ALLERGY- diphenhydramine hydrochloride capsule
Mckesson

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DIPHENHYDRAMINE HCl 25mg, USP

Active Ingredient

(per capsule)

Diphenhydramine HCl 25 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms of hay fever or other upper respiratory allergies:

- runny nose
- itching of the nose or throat
- sneezing
- itchy, watery eyes

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

- you may get very drowsy
- avoid alcoholic drinks
- alcohol, sedatives & 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
- do not take more than 6 tablets in 24 hours

Age (Yr)	Dose (capsules)
Adults and children 12 years of age and over	Take 1 to 2 capsules
Children 6 to under 12 years of age	Take 1 capsule
Children under 6 years of age	do not use

Other Information

- store at room temperature between 15°-30°C (59°-86°F)
- protect from light and moisture

Inactive Ingredients

Benzyl alcohol, Butyl paraben, D&C red # 28, FD&C blue # 1, FD&C red # 40, Edible black ink, gelatin, lactose, Magnesium stearate, Methyl paraben, polysorbate 80, Propyl paraben, Purified water, Sodium lauryl sulfate, Starch

Questions or Comments

Call 1-855-314-1850

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL


sunmark®
NDC 70677-0010-1
allergy relief
Antihistamine
For the temporary relief of sneezing, itchy & watery eyes, runny nose & itchy throat
DIPHENHYDRAMINE HCl 25 mg
100 CAPSULES
Drug Facts
Active ingredient (per capsule) Purpose
Diphenhydramine HCl 25 mg Antihistamine
Uses temporarily relieves these symptoms of hay fever or other upper respiratory allergies:
• runny nose • itchy nose or throat
• sneezing • itchy, watery eyes
Warnings
• Do not use with any other product containing diphenhydramine, even one used on skin
• to make a child sleepy.
Ask a doctor before use if you have
• glaucoma • trouble urinating due to 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 & 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
• do not take more than 6 capsules in 24 hours
• Adults and children 12 years and over: take 1 to 2 capsules every 4 to 6 hours • Children 6 to under 12 years of age: take 1 capsule • Children under 6 years of age: Do not use.
READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION
Made in India
Another Quality Product Distributed By McKesson
One Post Street, San Francisco, CA 94104
Money Back Guarantee
Please visit us at www.sunmarkbrand.com
McKESSON
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NDC 70677-0010-1

ALLERGY

diphenhydramine hydrochloride capsule

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70677-0010
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**
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
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 (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics			
Color	WHITE, PINK	**Score**	no score
Shape	CAPSULE	**Size**	14mm
Flavor		**Imprint Code**	AP;020
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70677-0010-1	100 in 1 BOTTLE	11/25/2016	
1		1 in 1 CARTON; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	11/25/2016	

Labeler - Mckesson (177667227)

Revised: 12/2016 Mckesson