

CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet, film coated
MedVantx, Inc.

Active ingredient (in each tablet)

Cetirizine hydrochloride USP, 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
- 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 right away.

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).

Inactive ingredients

corn starch, lactose monohydrate, povidone, magnesium stearate and opadry white. The components of opadry white are: hydroxypropyl methylcellulose, polyethylene glycol 400, titanium dioxide

Questions or comments?

call toll free 1-800-206-7821

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 16714-271-02

Cetirizine Hydrochloride Tablets, USP 10 mg

6 Years and Older

Allergy

Antihistamine

24 Hour Relief of

- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

NORTHSTAR

100 Tablets

<div style="background-color: #2e7d72; color: white; padding: 5px; text-align: center;">MEDVANTX®</div> <div style="text-align: center; padding: 10px;"> <p>Don't let your medication run out Call us now!</p> <hr style="width: 50%; margin: 0 auto;"/> <p style="font-size: 1.2em; font-weight: bold;">888.825.8474</p> <p style="font-weight: bold;">medvantx.com</p> </div>	<div style="text-align: center; padding: 10px;"> <p>Dosage: See package insert</p> <p>Take ___ tab(s) ___ time(s) daily or every ___ hours for ___ days.</p> <p>Exp Date: 00/00 Lot: 00000000 66116-417-30 SA-900991</p> <p>Mfg. For: Northstar Rx LLC, Memphis, TN 38141</p> </div>	<div style="background-color: #2e7d72; color: white; padding: 5px; text-align: center;">MEDVANTX®</div> <div style="text-align: center; padding: 10px;"> <p>Cetirizine Hydrochloride Tablets, 10mg</p> <p>30 Tablets</p> <p>Store at controlled room temperature 20°-25°C (68°-77°F)</p> <p style="background-color: black; color: white; border-radius: 50%; width: 20px; height: 10px; margin: 0 auto; display: inline-block;"></p> <p>Sample Only — Not for Sale</p> <p>Keep container tightly closed Keep out of reach of children</p> </div>
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CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66 116-417(NDC:16 714-271)
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
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
POVIDONE K29/32 (UNII: 390RMW2PEQ)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	WHITE	Score	no score
Shape	CAPSULE	Size	9mm
Flavor		Imprint Code	S;521
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66 116-417-30	30 in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078862	10/19/2009	

Labeler - MedVantx, Inc. (806427725)

Registrant - Northstar Rx LLC (830546433)

Establishment

Name	Address	ID/FEI	Business Operations
Blenheim Pharmacal, Inc.		171434587	REPACK(66 116-417)

