ALLERGY RELIEF- diphenhydramine hydrochloride tablet, film coated
Strategic Sourcing Services, LLC

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• 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
• trouble urinating due to an enlarged prostate gland
• glaucoma

Ask a doctor or pharmacist before use if you aretaking 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 of age and over - 1 to 2 tablets
• children 6 to under 12 years of age - 1 tablet
• children under 6 years of age - do not use

Other information• each tablet containscalcium 24 mg
• store between 20°-25° C (68°-77° F).
• Protect from light

Inactive ingredients
c olloidal silicon dioxide, croscarmellose sodium, dicalcium phosphate, D&C red # 27 aluminium lake, lecithin, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, talc, titanium dioxide

Questions or comments?
1-833-358-6431

PRINCIPAL DISPLAY PANEL

ALLERGY RELIEF

diphenhydramine hydrochloride tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70677-1238
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40)	DIPHENHYDRAMINE	25 mg

(DIPHENHYDRAMINE - UNII:8GTS82S83M)	HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)	
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:70677-1238-1	120 in 1 BOTTLE; Type 0: Not a Combination Product	01/24/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/24/2024	

Labeler - Strategic Sourcing Services, LLC (116956644)

Revised: 1/2024 Strategic Sourcing Services, LLC