TOPCARE ALL DAY ALLERGY- cetirizine hydrochloride tablet
Topco Associates LLC

Topco Associates LLC. All Day Allergy 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
- 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	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

- do not use if blister unit is broken or torn (Blister Only)
- do not use if printed foil under cap is broken or missing (Bottle Only)
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

corn starch, FD&C blue no. 1 aluminum lake, hypromellose, lactose monohydrate, magnesium stearate, polydextrose, polyethylene glycol, povidone, titanium dioxide, triacetin

Questions or comments?

1-888-423-0139

Principal Display Panel

ORIGINAL PRESCRIPTION STRENGTH

All Day Allergy

CETIRIZINE HYDROCHLORIDE TABLETS, 10 mg

ANTIHISTAMINE

24 Hour Relief of:

Sneezing

Runny Nose

Itchy, Watery Eyes

Itchy Throat or Nose

Indoor & Outdoor Allergies

actual size

COMPARE TO ZYRTEC® active ingredient

70 TABLETS

TopCare®

ORIGINAL PRESCRIPTION STRENGTH

All Day Allergy

CETIRIZINE HYDROCHLORIDE TABLETS, 10 mg

ANTIHISTAMINE

TopCare®

NDC 36800-458-06

ORIGINAL PRESCRIPTION STRENGTH

All Day Allergy

CETIRIZINE HYDROCHLORIDE TABLETS, 10 mg

ANTIHISTAMINE

24 Hour Relief of:

- **Sneezing**
- **Runny Nose**
- **Itchy, Watery Eyes**
- **Itchy Throat or Nose**

Indoor & Outdoor Allergies

actual size

QUALITY GUARANTEED

*COMPARE TO ZYRTEC® active ingredient**

70 TABLETS

Drug Facts

Active ingredient (in each tablet)	Purpose
Cetirizine HCl 10 mg	Antihistamine

Uses

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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 ▶

LOT NO.

EXP.

: 4H22L 88 C1

Drug Facts (continued)

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)

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Drug Facts (continued)

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Questions or comments?
1-888-423-0139

DISTRIBUTED BY
TOPCO ASSOCIATES LLC
ELK GROVE VILLAGE, IL 60007
1-888-423-0139 ©TOPCO PER913
topcare@topco.com

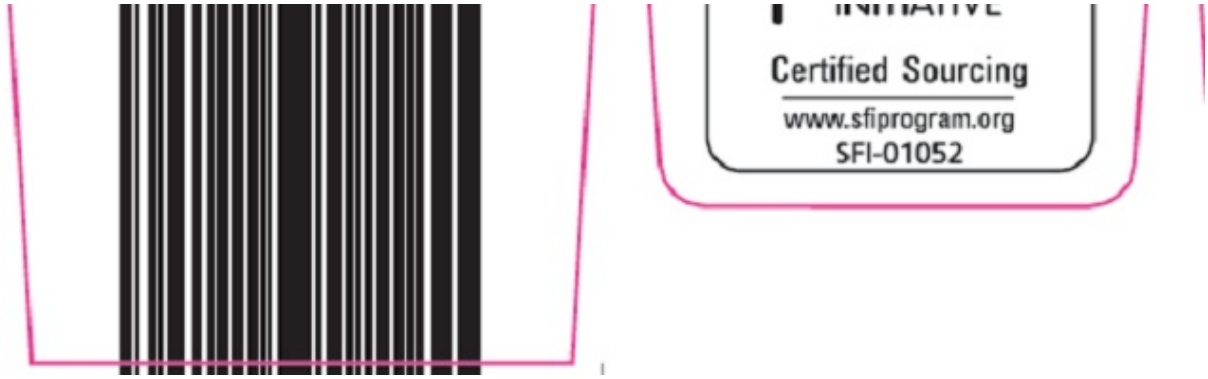
QUALITY GUARANTEED

This TOP CARE® product is laboratory tested to guarantee its highest quality. Your total satisfaction is guaranteed.

GLUTEN FREE

*This product is not manufactured or distributed by McNeil Consumer Healthcare, division of McNeil-PPC, Inc., distributor of Zyrtec®.

SUSTAINABLE FORESTRY INITIATIVE



TOPCARE ALL DAY ALLERGY

cetirizine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:36800-458
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)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POVIDONES (UNII: FZ989GH94E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	

Product Characteristics			
Color	WHITE	**Score**	no score
Shape	OVAL	**Size**	10mm
Flavor		**Imprint Code**	4H2
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:36800-458-13	5 in 1 CARTON	02/12/2008	

1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:36800-458-66	14 in 1 CARTON	01/10/2008	
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:36800-458-39	1 in 1 PACKAGE	04/16/2009	
3		30 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:36800-458-95	1 in 1 PACKAGE	09/08/2009	
4		45 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:36800-458-72	1 in 1 CARTON	01/28/2008	07/05/2015
5		60 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:36800-458-75	1 in 1 CARTON	09/19/2008	06/01/2015
6		90 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:36800-458-47	1 in 1 CARTON	05/26/2010	
7		150 in 1 BOTTLE; Type 0: Not a Combination Product		
8	NDC:36800-458-87	1 in 1 CARTON	04/28/2010	
8		300 in 1 BOTTLE; Type 0: Not a Combination Product		
9	NDC:36800-458-06	1 in 1 CARTON	03/20/2014	
9		70 in 1 BOTTLE; Type 0: Not a Combination Product		
10	NDC:36800-458-58	1 in 1 CARTON	02/05/2014	
10		40 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078336	01/10/2008	

Labeler - Topco Associates LLC (006935977)

Revised: 3/2017 Topco Associates LLC