ALLERGY RELIEF- cetirizine hcl tablet
NUVICARE LLC

Allergy Relief- Cetirizine HCl 10 mg

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 or pharmacist before use if you are taking sedatives or tranquilizers.

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

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 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 (1-800 222-1222) right away.

Directions

Adults and children 6 years of age and older: 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.

Children under 6 years of age: Ask a doctor.

Adults 65 years of age and older: Ask a doctor.

Consumers with liver or kidney disease: Ask a doctor

Other Information

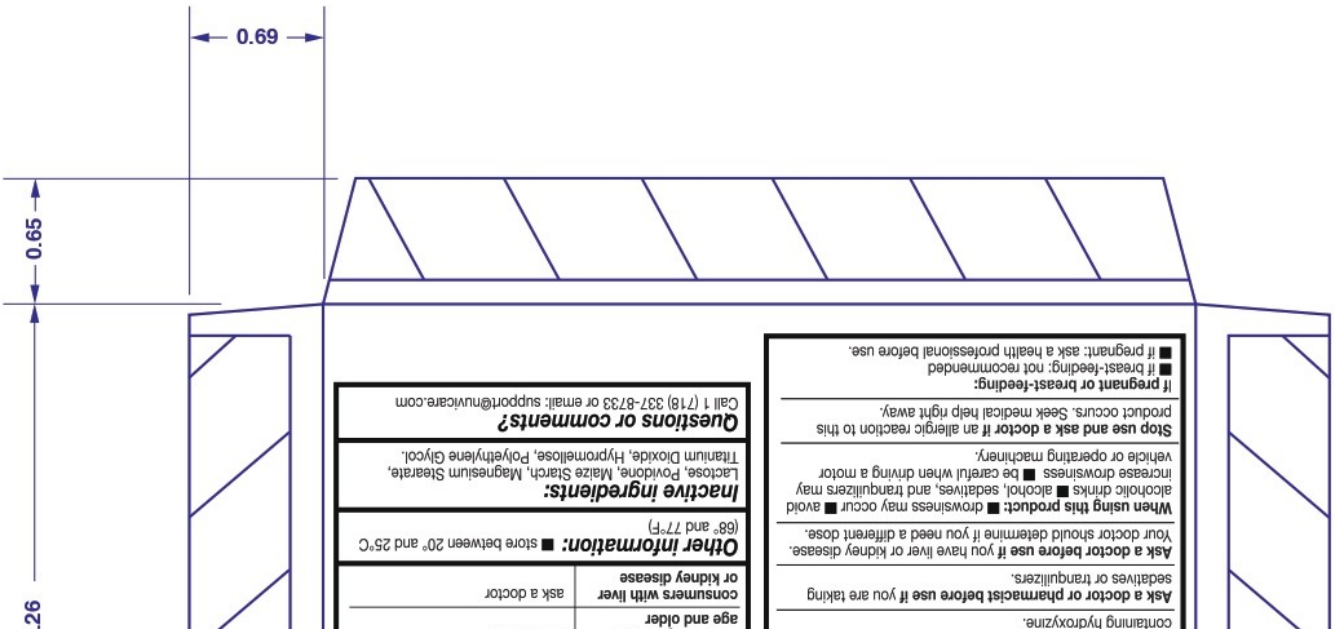
- Store between 20°Cto 25°C(68°F to 77°F)

Inactive Ingredients

Lactose, Povidone, Maize Starch, Magnesium Stearate, Titanium Dioxide, Hypromellose, Polyethylene Glycol

Questions or Comments?

Call 1 (718) 337-8733 or visit support@nuvicare.com



ALLERGY RELIEF

cetirizine hcl tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84324-001
Route of Administration	ORAL		

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

UNII:YO7261ME24)	HYDROCHLORIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics

Color	white	**Score**	no score
Shape	BULLET (Barrel shape)	**Size**	9mm
Flavor		**Imprint Code**	CTN;10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84324-001-02	14 in 1 BLISTER PACK; Type 0: Not a Combination Product	06/09/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077829	06/09/2025	

Labeler - NUVICARE LLC (119257565)

Registrant - NUVICARE LLC (119257565)

Revised: 6/2025 NUVICARE LLC