ALLERGY RELIEF- diphenhydramine hydrochloride tablet, film coated
Gobrands, Inc

Allergy Relief

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
• 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 • 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
• 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 • avoid alcoholic beverages

Stop use and ask a doctor if sleeplessness persists continuously for more than 2 weeks. Insomnia may be a symptom of serious underlying medical illness.

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 only one dose per day (24 hours)

adults & children 12 years & over	take 2 softgels at bedtime if needed or as directed by a doctor
children under 12 years	do not use

Other information

• store at 20-25° C (68-77° F) • avoid excessive heat above 40° C (104° F) and high humidity • Protect from light

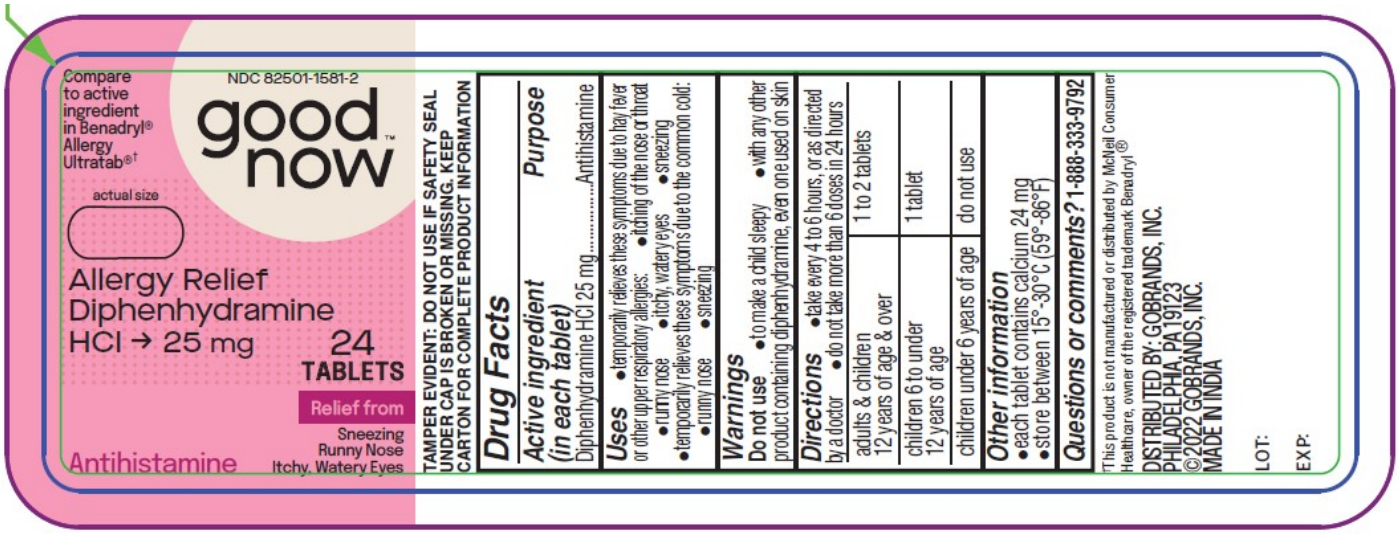
Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, dicalcium phosphate, D&C red # 27 aluminum lake, lecithin, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, talc, titanium dioxide

Questions or comments?

1-888-333-9792

PRINCIPAL DISPLAY PANEL



ALLERGY RELIEF

diphenhydramine hydrochloride tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82501-1581
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**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
CALCIUM PHOSPHATE (UNII: 97Z1WI3NDX)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	pink	**Score**	no score
Shape	OVAL	**Size**	11mm
Flavor		**Imprint Code**	S4
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82501-1581-2	1 in 1 CARTON	05/25/2022	
1		24 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	M017	05/25/2022	

Labeler - Gobrands, Inc (057499049)

Revised: 12/2023 Gobrands, Inc