

ALLERGY RELIEF 25 MG- diphenhydramine hcl 25mg capsule
Ulai Health LLC

Drug Facts

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
- 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

- drowsiness may occur
- avoid alcoholic drinks

- 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 (1-800-222-1222).

Directions

- take every 4 to 6 hours, or as directed by a doctor
- do not exceed 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	ask a doctor

Other information

- store at 20-25 °C (68-77 °F)

Inactive ingredients

black iron oxide, D&C red #28, FD&C blue #1, FD&C red #40, gelatin, lactose monohydrate, magnesium stearate, silicon dioxide, sodium lauryl sulfate

Questions or comments?

(866) 562-2756 (Mon- Fri 8 AM to 4 PM EST)

PHARBEST

NDC 73057-348-08

Manufactured in the USA

***COMPARE TO the active ingredient in**

BENADRYL ® ALLERGY ULTRATAB ®

ALLERGY RELIEF

Antihistamine - Diphenhydramine HCl 25 mg

- Itchy, Watery Eyes • Runny Nose

• Itchy Throat • Sneezing

100 CAPSULES



PHARBEST  Manufactured in the USA

NDC 73057-348-08

***COMPARE TO the active ingredient in BENADRYL® ALLERGY ULTRATAB®**

ALLERGY RELIEF

Antihistamine - Diphenhydramine HCl 25 mg

TAMPER EVIDENT: DO NOT USE IF SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Drug Facts

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Purpose
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

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When using this product ■ drowsiness may occur ■ avoid alcoholic drinks

Org. 06/19

Peel here for more drug facts



3 7 3 0 5 7 1 3 4 8 0 8 2

Distributed by:
ULAI HEALTH LLC., Alexandria, VA 22312

Drug Facts (continued)

■ alcohol, sedatives, and tranquilizers may increase drowsiness ■ use caution when driving a motor vehicle or operating machinery ■ excitability may occur, especially in children

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This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of registered trademark of Benadryl® Allergy Ultratab®.

STOP PEELING

ALLERGY RELIEF 25 MG

diphenhydramine hcl 25mg capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73057-348
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
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	
D&C RED NO. 28 (UNII: 7671P0Y5NH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Product Characteristics

Color	pink (Pink/ Clear)	Score	no score
Shape	CAPSULE	Size	14mm
Flavor		Imprint Code	PH014
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73057-348-08	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2019	
2	NDC:73057-348-03	24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2019	
3	NDC:73057-348-11	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M010	07/01/2019	

Labeler - Ulai Health LLC (081181535)

Registrant - Pharbest Pharmaceuticals, Inc. (557054835)

Establishment

Name	Address	ID/FEI	Business Operations
Pharbest Pharmaceuticals, Inc.		557054835	manufacture(73057-348) , analysis(73057-348) , pack(73057-348) , label(73057-348)

Revised: 1/2025

Ulai Health LLC