CAREONE ALLERGY RELIEF D- cetirizine hcl, pseudoephedrine hcl tablet, extended release American Sales Company

American Sales Company Allergy Relief-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.

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	ask a doctor
disease	

Other information

• store between 20° to 25°C (68° to 77°F)

Inactive ingredients

colloidal silicon dioxide, hydroxypropyl cellulose, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, talc, yellow iron oxide

Questions or comments?

1-800-719-9260

Principal Display Panel

Compare to the active ingredients in Zyrtec-D® ORIGINAL PRESCRIPTION STRENGTH 12hr Allergy Relief-D Cetirizine Hydrochloride and Pseudoephedrine Hydrochloride Extended Release Tablets, 5 mg/120 mg Antihistamine/Nasal Decongestant Actual Size 24 EXTENDED RELEASE TABLETS Allergy & Congestion INDOOR & OUTDOOR ALLERGIES OUR PHARMACISTS RECOMMEND 12-HOUR RELIEF OF: Sneezing – Runny Nose – Sinus Pressure Itchy, Watery Eyes – Itchy Throat or Nose – Nasal Congestion Gluten Free

	[↑] £98L£0Z5L7 £	Questions or comments? 1-800-719-9260 Made in India DisTRIBUTED BY FO ODHO LD U.S.A., LLC EXODVER, MD 20785-1-877-846-9949 G2017 54 5 Bands, LLC G uality guaranteed or your money back.	
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DO NOT USE IF BLISTER UNIT IS BROKEN OR TORN "This product is not manufactured or distributed by McNeil Consumer Healthcare, division of McNeil-PPC, Inc., distributor of Zyrtec-D [®] .	Drug Facts (continued)	Dong Facts (continued) Warnings Donotuse If you are even had an allergic reaction tothis myou are even had an allergic reaction tothis myou are one had an allergic reaction tothis myou are one toto rany of its ingredents or to an antihitatamine if you are now taking a prescription containing hydroxyzine. Mod taug stor totase inhibor (MAC) (ertain drugs for mod depression, psychiatric, or remodioral conditions, or depression, psychiatric, or remodioral conditions depression, psychiatric, or remodioral conditional first or tothor and aff erent dose. Misen using this product drowsiness moreth an directed drowsiness moreth and tranquilizers may increase absonol, sadatives, and tranquilizers may increase absonol, sadatives, drowsiness absonol, sadatives. Towsiness absonol, sadatives, absonol, sadatives, absonol, sadatives, drowsiness absonol, sadatives, absonol, sadatives, abs	: 17662 OF C1
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		NDC 41520-091-62 Compare to the active ingredients in Zyrtec-D®* AL PRESCRIPTION STRENGTH STRENGT-D	
	Cetin Pseudoe Extended Relea Anti Actual Size EXTENDED		



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CAREONE	ALLERGY R	FI IFF D				
		tablet, extended release				
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Product Info	mation					
Product T ype		HUMAN OTC DRUG	Item Code (Source)		NDC:41520-091	
Route of Admin	stration	ORAL				
Active Ingred	ient/Active Moi	5				Strength
	Ingr	edient Name		Basis of Str	Basis of Strength	
CETIRIZINE HYDRO CHLO RIDE (UNII: 640047KTOA) (CETIRIZINE - UNII: Y07261ME24)			CETIRIZINE HYDROCHLORIDE		5 mg	
PSEUDO EPHEDR - UNII:7CUC9DDI		IDE (UNII: 6 V9 V2RYJ8 N) (PSEUD	OEPHEDRINE	PSEUDOEPHEDRIN HYDROCHLORIDE	Е	120 mg
Inactive Ingr	dianta					
macuve mgro	Inactive Ingredients Ingredient Name			Star		Strength
SILICON DIO XII	E (UNII: ETJ7Z6 XBU	-				Strength
		500000 WAMW) (UNII: RFW2ET6	71P)			
	ES (UNII: 3NXW29V3	, .	/ 11)			
	DHYDRATE (UNII: EV					
	E ARATE (UNII: 7009					
		(UNII: OP1R32D61U)				
TALC (UNII: 7SEV						
	ELLOW (UNII: EX43	802MRT)				
	× ·	,				
Product Char	acteristics					
Color	Color WHITE (one side white one side light yellow)		Score no		core	
Shape	hape ROUND		Size 12mm		n	
Flavor	Flavor		Imprint Code 5029;		;5;120	
Contains						
Packaging						
# Item Code		Package Description	Ma	keting Start Date	Marketin	ig End Date
1 NDC:41520-091	-62 24 in 1 CARTON		07/20/2016			
1	1 in 1 BLISTER I	PACK; Type 0: Not a Combination	Product			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
ANDA	ANDA077170	07/20/2016				

Labeler - American Sales Company (809183973)

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American Sales Company