

**ALLERGY RELIEF- diphenhydramine hcl capsule
DOLGENCORP, LLC**

Dollar General 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
 - 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

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- 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
- alcohol, sedatives, and tranquilizers may increase drowsiness
- 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.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

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

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 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- 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?

1-888-309-9030

Principal display panel

DG™ | health

Compare to
active ingredient
of Benadryl®
Allergy*

Allergy Relief

**Diphenhydramine HCl, 25 mg
Antihistamine**

Temporarily Relieves:

- Sneezing
- Itchy, watery eyes
- Runny nose
- Itchy nose or throat

Each capsule individually
banded for your protection

24 Capsules

Actual Capsule Size

**25
mg each**

**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**

**100%
Satisfaction
Guaranteed!**
(888) 309-9030

DISTRIBUTED BY
OLD EAST MAIN CO.
100 MISSION RIDGE
GOODLETTSVILLE, TN 37072

*This product is not manufactured or distributed by
Johnson & Johnson Corporation, owner of the
registered trademark Benadryl® Allergy.
50844 ORG111719008

Allergy Relief

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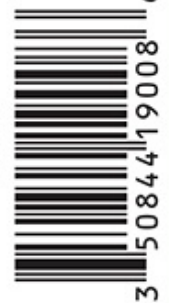
Actual Capsule Size

DG™ | health

Compare to active ingredient of Benadryl® Allergy*

Allergy Relief

B-0315-190-08
ORG111719008



A0505

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

No print/No varnish
Lot & Exp date

Drug Facts

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

Active ingredient (in each banded capsule)
Diphenhydramine HCl 25 mg 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
a breathing problem such as emphysema or chronic bronchitis
glaucoma
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
alcohol, sedatives, and tranquilizers may increase drowsiness
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.

Drug Facts (continued)

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions
take every 4 to 6 hours
do not take more than 6 doses in 24 hours
adults and children 12 to 2 capsules
children 6 to under 12 1 capsule
children under 6 years do not use

Other information
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store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
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? 1-888-309-9030
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50844 ORG111719008

083000

100% Satisfaction Guaranteed!
(888) 309-9030

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OLD EAST MAIN CO.
100 MISSION RIDGE
GOODLETTSVILLE, TN 37072

DG Health 44-190

ALLERGY RELIEF

diphenhydramine hcl capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-191
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

		Marketing Start	Marketing End
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-191-08	2 in 1 CARTON	07/30/2020	
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	07/30/2020	

Labeler - DOLGENCORP, LLC (068331990)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(55910-191)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	pack(55910-191)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(55910-191)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(55910-191)

Revised: 6/2024

DOLGENCORP, LLC