TOPCARE ALL DAY ALLERGY D- cetirizine hydrochloride, pseudoephedrine hydrochloride tablet, film coated, extended release Topco Associates LLC

Topco Associates LLC. All Day Allergy-D Drug Facts

Active ingredients (in each extended release tablet)

Cetirizine HCl 5 mg

Pseudoephedrine HCl 120 mg

Purpose

Antihistamine

Nasal decongestant

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
- nasal congestion
- reduces swelling of nasal passages
- temporarily relieves sinus congestion and pressure
- temporarily restores freer breathing through the nose

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.
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- heart disease
- thyroid disease

- diabetes
- glaucoma
- high blood pressure
- trouble urinating due to an enlarged prostate gland
- 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

- · do not use more than directed
- 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.
- · you get nervous, dizzy, or sleepless
- symptoms do not improve within 7 days or are accompanied by fever

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

do not break or chew tablet; swallow tablet whole

adults and children 12 years and over	take 1 tablet every 12 hours; do not take more than 2 tablets in 24 hours.
adults 65 years and over	ask a doctor
children under 12 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)
- do not use if blister unit is broken or torn
- see side panel for lot number and expiration date
- meets USP Dissolution Test 2

Inactive ingredients

colloidal silicon dioxide, hypromellose, lactose monohydrate, low-substituted hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, talc, titanium dioxide

Questions or comments?

1-888-423-0139

Package/Label Principal Display Panel

TopCare® health

COMPARE TO ZYRTEC-D® ACTIVE INGREDIENTS

ORIGINAL PRESCRIPTION STRENGTH

ALLERGY & CONGESTION

All Day Allergy - D

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE

HYDROCHLORIDE EXTENDED RELEASE TABLETS, 5 mg/120 mg

ANTIHISTAMINE/NASAL DECONGESTANT

INDOOR & OUTDOOR ALLERGIES

12 HOUR

RELIEF OF:

- Sneezing
- Itchy, Watery Eyes
- Runny Nose
- Itchy Throat or Nose
- Sinus Pressure
- Nasal Congestion

24 EXTENDED RELEASE TABLETS

actual size

- See side panel for lot number and expiration date mæts USP Diss d ution Test S
 - don otu se if blisteru mitis brok en or torn
 - ■store between 20° to 25°C (68° to 77°F) notre mo ini 1911 10

Daung Facts (continued)





- liver or kidneydisease. Your doctor should determine if you need a bns lg dist sond begris ins of sub gnitisninu e iduoti 🔳 арапсоша
 - enuesenq boold rigiri∎ Ask a doctorh efore u se if youh ave ■ heart disease ■ thyroid disease ■ diabetes

the MAOI drug, if you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product. conditions, or Parkins on sidis ease), or for 2 weeks after stopping (MAM) (certain drugs for depression, psychiatric, or emotional

- ify ou are now taking a prescription monoamine oxidase inhibitor ingredients or to an antihistamine containing hydroxy zine.
- ifyou hav eev erhad an all ergic reaction tothis productoranny of its 9 su 1 ou o U

Warnings

- temporarily restoresfreer breathing through the nose ■ temporarilyrelieves sinus congestion and pressure
 - reduces swelling of nasal passages
- ithing of the nose or throat næal congestion ■itdny, wateryeyes Buizaans soon ynnun upper respiratory all ergies:
- temporani y relieves fnese symptoms due to hayfever or other ജ്വ

nimstärlfin A......pm 72 DH anitzititsÖ mersegnoseb lazsM......pm 02 F DH anitbahqaobuse9

(fəl dat əsaələr bəb nə ixə dəsə rii)

Active ingredients

Bring Facts

or kidney disease ask a doctor consumers with liver 12 years ofage children under aska doctor ask a doctor adults 65 years and over take 1 tablet every1 Zhours;do not take more than 2 tablets in 24 hours. 12 years and over adults and children

■ do not break or chew tablet swallow tablet whole Directions

Keep out of reach of children, In case of over dose, get medical help or contact a Poison Control Center inght away. (1-80 0-222-1 222) ■ if pregnant ask a health professional before use. ■ if breast-feeding: not recommended : gn ibə əf-tasəxi no in an gərq fi

eso dun_d

- you getn ervous ,diz zy, or sleepless you getn ervous, din prove within 7 days or are accompanied
- an aller gic reaction to this product occurs. Seek medical help Stop use a nd aska do ctor if
- be careful when driving a motor vehicle or operating machinery drowsi ness may occur
 avoid alcoholic drinke
 alcohol, sedat ves , and tranguil izers may in crease drows iness

When using this product ■ don ot use more than directed aevits be a ro and siliupnant

Ask a doctor or pharma cist before use if you are taking

Drug Facts (continued)



This TopCare® product is laboratory tested to guarantee its highest qualit

uarantee its highest quality satisfaction is guaranteed.

Scan here for more information or call 1-888-423-0139

GF

GLUTEN FREE

Your total s



*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., distributor of Zyrtec-D®.

36800-744-62



ALLERGY & CONGESTION

I Day Allergy-D

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE EXTENDED RELEASE TABLETS, 5 mg/120 mg ANTIHISTAMINE/NASAL DECONGESTANT

> COMPARE TO ZYRTEC-D® ACTIVE INGREDIENTS*



ORIGINAL PRESCRIPTION STRENGTH

ALLERGY & CONGESTION

II Day Allergy-D

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE EXTENDED RELEASE TABLETS, 5 mg/120 mg ANTIHISTAMINE/NASAL DECONGESTANT

INDOOR & OUTDOOR ALLERGIES



- · Itchy, Watery Eyes
- · Itchy Throat or Nose Runny Nose Sinus Pressure
 Nasal Congestion

EXTENDED RELEASE TABLETS

actual size







TOPCARE ALL DAY ALLERGY D

cetirizine hydrochloride, pseudoephedrine hydrochloride tablet, film coated, extended release

Product	Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:36800-744

Route of Administration ORAL

	Active Ingredient/Active Moiety				
ı	Ingredient Name	Basis of Strength	Strength		
	CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	5 mg		
		PSEUDOEPHEDRINE HYDROCHLORIDE	120 mg		

Inactive Ingredients		
Ingredient Name	Strength	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)		
LOW-SUBSTITUTED HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 2165RE0K14)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)		
TALC (UNII: 7SEV7J4R1U)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

Product Characteristics			
Color	WHITE	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	L147
Contains			

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:36800-744- 62	24 in 1 CARTON	03/27/2020		
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210719	03/27/2020	

Labeler - Topco Associates LLC (006935977)

Revised: 2/2022 Topco Associates LLC