

DIPHENHYDRAMINE HYDROCHLORIDE- diphenhydramine hydrochloride tablet, film coated

HF Acquisition Co LLC, DBA HealthFirst

Diphenhydramine Hydrochloride 25 mg

Antihistamine

USES

Temporarily relieves these symptoms due to hay fever or other respiratory allergies

runny nose

sneezing

itching of the nose or throat

itchy, watery eyes

Temporarily relieves these symptoms due to the common cold

runny nose

sneezing

Do not use – Diphen Allergy Caplet

to make a child sleepy

with any other product containing diphenhydramine, even one that is used on skin.

Ask a doctor before use if you have – Diphen Allergy Caplet

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 are – Diphen Allergy Caplet

taking sedatives or tranquilizers

When using this product – Diphen Allergy Caplet

marked drowsiness may occur

avoid alcoholic drinks

alcohol, sedatives and tranquilizers may increase the drowsiness effect

use caution when driving a motor vehicle or operating machinery

excitability may occur, especially in children

If pregnant or breast-feeding, – Diphen Allergy Caplet

ask a health care professional before use.

Keep out of reach of children. – Diphen Allergy Caplet

In case of overdose, contact a physician or poison control center right away (1-800-222-1222).

DIRECTIONS

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protect from light

store at room temperature 68°-77°F (20°-25°C)

tamper-evident sealed packets

do not use any opened or torn packets

croscarmellose sodium, D&C red #27 aluminum lake, hypromellose, lactose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, silicon dioxide, titanium dioxide

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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
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I3O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	pink	Score	no score
Shape	OVAL	Size	11mm
Flavor		Imprint Code	048;D
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51662-1674-1	1 in 1 PACKET; Type 0: Not a Combination Product	01/01/2012	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/01/2012	

Labeler - HF Acquisition Co LLC, DBA HealthFirst (045657305)

Registrant - HF Acquisition Co LLC, DBA HealthFirst (045657305)

Establishment			
Name	Address	ID/FEI	Business Operations
HF Acquisition Co LLC, DBA HealthFirst		045657305	repack(51662-1674)

