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

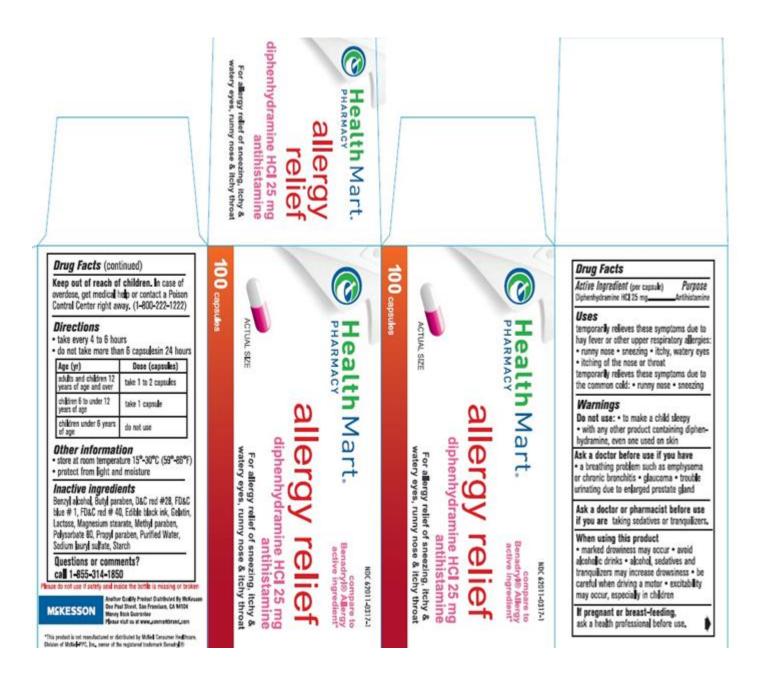
Benzyl alcohol, Butyl paraben, D&C red # 28, FD&C blue # 1, FD&C red # 40, Edible black ink, gelatin, lactose, Magnesium stearate, Methyl paraben, polysorbate 80, Propyl paraben, Purified water, Sodium lauryl sulfate, Starch

Questions or Comments

Call 1-855-314-1850

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL





NDC 62011-0317-1

ALLERGY diphenhydramine hydrochlori	ide capsule					
	-					
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (So	Item Code (Source)		NDC:62011-0317	
Route of Administration	ORAL					
Active Ingredient/Active	Moiety					
Ingredient Name		Basis of Strength		Strengt		
	DIPHENHYDRAMINE HYDRO CHLO RIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)			DIPHENHYDRAMINE HYDROCHLORIDE		
DIPHENHYDRAMINE HYDROC	HLORIDE (UNII: TC2D6JAD40) (E				25 mg	

Inactive Ingredien	165				
	Ingredient Name			Strength	
BENZYL ALCOHOL (U	NII: LKG8494WBH)				
BUTYLPARABEN (UNII	: 3QPI1U3FV8)				
D&C RED NO.28 (UNII	: 767IP0 Y5NH)				
FD&C BLUE NO. 1 (UN	II: H3R47K3TBD)				
FD&C RED NO.40 (UN	II: WZB9127XOA)				
GELATIN (UNII: 2G86Q	N327L)				
LACTOSE (UNII: J2B2A	4N98G)				
MAGNESIUM STEARA	ГЕ (UNII: 70097M6I30)				
METHYLPARABEN (UN	₩II: А2I8С7HI9Т)				
PROPYLPARABEN (UN					
POLYSORBATE 80 (UI					
SODIUM LAURYL SUL	FATE (UNII: 368GB5141J)				
STARCH, CORN (UNII:					
Product Character Color	r istics WHITE, PINK	Score		no score	
Shape	CAPSULE	Size		14mm	
Flavor		Imprint Cod	da	AP;020	
Contains			ue	AI,020	
Contains					
Packaging					
	Package Description				
	Package Desc	rintion	Marketing Start Date	Marketing End Date	
# Item Code	Package Desc	ription	Marketing Start Date	Marketing End Date	
Item Code 1 NDC:62011-0317-1 1	100 in 1 BOTTLE	-	Marketing Start Date 11/25/2016	Marketing End Date	
# Item Code 1 NDC:62011-0317-1 1	C C	-		Marketing End Dat	
Item Code 1 NDC:62011-0317-1 1 1 1 1	l00 in 1 BOTTLE l in 1 CARTON; Type 0: Not a C	-		Marketing End Date	
# Item Code 1 NDC:62011-0317-1 1 1 1 1	100 in 1 BOTTLE L in 1 CARTON; Type 0: Not a C rmation	Combination Product	11/25/2016	Marketing End Date	
Item Code 1 NDC:62011-0317-1 1 1 1 1	100 in 1 BOTTLE L in 1 CARTON; Type 0: Not a C rmation	Combination Product		Marketing End Date	

Labeler - Mckesson (177667227)

Revised: 12/2016

Mckesson