

CETIRIZINE HYDROCHLORIDE 10 MG- cetirizine hydrochloride tablet
YYBA CORP

YYBA (as PLD) - WELMATE - CETIRIZINE HYDROCHLORIDE TABLETS, 10 MG
(73581-202)

Active ingredient (in each tablet)

Cetirizine Hydrochloride 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 hydroxyine.

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 (1-800-222-1222)

Directions

| | |
|---|--|
| adults and children 6 years and over | take one 10 mg tablet oncedaily; do not take more than one 10mg tablet in 24 hours. A 5mg product may be appropriate for less severe symptoms. |
| adults 65 years and older | 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° and 25°C (68° and 77°F)

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide

Questions?

call toll-free 1-866-933-6337



Compare to the
active ingredient
of Zyrtec®*

NDC 73581-202-05

Cetirizine Hydrochloride Tablets

For indoor &
outdoor allergies

10 MG | ORIGINAL PRESCRIPTION
STRENGTH

24 hour relief of:

Sneezing
Runny nose
Itchy, watery eyes
Itchy nose or throat

VALUE SIZE!

500 TABLETS

DO NOT USE IF PRINTED SEAL UNDER CAP IS MISSING OR DAMAGED

Drug Facts

Active ingredient

(in each tablet)

Cetirizine hydrochloride 10 mg.....Antihistamine

Purpose

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.

➡ PEEL FOR DIRECTIONS 67032-500-103-0

LIFT HERE



Why pay more?
wellspringmeds.com
866-933-6337

Distributed by:
Wellspring
Airmont, NY 10952, U.S.A.

*This product is not manufactured or distributed by the owner of the registered trademark Zyrtec®

Questions? call toll free 1-866-933-6337

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide.

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

Directions

| | |
|--|--------------|
| 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 |

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.

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]

If pregnant or breast-feeding:

- if breast-feeding: not recommended
- if pregnant: ask a health professional before use

Drug Facts, cont.

CETIRIZINE HYDROCHLORIDE 10 MG

cetirizine hydrochloride tablet

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:73581-202 |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|----------|
| CETIRIZINE HYDROCHLORIDE (UNII: 64O047KTOA) (CETIRIZINE - UNII:Y07261ME24) | CETIRIZINE | 10 mg |

Inactive Ingredients

| Ingredient Name | Strength |
|-----------------|----------|
| | |

| | |
|---|--|
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | |
| CROSCARMELOSE SODIUM (UNII: M28OL1HH48) | |
| HYPROMELLOSE 2208 (100 MPA.S) (UNII: B1QE5P712K) | |
| LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X) | |
| MAGNESIUM STEARATE (UNII: 70097M6I30) | |
| MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) | |
| POLYETHYLENE GLYCOL 1000 (UNII: U076Q6Q621) | |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) | |

Product Characteristics

| | | | |
|-----------------|-----------|---------------------|----------|
| Color | white | Score | 2 pieces |
| Shape | RECTANGLE | Size | 10mm |
| Flavor | | Imprint Code | G4 |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:73581-202-01 | 100 in 1 BOTTLE; Type 0: Not a Combination Product | 06/16/2021 | |
| 2 | NDC:73581-202-05 | 500 in 1 BOTTLE; Type 0: Not a Combination Product | 06/17/2021 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| ANDA | ANDA209274 | 06/16/2021 | |

Labeler - YYBA CORP (006339772)

Revised: 1/2024

YYBA CORP