

**GOOD SENSE ALLERGY RELIEF- diphenhydramine hydrochloride capsule
Proficient Rx LP**

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.

Perrigo Allergy Relief 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

- with any other product containing diphenhydramine, even one used on skin
- to make a child sleepy

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

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
- 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 this product in children under 6 years of age

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

Questions or comments?

1-800-719-9260

Principal Display Panel

Easy to Swallow

Allergy Relief

Capsules

Diphenhydramine Hydrochloride

Antihistamine

Actual Size

Sneezing

Itchy, Watery Eyes

Runny Nose

Itchy Throat

Compare to active ingredient of Benadryl® Allergy



NDC 63187-167-24

Lot #:00000
Exp. 00/00/00
SN# MASTER

AllergyRelief 25mg

#24 CapsulesEach capsule contains: Diphenhydramine HCl 25mg
Antihistamine

See box

Product ID: RD016724

Dist. By: Perrigo Allegan, MI 49010

Store at 20°-25°C (68°-77°F)

Keep medication out of the reach of children

Relabeled By: Proficient Rx LP
Thousand Oaks, CA 91320AllergyRelief 25mg
#24 Capsules
Lot #:00000 SN#MASTER
NDC 63187-167-24 Exp:00/00/00AllergyRelief 25mg
#24 Capsules
Lot #:00000 SN#MASTER
NDC 63187-167-24 Exp:00/00/00AllergyRelief 25mg
#24 Capsules
Lot #:00000 SN#MASTER
NDC 63187-167-24 Exp:00/00/00

GOOD SENSE ALLERGY RELIEF

diphenhydramine hydrochloride capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63187-167(NDC:0113-0462)
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)	
BUTYL PARABEN (UNII: 3QPI1U3FV8)	
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, UNSPECIFIED (UNII: 2G86QN327L)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYL PARABEN (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:63187-167-24	2 in 1 CARTON	01/01/2019	
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	09/15/1989	

Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(63187-167) , RELABEL(63187-167)

Revised: 1/2021

Proficient Rx LP