

ALLERGY RELIEF- diphenhydramine hydrochloride capsule
RETAIL BUSINESS SERVICES , 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.

CAR-1113-2020-0817

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

- 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 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 times 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

- store between 20-25°C (68-77°F)
- retain carton for complete product information

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 lauryl sulfate

PRINCIPAL DISPLAY PANEL

CAREONE®

NDC 72476-113-01

Compare to the active ingredient in Benadryl® Allergy†

ALLERGY RELIEF

Diphenhydramine HCl, 25 mg

Antihistamine

FOR RELIEF OF:

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

Actual Size

24 CAPSULES



ALLERGY RELIEF

diphenhydramine hydrochloride capsule

Product Information

| | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:72476-113 |
| 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: 3QP1U3FV8) | |
| D&C RED NO. 28 (UNII: 767IP0Y5NH) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| 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, white | Score | no score |
| Shape | OVAL | Size | 14mm |
| Flavor | | Imprint Code | A;20 |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:72476-113-01 | 2 in 1 CARTON | 08/01/2020 | |
| 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 | 08/01/2020 | |

Labeler - RETAIL BUSINESS SERVICES , LLC. (967989935)

Revised: 9/2020

RETAIL BUSINESS SERVICES , LLC.