ALLEXPERT 24HR ANTIHISTAMINE CETIRIZINE HYDROCHLORIDE ORAL SOLUTION- allexpert 24hr antihistamine cetirizine hydrochloride solution
Bowong Technology (ShenZhen) Co.,Ltd

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Active Ingredient(s)

Cetirizine HCl 5mg.(in each 5ml)

Purpose

Antihistamine

Use

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

1.ASK A DOCTOR before use if you have Iver or kidney disoaso. Your doctor should dotermine il you need a dilferent dose.
2.ASK A DOCTOR or phamaclst belore use i you aro taking tranqulzors or sodativcs.

Do not use

if you have over had an allorgic reaclion lo ths product or any ol ils ingredients or to an anlihislamine conlaining hydroxydine
or other Piperazine derivatives .

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

if an allergic reoclion lo this product occurs. Seek medical help right away.

In cose of overdose. get medical help or contact a Poison Control Center right away.

Directions

See Outer box

Other information

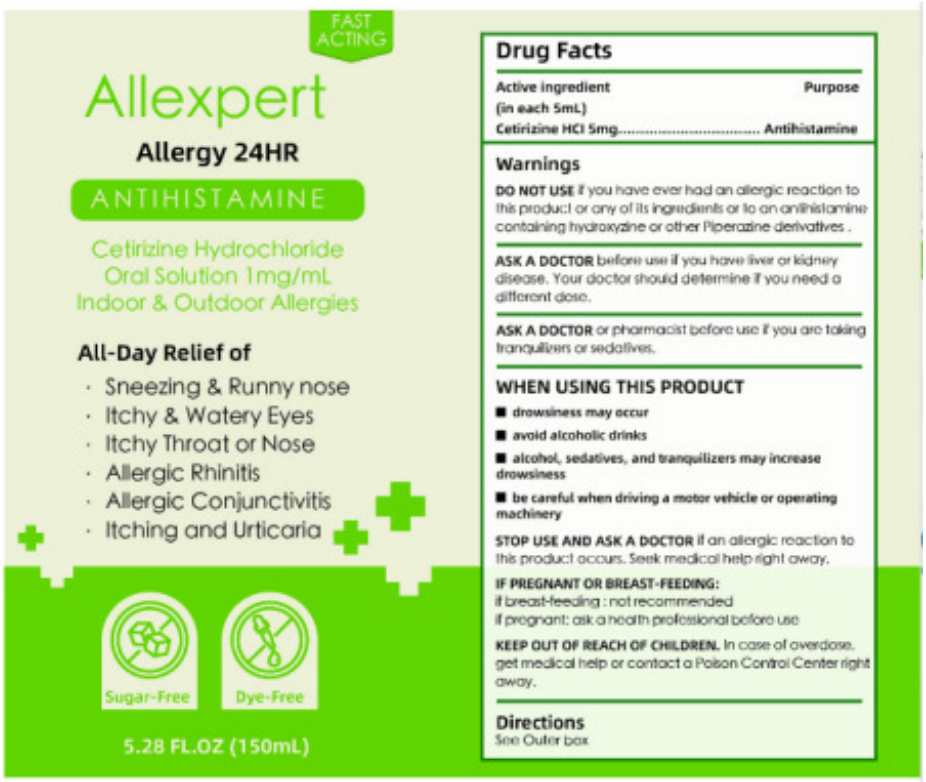
■store between 20* to 25C (68* to 77*F)

■see bottom panel for lot number and expiration date.

Inactive ingredients

Sodium Dihydrogen Phosphate, Dibasic
Sodium Phosphate, Sodium Benzoate,
Aspartame, purified water.

Package Label - Principal Display Panel



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allexpert 24hr antihistamine cetirizine hydrochloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84264-001
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BENZOATE (UNII: OJ245FE5EU)	
ASPARTAME (UNII: Z0H242BBR1)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
WATER (UNII: 059QF0KO0R)	
AZITHROMYCIN SODIUM DIHYDROGEN PHOSPHATE (UNII: ADV7FS82PV)	

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End

#	Item Code	Package Description	Date	Date
1	NDC:84264-001-01	150 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/17/2024	

Labeler - Bowong Technology (ShenZhen) Co.,Ltd (844456666)

Establishment

Name	Address	ID/FEI	Business Operations
Bowong Technology (ShenZhen) Co.,Ltd		844456666	manufacture(84264-001)

Revised: 4/2024 Bowong Technology (ShenZhen) Co.,Ltd