

**ALLERGY RELIEF DYE-FREE- diphenhydramine hcl capsule, liquid filled**  
**Cardinal Health**

*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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**Leader 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
- difficulty in urination due to enlargement of the 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 beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- excitability may occur, especially in children
- use caution when driving a motor vehicle or operating machinery

**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 and over	1 to 2 capsules
children 6 to under 12 years	1 capsule
children under 6 years	do not use

**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, polyethylene glycol, purified water, sorbitol

**Questions or comments?**

**1-800-426-9391**

**Principal display panel**

**LEADER™**

NDC 70000-0144-1

Dye-Free

**Allergy Relief**

Diphenhydramine HCl, 25 mg  
Antihistamine

**For Allergy Relief**

Sneezing, Runny Nose,  
Itchy, Watery Eyes,  
Itchy Throat

**24 LIQUID GELS**

Liquid Filled Capsules

**COMPARE TO  
BENADRYL® DYE-FREE**

# ALLERGY LIQUI-GELS®

active ingredient\*

100% Money

Back Guarantee

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

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

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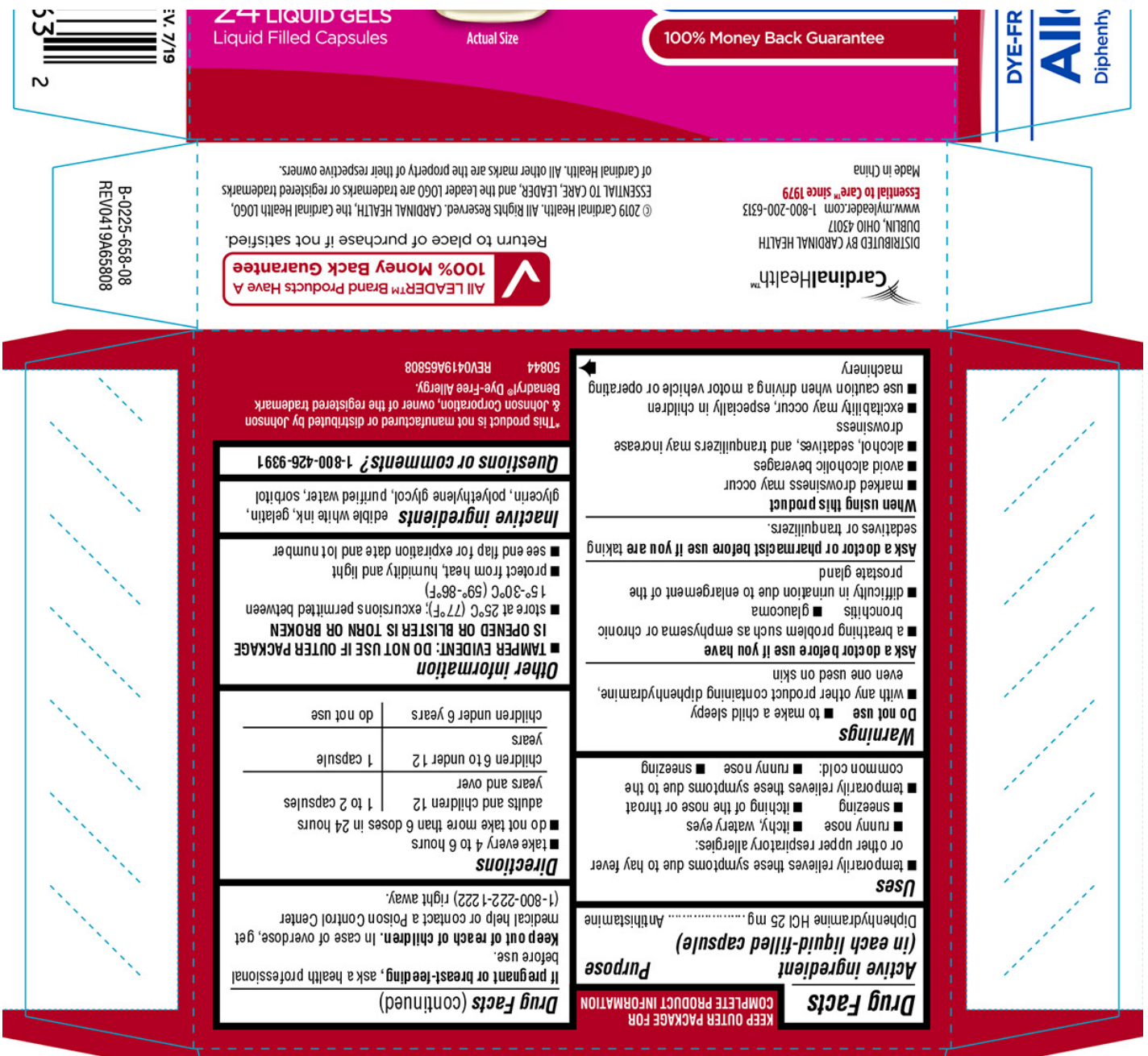
Made in China

All LEADER™ Brand Products Have A

**100% Money Back Guarantee**

Return to place of purchase if not satisfied.





Leader 44-658

ALLERGY RELIEF DYE-FREE			
diphenhydramine hcl capsule, liquid filled			
<b>Product Information</b>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0144
Route of Administration	ORAL		
<b>Active Ingredient/Active Moiety</b>			
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
	DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

## Inactive Ingredients

Ingredient Name	Strength
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
SORBITOL (UNII: 506T60A25R)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
WATER (UNII: 059QF0K00R)	

## Product Characteristics

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

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0144-1	2 in 1 CARTON	03/01/2015	
1		12 in 1 BLISTER PACK; 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	03/01/2015	

**Labeler** - Cardinal Health (097537435)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	PACK(70000-0144)

Revised: 10/2019

Cardinal Health