ALLERGY ANTIHISTAMINE- diphenhydramine hydrochloride tablet
P & L Development, 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.

Drug Facts

Active ingredient (in each tablet)

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	take 1 to 2 tablets
children 6 to under 12 years	take 1 tablet
children under 6 years	do not use this

Other information

- each tablet contains: **calcium 25 mg**
- store between 20-25°C (68-77°F)
- protect from light and moisture

Inactive ingredients

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, D&C Red #27 Aluminum Lake, dibasic calcium phosphate dihydrate, hypromellose, lecithin*, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, polyvinyl alcohol*, purified water*, talc*, titanium dioxide

* contains one or more of these ingredients

Questions or comments?

Call **1-877-753-3935 Monday-Friday 9AM-5PM EST**

Principal Display Panel

Compare to the active ingredient in **Benadryl® Allergy****

Allergy relief

Diphenhydramine HCl, 25 mg

Antihistamine

for allergy relief

- runny nose
- sneezing
- itchy, watery eyes
- Itchy nose or throat

TABLETS

†This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Benadryl® Allergy.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Distributed by:

PL Developments

200 Hicks Street

Westbury, NY 11590

Product Label

ready in case®

allergy relief

diphenhydramine HCl 25 mg

Compare to the active ingredient in
Benadryl® Allergy**

NDC 59726-691-48

allergy relief

diphenhydramine HCl 25 mg

antihistamine
for allergy relief:

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Drug Facts (continued)

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ReadyinCase Allergy Relief

ALLERGY ANTIHISTAMINE

diphenhydramine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59726-691
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**
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
CALCIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: O7TSZ97GEP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
ALUMINUM OXIDE (UNII: LMI26O6933)	

EGG PHOSPHOLIPIDS (UNII: 1Z74184RGV)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	pink	**Score**	no score
Shape	CAPSULE	**Size**	12mm
Flavor		**Imprint Code**	T;061;V;25;S4
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59726-691-12	2 in 1 CARTON	08/27/2013	01/05/2024
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:59726-691-01	1 in 1 BOX	08/27/2013	01/05/2024
2		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:59726-691-04	400 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/27/2013	01/05/2024
4	NDC:59726-691-48	48 in 1 CARTON	08/27/2013	01/05/2024
4		1 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	08/27/2013	01/05/2024

Labeler - P & L Development, LLC (800014821)

Revised: 10/2021 P & L Development, LLC