

**ALLERGY- diphenhydramine hydrochloride capsule**  
**Publix Super Markets Inc**

*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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**Publix Super Markets, Inc. Allergy Capsules Drug Facts**

**Active ingredient (in each capsule)**

Diphenhydramine HCl 25 mg

**Purpose**

Antihistamine

**Uses**

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

- 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**

- 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 right away (1-800-222-1222).

**Directions**

- take every 4 to 6 hours, or as directed by a doctor
- 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          |

**Other information**

- store at 20-25°C (68-77°F). Avoid high humidity. Protect from light.
- **each capsule is sealed with a Uni Band® seal which bonds the two capsule halves together. Do not use if seal is broken or missing. Do not use if blister unit is broken or torn.**

**Inactive ingredients**

anhydrous lactose, benzyl alcohol, butylparaben, D&C red no. 28, edetate calcium disodium, edible ink, FD&C blue no. 1, FD&C red no. 40, gelatin, magnesium stearate, methylparaben, polysorbate 80, propylparaben, sodium lauryl sulfate, sodium propionate

**Principal Display Panel**

allergy capsules

DIPHENHYDRAMINE HCl 25 mg

ANTIHISTAMINE

Relieves:

Sneezing

Itchy, watery eyes

Runny nose

Itchy throat

ACTUAL SIZE

48 CAPSULES

Compare to the active ingredient in Benadryl®



# allergycapsules

DIPHENHYDRAMINE HCl 25 mg  
ANTIHISTAMINE



NDC 56062-462-67

# allergycapsules

DIPHENHYDRAMINE HCl 25 mg  
ANTIHISTAMINE

### Relieves:

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Important: Read all product information before using. Keep this box for important information.

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#### Drug Facts (continued)

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This product is not manufactured or distributed by the owner of the registered trademark Benadryl®.

CONVENIENT RECLOSING TAB



OPEN OTHER END

46267 63 07



edible ink, FD&C blue no. 1, FD&C red no. 40, gelatin, magnesium stearate, methylparaben, polysorbate 80, propylparaben, sodium lauryl sulfate, sodium propionate

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**Publix.**



## ALLERGY

diphenhydramine hydrochloride capsule

### Product Information

|                         |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:56062-462 |
| 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 |
|---|----------|
| ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)        |          |
| BENZYL ALCOHOL (UNII: LKG8494WBH)           |          |
| BUTYLPARABEN (UNII: 3QP1U3FV8)              |          |
| D&C RED NO. 28 (UNII: 767IP0Y5NH)           |          |
| EDETATE CALCIUM DISODIUM (UNII: 25IH6R4SGF) |          |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD)          |          |
| FD&C RED NO. 40 (UNII: WZB9127XOA)          |          |
| GELATIN (UNII: 2G86QN327L)                  |          |
| MAGNESIUM STEARATE (UNII: 70097M6I30)       |          |
| METHYLPARABEN (UNII: A2I8C7HI9T)            |          |
| POLYSORBATE 80 (UNII: 6OZP39ZG8H)           |          |
| PROPYLPARABEN (UNII: Z8IX2SC1OH)            |          |
| SODIUM LAURYL SULFATE (UNII: 368GB5141J)    |          |
| SODIUM PROPIONATE (UNII: DK6Y9P42IN)        |          |

### Product Characteristics

|          |   |              |          |
|----------|---|--------------|----------|
| Color    | PINK (clear) , WHITE (clear) , RED (band) | Score        | no score |
| Shape    | CAPSULE                                   | Size         | 14mm     |
| Flavor   |   | Imprint Code | L462     |
| Contains |   |              |          |

### Packaging

| # | Item Code        | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---------------------|----------------------|--------------------|
| 1 | NDC:56062-462-62 | 24 in 1 CARTON      | 08/31/1993           |                    |

|                              |   |  |                             |                           |
|------------------------------|---|--|-----------------------------|---------------------------|
| 1                            |   | 1 in 1 BLISTER PACK; Type 0: Not a Combination Product |                             |                           |
| 2                            | NDC:56062-462-67                                | 48 in 1 CARTON   | 04/18/2002                  |                           |
| 2                            |   | 1 in 1 BLISTER PACK; Type 0: Not a Combination Product |                             |                           |
| <b>Marketing Information</b> |   |  |                             |                           |
| <b>Marketing Category</b>    | <b>Application Number or Monograph Citation</b> |  | <b>Marketing Start Date</b> | <b>Marketing End Date</b> |
| OTC monograph final          | part341   |  | 08/31/1993                  |                           |

**Labeler** - Publix Super Markets Inc (006922009)

Revised: 1/2019

Publix Super Markets Inc