

**ALLERGY RELIEF- diphenhydramine hcl tablet
TARGET CORPORATION**

160R Target 11673-494 Diphenhydramine HCL 25 mg Tablets

Drug Facts

Active ingredients (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 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 times in 24 hours

adults and children 12 years and over	1 to 2 tablets
children 6 to under 12 years	1 tablet
children under 6 years	do not use

Other information

- **each tablet contains:**calcium 25 mg
- protect from light
- store at 25°C (77° F) excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package

Inactive ingredients carnauba wax, colloidal silicon dioxide, croscarmellose sodium, D&C red #27 aluminum lake, dibasic calcium phosphate dihydrate, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol (PEG) 400, polysorbate 80, titanium dioxide

Questions or comments?

Call **1-800-910-6874**

Compare to active ingredient in Benadryl® Allergy Ultratab®*

Allergy Relief

Diphenhydramine HCl, 25 mg / Antihistamine



300 TABLETS

For the temporary relief of:
sneezing, itchy and watery eyes,
runny nose, itchy throat

300
Count

Adhesive Area

TAMPER EVIDENT: DO NOT USE THIS PRODUCT IF THE IMPRINTED FOIL SEAL OVER THE MOUTH OF THE BOTTLE IS CUT, TORN, BROKEN OR MISSING

Drug Facts

Active ingredient (in each tablet)

Diphenhydramine HCl 25 mg.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 enlarged prostate gland

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers

Drug Facts (continued under label)

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

NDC 11673-494-32

160R 0923

Dist. by Target Corp.
Mpls., MN 55403
TM & © 2024
Target Brands, Inc.

094 04 0156 R00 C-002262-01-009-0000



3 11673 49432 5

Lot No.:

Exp. Date:

Varnish Omit Area

Drug Facts (continued)

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

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Allergy Relief



Diphenhydramine HCl, 25 mg / Antihistamine

100 TABLETS

Adhesive Area

READ AND KEEP CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

TAMPER EVIDENT: DO NOT USE THIS PRODUCT IF THE IMPRINTED FOIL SEAL OVER THE MOUTH OF THE BOTTLE IS CUT, TORN, BROKEN OR MISSING

Drug Facts

Active ingredient (in each tablet) Purpose
Diphenhydramine HCl, 25 mg 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

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Drug Facts (continued under label)

160R 0923 NDC 11673-494-01
094040375 R00 C-002262-01-009-0000
Distributed by Target Corporation
Minneapolis, MN 55403
TM & ©2024 Target Brands, Inc.

Lot No.:

Exp. Date:

Varnish Omit Area

Drug Facts (continued)

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adults and children 12 years and over

1 to 2 tablets

children 6 to under 12 years

1 tablet

children under 6 years

do not use

Other information

■ each tablet contains calcium 25 mg

■ protect from light

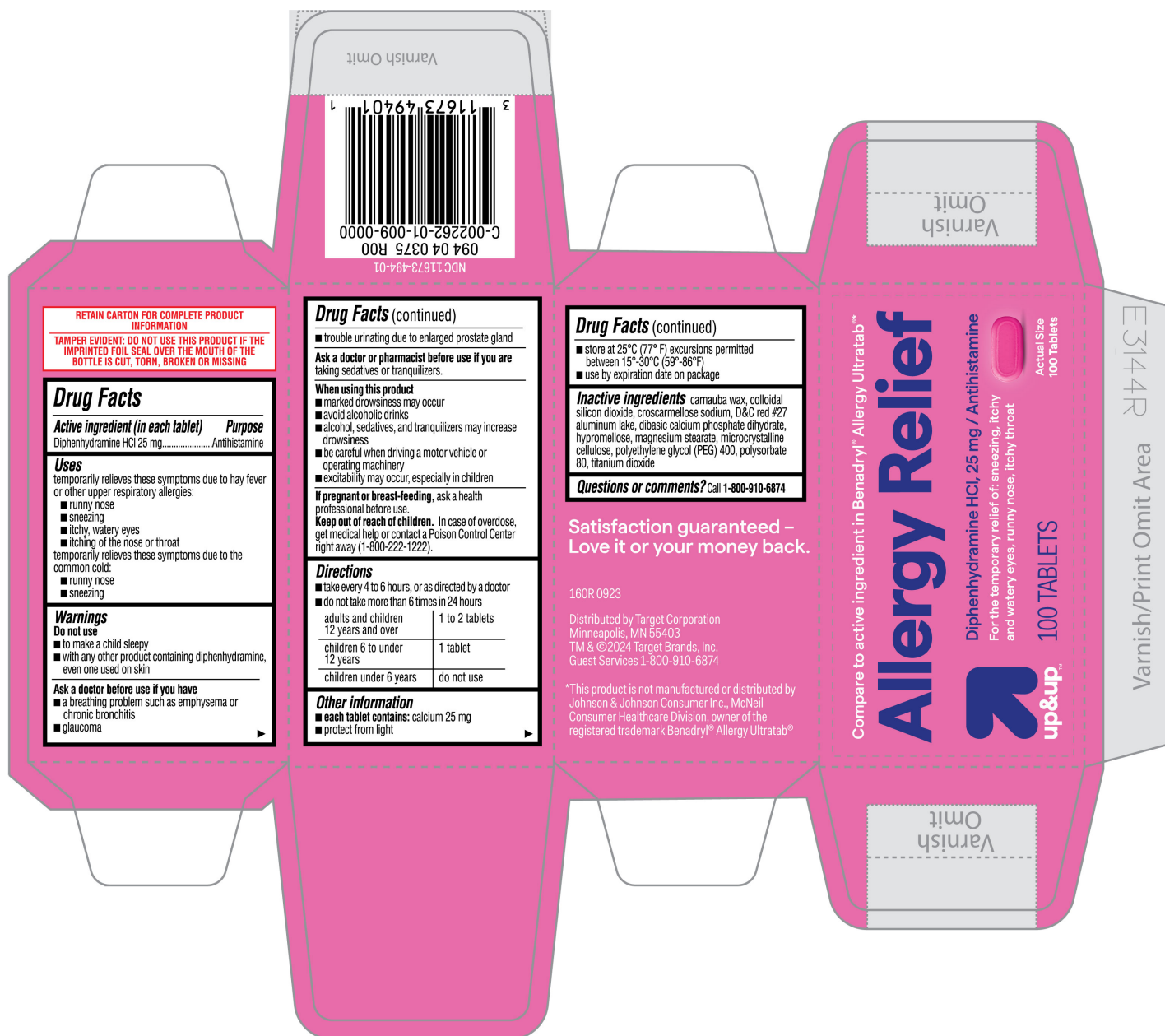
■ store at 25°C (77° F); excursions permitted between 15°-30° C (59°-86° F)

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Questions or comments?

Call 1-800-910-6874





ALLERGY RELIEF

diphenhydramine hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-494
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
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	

Product Characteristics

Color	pink	Score	no score
Shape	CAPSULE	Size	11mm
Flavor		Imprint Code	T;061
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-494-32	300 in 1 BOTTLE; Type 0: Not a Combination Product	10/20/2023	
2	NDC:11673-494-01	1 in 1 CARTON	10/20/2023	
2		100 in 1 BOTTLE; 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	10/20/2023	

Labeler - TARGET CORPORATION (006961700)

Registrant - TIME CAP LABORATORIES, INC. (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
TIME CAP LABORATORIES, INC.		037052099	manufacture(11673-494)