

**DIPHENHYDRAMINE HYDROCHLORIDE- diphenhydramine
hydrochloride capsule
ST. MARY'S MEDICAL PARK PHARMACY**

0835K- Major

Drug Facts

Active ingredient (in each 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
- 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 (1-800-222-1222).

Directions

- take every 4 to 6 hours, or as directed by a doctor
- 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 in a dry place at 15° - 30°C (59° - 86°F)

corn starch, D&C red #28, FD&C blue #1, FD&C red #40, gelatin, lactose monohydrate, magnesium stearate, sodium lauryl sulfate

Questions or comments?

1-800-616-2471

Distributed by: MAJOR® PHARMACEUTICALS, Indianapolis, IN 46268

Product of China. Manufactured and packaged in the USA using domestic and imported ingredients.

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

To preserve quality and freshness, keep bottle tightly closed.

KEEP OUT OF REACH OF CHILDREN. DO NOT USE IF PRODUCT APPEARS TO BE TAMPERED WITH OR IMPRINTED SEAL UNDER CAP IS BROKEN OR MISSING. DO NOT USE IF RED CAPSULE BAND IS BROKEN OR MISSING.

<p>NDC 60760-697-30 Diphenhydramine HCl (BANOPHEN) CAPSULES 25 mg QTY: 30 LOT# XXXXXXX EXP XX-XX RX# ??????? MANUFACTURED BY: MAJOR PHARMACEUTICALS Indianapolis, IN 46268</p>	<p>Diphenhydramine HCl (BANOPHEN) CAPSULES 25 mg QTY: 30 RX# ??????? NDC 60760-697-30 LOT# XXXXXXX EXP XX-XX</p>	<p>Diphenhydramine HCl (BANOPHEN) CAPSULES 25 mg QTY: 30 RX# ??????? NDC 60760-697-30 LOT# XXXXXXX EXP XX-XX</p>	<p>Diphenhydramine HCl (BANOPHEN) CAPSULES 25 mg QTY: 30 RX# ??????? NDC 60760-697-30 LOT# XXXXXXX EXP XX-XX</p>	<p>Diphenhydramine HCl (BANOPHEN) CAPSULES 25 mg QTY: 30 RX# ??????? NDC 60760-697-30 LOT# XXXXXXX EXP XX-XX</p>
<p>PACKAGED BY:  St. Mary's 10860 MAVINEE DR. ORO VALLEY, AZ 85737 MANAGED PHARMACY PROGRAMS</p>		<p>TAKE AS DIRECTED MAY CAUSE DROWSINESS</p>  <p>Rx only STORE AT CONTROLLED ROOM TEMPERATURE 15°- 30° C (59°-86°F)</p>		

DIPHENHYDRAMINE HYDROCHLORIDE
 diphenhydramine hydrochloride capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60760-697(NDC:0904-7237)
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)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	pink (Half pink and half clear with white powder inside and sealed with red band)	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:60760-697-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/19/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	04/14/2022	

Labeler - ST. MARY'S MEDICAL PARK PHARMACY (063050751)

Establishment

Name	Address	ID/FEI	Business Operations
ST. MARY'S MEDICAL PARK PHARMACY		063050751	relabel(60760-697) , repack(60760-697)

Revised: 12/2023

ST. MARY'S MEDICAL PARK PHARMACY