

**ALLERGY RELIEF ANTIHISTAMINE- diphenhydramine hydrochloride capsule,  
liquid filled  
CDMA Inc**

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**Allergy Relief Antihistamine**

***Drug Facts***

***Active ingredient (in each softgel)***

Diphenhydramine HCl 25mg

***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 • trouble urinating due to an 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, 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**

**Overdose warning:** In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as children even if you do not notice any signs or symptoms.

## **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 over	1 to 2 softgel
children 6 to under 12 years	1 softgel
children under 6 years of age	do not use this product

## **Other information**

- store at room temperature between 20-25°C (68-77°F) • avoid excessive heat, humidity and light

## **Inactive ingredients**

edible white ink, gelatin, glycerin, polyethylene glycol, povidone, purified water, sorbitol sorbitan solution

## **Questions or Comments?**

Call: **1-888-577-8033** Monday - Friday 9am-4:30pm EST

## **QC<sup>®</sup> QUALITY CHOICE**

**\*Compare to the active Ingredient in BENADRYL<sup>®</sup> Allergy LIQUI-GELS<sup>®</sup>**

### **Dye-Free**

- Relieves sneezing
- Itchy, watery eyes
- Runny nose
- Itchy throat

**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 McNeil Consumer Healthcare, distributor of Benadryl<sup>®</sup>, or Catalent Pharma Solutions, Inc., owner of the registered trademark LIQUI-GELS<sup>®</sup>.

## **QC 100% SATISFACTION GUARANTEED**

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Questions: 800-935-2362

Product of UAE

Packaged and Quality assured in USA

**READ AND KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION**

REV.01-092022

**Packaging**



<b>ALLERGY RELIEF ANTIHISTAMINE</b>			
diphenhydramine hydrochloride capsule, liquid filled			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63868-474
<b>Route of Administration</b>	ORAL		
<b>Active Ingredient/Active Moiety</b>			
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
	<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>GELATIN, UNSPECIFIED</b> (UNII: 2G86QN327L)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ05DW1A)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SORBITAN</b> (UNII: 6O92ICV9RU)	

## Product Characteristics

<b>Color</b>	WHITE (clear)	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	15mm
<b>Flavor</b>		<b>Imprint Code</b>	783
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-474-24	2 in 1 CARTON	05/01/2021	
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 Drug	M012	05/01/2021	

**Labeler** - CDMA Inc (011920774)

Revised: 12/2024

CDMA Inc