# ALLERGY RELIEF- diphenhydramine hcl capsule, liquid filled Topco Associates, LLC

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.

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## **TopCare 44-658**

### Active ingredient (in each liquid-filled capsule)

Diphenhydramine HCl 25 mg

#### Purpose

Antihistamine

#### Uses

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

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- 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 (1-800-222-1222) right away.

#### 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	1 to 2 capsules
over	
children 6 to under 12 years of age	1 capsule
	do not use this
children under 6	product in
years	children
of age	under 6 years of
	age

#### Other information

- TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- protect from heat, humidity and light
- see end flap for expiration date and lot number

## **Inactive ingredients**

edible white ink, gelatin, glycerin, lecithin, mineral oil, polyethylene glycol, purified water, sorbitol

#### Questions or comments?

1-888-423-0139

## Principal display panel

## **TopCare**

Healthтм

NDC 36800-658-08

COMPARE TO BENADRYL®
DYE-FREE ALLERGY LIQUI-GELS®
ACTIVE INGREDIENT\*

**DYE-FREE** 

## **Allergy Relief**

DIPHENHYDRAMINE HCl 25 mg - Antihistamine

#### **RELIEF OF:**

- Sneezing
- Runny nose
- Itchy, Watery Eyes
- Itchy, Nose or Throat

Our Pharmacists Recommend

#### 24 SOFTGELS\*\*

\*\* Liquid Filled Capsules

actual size

DISTRIBUTED BY TOPCO ASSOCIATES LLC ELK GROVE VILLAGE, IL 60007 ©TOPCO LNKA0219 QUESTIONS? 1-888-423-0139 topcare@topco.com www.topcarebrand.com

PRODUCT OF CHINA PACKAGED AND QUALITY ASSURED IN THE USA

\*This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Benadryl® Dye-Free Allergy LIQUI-GELS®.

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TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING





**Topcare 44-658** 

#### ALLERGY RELIEF

diphenhydramine hcl capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:36800-658
Route of Administration	ORAL		

	Active Ingredient/Active Moiety		
l	Ingredient Name	Basis of Strength	Strength
	$ \begin{tabular}{ll} \textbf{DIPHENHYDRAMINE HYDRO CHLO RIDE} & (UNII: TC2D6 JAD40) & (DIPHENHYDRAMINE -UNII: 8GTS82S83M) \\ \end{tabular} $	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients			
Ingredient Name	Strength		
GELATIN (UNII: 2G86QN327L)			
GLYCERIN (UNII: PDC6A3C0OX)			
LECITHIN, SO YBEAN (UNII: 1DI56 QDM62)			
MINERAL OIL (UNII: T5L8T28FGP)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
WATER (UNII: 059QF0KO0R)			
SORBITOL (UNII: 506T60A25R)			

Product Characteristics			
Color	YELLOW (clear)	Score	no score
Shape	OVAL	Size	15mm
Flavor		Imprint Code	658
Contains			

Packaging						
# Item Co	o de	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>		
1 NDC:36800	2 in 1 CARTON		06/15/2019			
1	12 in 1 BLISTE Product	R PACK; Type 0: Not a Combination				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH FINAL	part341	06/15/2019		

## Labeler - Topco Associates, LLC (006935977)

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		832867837	PACK(36800-658)	

Revised: 3/2020 Topco Associates, LLC