

DIPHENHYDRAMINE HYDROCHLORIDE - diphenhydramine hydrochloride capsule
Unit Dose Services

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 HYDROCHLORIDE CAPSULES, USP 25mg

Drug Facts

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Active Ingredient
(in each capsule)

Diphenhydramine HCl 25 mg

Purpose

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

with any other product containing diphenhydramine, even one used on skin **Do not use**

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
- taking sedatives or tranquilizers **Ask a doctor or pharmacist before use if you are**

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
- ask a health professional before use. **If pregnant or breast-feeding,**

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- take 1 to 2 capsules every 4-6 hours; not more than 6 doses in 24 hours **adults and children 12 years and over:**
- ask a doctor **children under 12 years:**

Other Information

- store at 15-30 °C (59-86 °F)
- protect from moisture
- For 1000 Count: This is a bulk package. Dispense contents in a tight, light-resistant container with a child-resistant closure as defined in the USP

Inactive Ingredients

benzyl alcohol, butylparaben, D&C red# 28, edible black ink, FD&C blue #1, FD&C red# 40, gelatin, lactose, magnesium stearate, methylparaben, polysorbate 80, propylparaben, sodium laurel sulfate

Questions or Comments

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP OR BAND AROUND ANY CAPSULE IS MISSING OR DAMAGED

Distributed by: Qualitest Pharmaceuticals, Inc.

DIPHENHYDRAMINE HYDROCHLORIDE CAPSULE

NDC: 50436-3594-1

**DIPHENHYDRAMINE
HCL
25 MG
30 CAP**

Rx

WARNING:
KEEP OUT OF REACH OF CHILDREN
STORE AT 20-25°C (68-77°F)
CONTROLLED ROOM TEMPERATURE



MFG BY: QUALITEST
XXXXXXXXXX
MFG NDC: 00603-3339-32
MFG LOT: XXXXXXXX
LOT: XXXXXXXX EXP: XXXXXXXX
Pkg by: Unit Dose Services, LLC
Miami, FL 33179

NDC: 50436-3594-1 30 CAP
DRUG: DIPHENHYDRAMINE
HCL
LOT: XXXXXXXX EXP: XXXXXXXX

NDC: 50436-3594-1 30 CAP
DRUG: DIPHENHYDRAMINE
HCL
LOT: XXXXXXXX EXP: XXXXXXXX

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DRUG: DIPHENHYDRAMINE
HCL
LOT: XXXXXXXX EXP: XXXXXXXX

DIPHENHYDRAMINE HYDROCHLORIDE

diphenhydramine hydrochloride capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50436-3594(NDC:0603-3339)
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: 3QP1IU3FV8)	
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Product Characteristics

Color	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:50436-3594-1	30 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	05/24/2007	

Labeler - Unit Dose Services (831995316)

Registrant - Unit Dose Services (831995316)

Establishment

Name	Address	ID/FEI	Business Operations
Unit Dose Services		831995316	REPACK(50436-3594)