ALLERGY RELIEF- diphenhydramine hcl capsule
Geiss, Destin & Dunn Inc.

GoodSense 44-190

Active ingredient (in each banded capsule)

Diphenhydramine HCl 25 mg

Purpose

Antihistamine

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
  - runny nose
  - itching of the nose or throat
  - itchy, watery eyes
  - sneezing
- 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
- use caution 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.

Directions

- **do not take more than directed**
- 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

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN OR IF RED BAND AROUND CAPSULE IS BROKEN OR MISSING**
- store at 25ºC (77ºF); excursions permitted between 15º-30ºC (59º-86ºF)
- protect from moisture
- see end flap for expiration date and lot number

Inactive ingredients

butylparaben, corn starch, D&C red #28, edible ink, FD&C blue #1, FD&C red #40, gelatin, lactose anhydrous, magnesium stearate, methylparaben, polysorbate 80, propylparaben, silicon dioxide

Questions or comments?

1-800-426-9391

Principal Display Panel

GOODSENSE®

NDC 50804-091-08

Allergy Relief

Diphenhydramine HCl, 25 mg
Antihistamine

Sneezing,
Itchy, Watery Eyes

Runny Nose,
Itchy Throat

*Compare to the active ingredient of
Benadryl® Allergy

actual size

24 Capsules

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN OR BROKEN OR IF RED BAND AROUND CAPSULE IS BROKEN OR MISSING

*This product is not manufactured or distributed by Kenvue Inc., owner of the registered trademark Benadryl® Allergy.
50844 REV0524C19008

Distributed by: Perrigo Direct, Inc.
Peachtree City, GA 30269
www.PerrigoDirect.com (1-800-426-9391)
GoodSense® is a registered trademark of
L. Perrigo Company.

GOODSENSE®

Allergy Relief

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NDC 50804-091-08

Allergy Relief

Diphenhydramine HCl, 25 mg
Antihistamine

actual size

Sneezing,
Itchy, Watery Eyes
Runny Nose,
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no print / no varnish area
lot no. & exp. date

8 46036 00911 8

GOODSENSE®

Allergy Relief

B-0560-190-08-RD
REV0524C19008

Distributed by: Perrigo Direct, Inc.
Peachtree City, GA 30269
www.PerrigoDirect.com (1-800-426-9391)
GoodSense® is a registered trademark of L. Perrigo Company.

Warnings

Do not use

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- 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

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- avoid alcoholic beverages
- use caution 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.

children 6 to under 12 years	1 capsule
children under 6 years	do not use

Other information

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Good Sense 44-190

ALLERGY RELIEF

diphenhydramine hcl capsule

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-091
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**
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
STARCH, CORN (UNII: O8232NY3SJ)	
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics			
Color	pink, white	**Score**	no score
Shape	CAPSULE	**Size**	14mm
Flavor		**Imprint Code**	44;107
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50804-091-08	2 in 1 CARTON	06/02/2020	
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 Drug	M012	06/02/2020	

Labeler - Geiss, Destin & Dunn Inc. (076059836)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(50804-091)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	pack(50804-091)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(50804-091)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(50804-091)

Revised: 6/2025 Geiss, Destin & Dunn Inc.