 to 6 hours
- do not take more than 6 doses in 24 hours

25 mg Dosage:

Age	**Dose**
adults and children 12 years 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

50 mg Dosage:

Age	**Dose**
adults and children 12 years and over	1 capsules
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 temperature 15° to 30° C (59° to 86°F)
- do not use if either capsule band or imprinted safety seal under cap is broken or missing
- protect from moisture

Inactive ingredients

Colloidal silicon dioxide, Gelatin, Lactose Monohydrate, Magnesium Stearate, Microcrystalline cellulose, Sodium lauryl sulfate

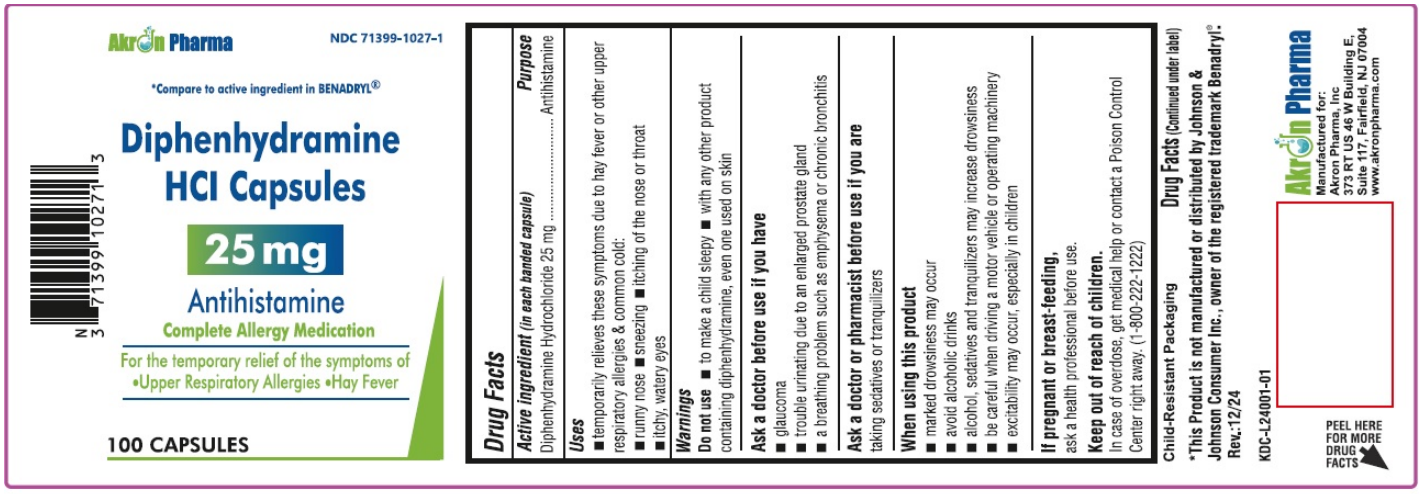
Questions or comments?

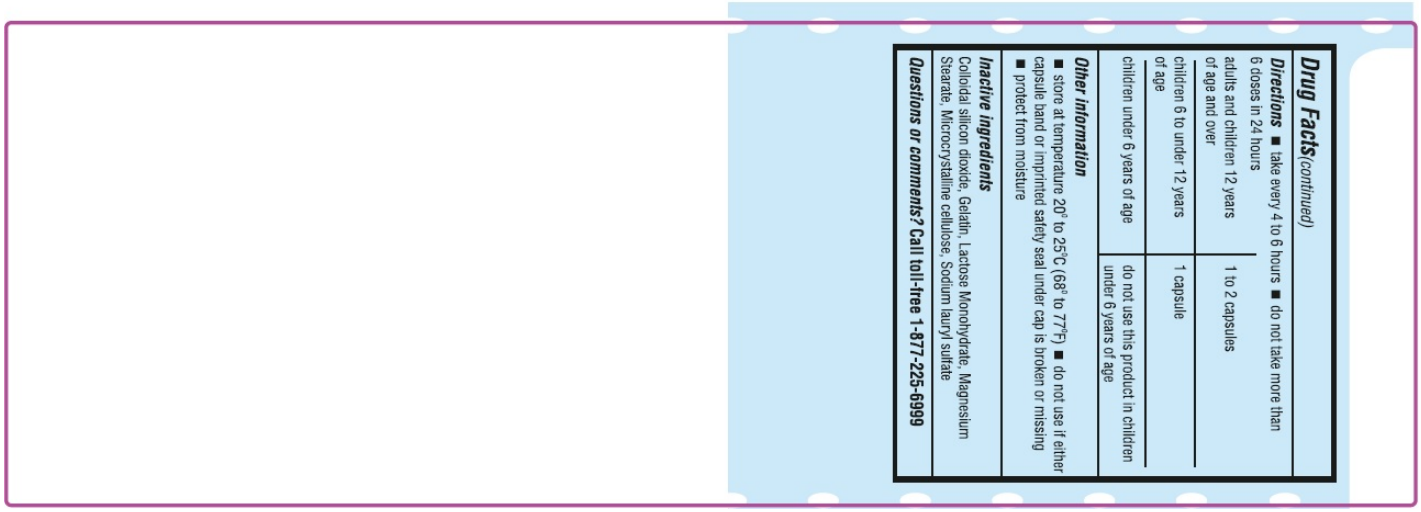
Call toll-free 1-877-225-6999

Manufactured for:
Akron Pharma, Inc,

373 RT US 46 W, Building E,

Suite 117, Fairfield, NJ 07004





Akron Pharma

NDC 71399-1027-2

THIS LARGE PACKAGE SIZE FOR DISPENSING PURPOSE ONLY

*Compare to active ingredient in BENADRYL®

Diphenhydramine HCl Capsules

25 mg

Antihistamine

Complete Allergy Medication

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

1000 CAPSULES

Drug Facts

Active ingredient (in each banded capsule) — **Purpose**
Diphenhydramine Hydrochloride 25 mg ..... Antihistamine

