ALL DAY ALLERGY RELIEF- cetirizine hcl tablet
P & L Development, LLC

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 use

if 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 have

liver or kidney disease. Your doctor should determine if you need a different dose.

Ask a doctor or pharmacist before use if you are

taking tranquilizers or sedatives.

When using this product

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

Stop use and ask a doctor if

an 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.

Directions

adults and children 6 years and over	Take 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 20º to 25ºC (68º to 77ºF)

Inactive ingredients

lactose monohydrate, magnesium stearate, polyethylene glycol, polyvinyl alcohol, povidone, starch, talc, titanium dioxide

Questions or comments?

Call **1-877-753-3935 Monday-Friday 9AM-5PM EST**

Principal Display Panel

Compare to the active ingredient in **Zyrtec**®†

all day

allergy relief

cetirizine HCl tablets 10 mg

antihistamine

24 hour relief of:

- sneezing
- itchy, watery eyes
- runny nose
- itchy throat or nose

tablets

†This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Zyrtec®.

Distributed by:

PL Developments

200 Hicks Street

Westbury, NY 11590

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Package Labeling

ready in case®

all day allergy relief

cetirizine HCl 10 mg
antihistamine

Compare to the active ingredient in **Zyrtec®***

NDC 59726-129-05

ready in case®

all day
allergy relief

cetirizine HCl 10 mg
antihistamine

treats indoor & outdoor allergies

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

SEE CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Drug Facts

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Cetirizine HCl 10 mg	Antihistamine

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Drug Facts (continued)

If pregnant or breast-feeding:

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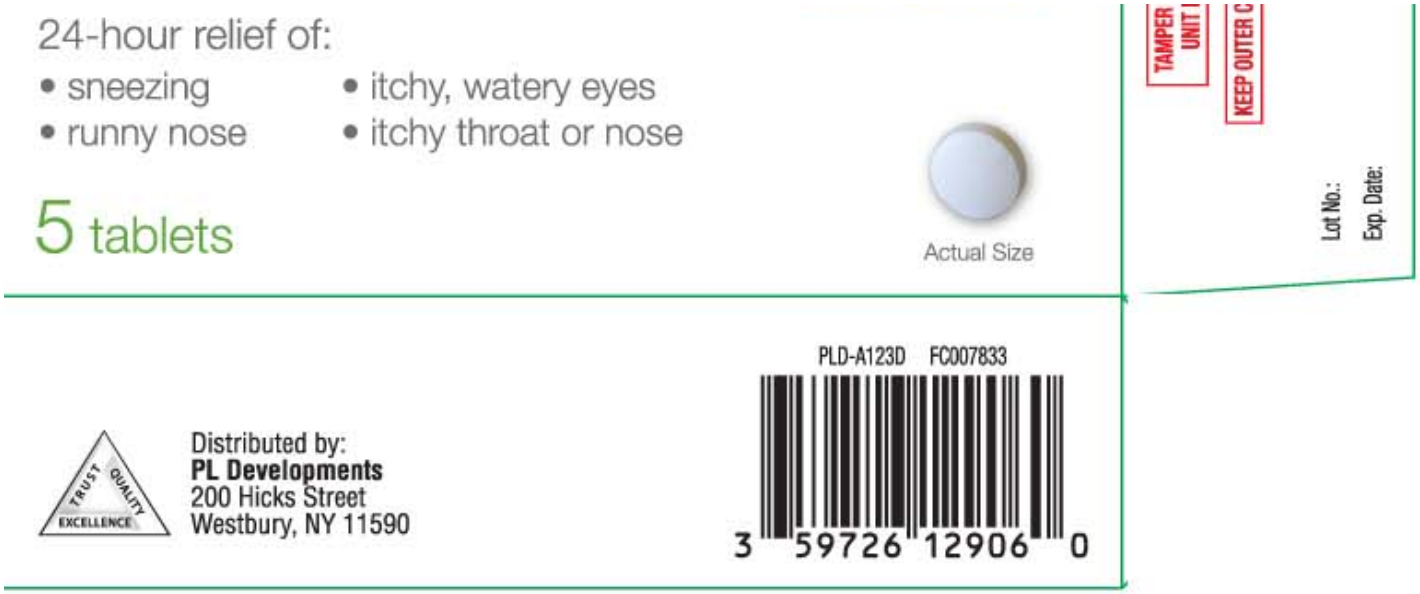

READYinCASE All Day Allergy Relief

ALL DAY ALLERGY RELIEF

cetirizine hcl tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59726-129
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CETIRIZINE HYDROCHLORIDE (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	10 mg

Inactive Ingredients	
Ingredient Name	**Strength**
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	

Product Characteristics			
Color	white (white)	**Score**	no score
Shape	OVAL (oval)	**Size**	8mm
Flavor		**Imprint Code**	IP;46

Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59726-129-14	1 in 1 CARTON	09/01/2010	
1		14 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:59726-129-05	1 in 1 CARTON	09/01/2010	
2		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:59726-129-03	1 in 1 BOX	09/01/2010	
3		300 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078780	09/01/2010	

Labeler - P & L Development, LLC (800014821)

Revised: 12/2022 P & L Development, LLC