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

Allergy Relief (Cetirizine HCI Tablets)

Drug Facts

Active ingredient (in each tablet)
Cetirizine HCl 10 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

Warnings

Do not useif you have ever had an allergic reaction to this product or any of its ingredients or to an antihistamine containing hydroxyzine.

Ask a doctor before use if you haveliver or kidney disease. Your doctor should determine if you need a different dose.

Ask a doctor or pharmacist before use if you aretaking tranquilizers or sedatives.

When using this product

- drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor ifan allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding:

- if breast-feeding: not recommended
- if pregnant: 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

- adults and children 6 years and over: One 10 mg tablet once daily; do not take more than one 10 mg tablet in 24 hours. A 5 mg product may be appropriate for less severe symptoms
- adults 65 years and over : ask a doctor
- children under 6 years of age : ask a doctor
- consumers with liver or kidney disease : ask a doctor

Other information store between 20° to 25°C (68° to 77°F)

**containes one or more of these ingrdients

Inactive ingredients colloidal silicon dioxide**, croscarmellose sodium**, hypromellose, lactose, magnesium stearate, maize starch**, microcrystalline cellulose**, polyethylene glycol, povidone**, titanium dioxide

Questions or comments?

833-358-6431



ALLERGY RELIEF

cetirizine hydrochloride tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-1241
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	

ı	CETIRIZINE HYDROCHLORIDE (UNII: 64O047KTOA) (CETIRIZINE -	CETIRIZ INE	10 mg
ı	UNII:YO7261MF24)	CETIKIZINE	10 mg

Inactive Ingredients			
Ingredient Name	Strength		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
MAIZE INVERT SUGAR (UNII: ED959S6ACY)			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POVIDONE (UNII: FZ989GH94E)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			

Product Characteristics			
Color	white (white to off-white)	Score	no score
Shape	BULLET (barrel shaped, biconvex)	Size	8mm
Flavor		Imprint Code	CTN;10
Contains			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:70677- 1241-1	120 in 1 BOTTLE; Type 0: Not a Combination Product	01/24/2024	

Marketing I	ng Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA077829	01/23/2024		

Labeler - Strategic Sourcing Services LLC (116956644)

Revised: 2/2024 Strategic Sourcing Services LLC