Uses
- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies & common cold:
- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes

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

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. (1-800-222-1222)

Drug Facts (continued)

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 temperature 20° to 25°C (68° to 77°F)
- do not use if either capsule band or imprinted safety seal under cap is broken or missing
- protect from moisture

Inactive ingredients
Colloidal silicon dioxide, Gelatin, Lactose Monohydrate, Magnesium Stearate, Microcrystalline cellulose, Sodium lauryl sulfate

Questions or comments? Call toll-free 1-877-225-6999

KDC-L24003-01

Rev.:12/24

Child-Resistant Packaging

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

Akron Pharma
Manufactured for:
Akron Pharma, Inc
373 RT US 46 W Building E,
Suite 117, Fairfield, NJ 07004
www.akronpharma.com

Akron Pharma

NDC 71399-1028-1

*Compare to active ingredient in BENADRYL®

Diphenhydramine HCl Capsules

50 mg

Antihistamine

Complete Allergy Medication

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

100 CAPSULES

Drug Facts

Active ingredient (in each banded capsule) — **Purpose**
Diphenhydramine Hydrochloride 50 mg ..... Antihistamine

Uses
- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies & common cold:
- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes

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

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. (1-800-222-1222)

Drug Facts (Continued under label)

Child-Resistant Packaging

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

Rev.:12/24

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PEEL HERE FOR MORE DRUG FACTS

Drug Facts (continued)

Directions
- take every 4 to 6 hours
- do not take more than 6 doses in 24 hours

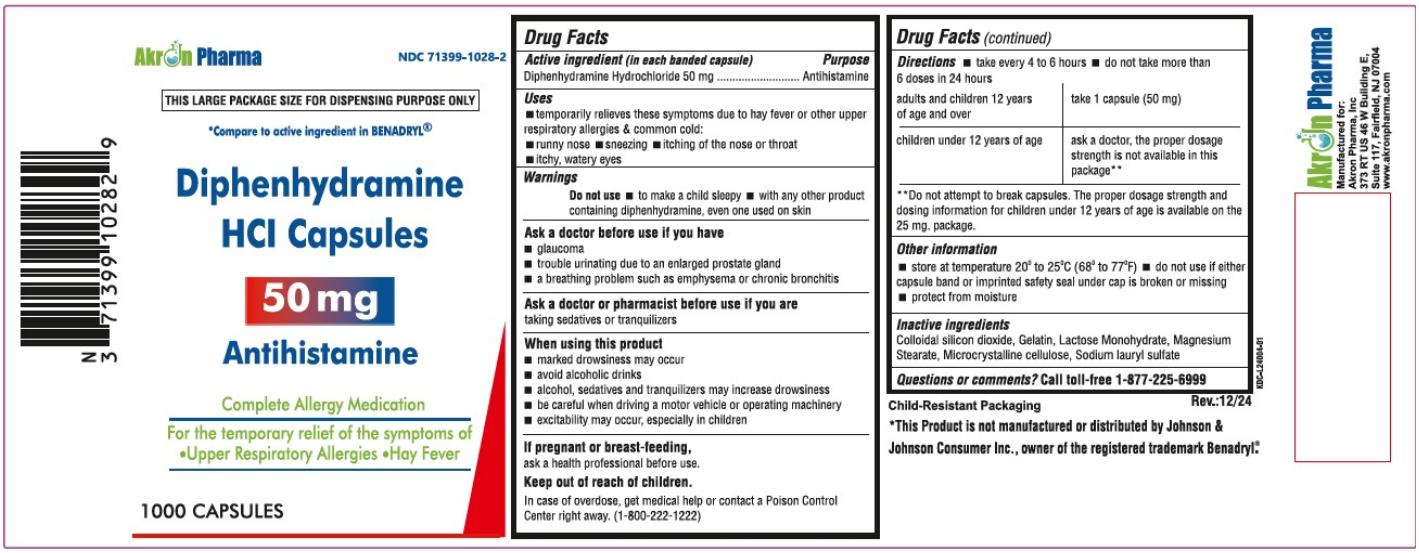
adults and children 12 years of age and over	take 1 capsule (50 mg)
children under 12 years of age	ask a doctor, the proper dosage strength is not available in this package**

**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 at temperature 20° to 25°C (68° to 77°F)
- do not use if either capsule band or imprinted safety seal under cap is broken or missing
- protect from moisture

Inactive ingredients
Colloidal silicon dioxide, Gelatin, Lactose Monohydrate, Magnesium Stearate, Microcrystalline cellulose, Sodium lauryl sulfate

Questions or comments? Call toll-free 1-877-225-6999



DIPHENHYDRAMINE HCL

diphenhydramine hcl capsule

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71399-1027
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Product Characteristics

Color	pink (PINK WHITE)	**Score**	no score
Shape	capsule (oblong)	**Size**	15mm
Flavor		**Imprint Code**	AD25
Contains			

Packaging

			Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-1027-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2024	
2	NDC:71399-1027-2	1000 in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/06/2024	

DIPHENHYDRAMINE HCL

diphenhydramine hcl capsule

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71399-1028
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Product Characteristics

Color	pink (PINK WHITE)	**Score**	no score
Shape	capsule (oblong)	**Size**	15mm
Flavor		**Imprint Code**	AD50
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:71399-1028-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2024	
2	NDC:71399-1028-2	1000 in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/06/2024	

Labeler - Akron Pharma Inc. (067878881)

Revised: 12/2024 Akron Pharma Inc.