ALLERGY RELIEF- diphenhydramine hcl tablet
DOLGENCORP INC

Diphenhydramine HCl 50 mg tablet, Dollar General, allergy relief

Active ingredient (in each capsule)

Diphenhydramine HCl 50 mg

Purpose

Antihistamine

Uses

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

Warnings

■ May cause excitability especially in children.

■ May cause marked drowsiness; alcohol, sedatives, and tranquilizres may increase the drowsiness effect.

Do not use

■ for children under 12 years of age ■ 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

■ avoid alcoholic beverages ■ be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

sleeplessness persists continuously for more than 2 weeks

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 Centre right away.

Directions

adults and children 12 years and over	■ take 1 tablet every 4 to 6 hours, or as directed by a doctor ■ do not take more than 6 tablets in 24 hours
children under 12 years	do not use

Other information

■ each tablet contains: calcium 36 mg ■ store between 20-25°C (68-77°F). Protect from light.

■ **do not use if blister unit is torn or broken**

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide

Questions or comments?

call toll-free 1-888-235-2466 (Mon - Fri 9AM - 5PM EST)

†This product is not manufactured or distributed by the owners of Benadryl® Allergy Extra Strength.

THIS PRODUCT IS PACKAGED IN A CHILD-RESISTANT AND

TAMPER-EVIDENT PACKAGE. USE ONLY IF BLISTERS ARE INTACT.

KEEP OUTER CARTON FOR COMPLETE

WARNINGS AND PRODUCT INFORMATION.

DISTRIBUTED BY OLD EAST MAIN CO.

100 MISSION RIDGE GOODLETTSVILLE, TN 37072

MADE IN INDIA

CODE: TN/DRUGS/TN00002222/2006

100%

Satisfaction Guaranteed!

(888) 309-9030

949129331

L0000959

R0225

carton

†Compare to the

active ingredient of
Benadryl® Allergy

Extra Strength

DG ™ health

Extra Strength

Allergy Relief

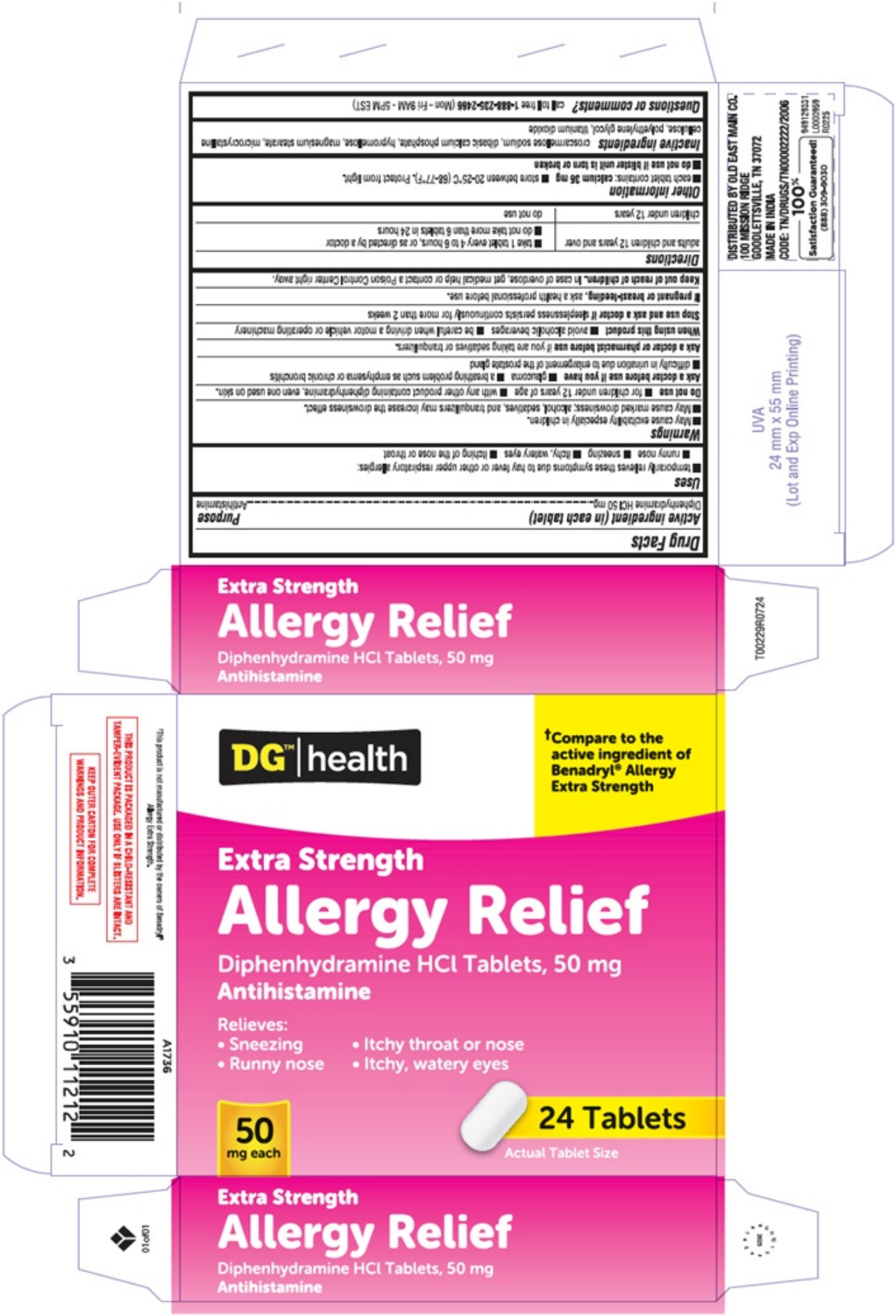
diphenhydramine HCl Tablets, 50 mg
Antihistamine

Relieves:
• Sneezing • Itchy throat or nose
• Runny nose • Itchy, watery eyes

50

mg each

24 Tablets



Extra Strength
Allergy Relief
Diphenhydramine HCl Tablets, 50 mg
Antihistamine

DG health

†Compare to the active ingredient of Benadryl® Allergy Extra Strength

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Actual Tablet Size

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Antihistamine

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UVA
24 mm x 55 mm
(Lot and Exp Online Printing)

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100% Satisfaction Guaranteed! (888) 309-9030

849123231
L0002869
R0225

Drug Facts

Active ingredient (in each tablet)	Purpose
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  - runny nose
  - sneezing
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ALLERGY RELIEF

diphenhydramine hcl tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-450
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	50 mg

Inactive Ingredients	
Ingredient Name	**Strength**
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

Product Characteristics			
Color	white	**Score**	no score
Shape	CAPSULE	**Size**	14mm
Flavor		**Imprint Code**	DH50
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55910-450-12	2 in 1 CARTON	06/24/2025	
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	06/24/2025	

Labeler - DOLGENCORP INC (068331990)

Registrant - Bionpharma Inc. (079637826)

Establishment

Name	Address	ID/FEI	Business Operations
OrBion Pharmaceuticals Private Limited		854403569	manufacture(55910-450) , pack(55910-450)

Revised: 6/2025 DOLGENCORP INC