GOOD NEIGHBOR PHARMACY ALL DAY ALLERGY D- cetirizine hydrochloride, pseudoephedrine hydrochloride tablet, film coated, extended release Amerisource Bergen

Amerisource Bergen 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?

1-800-719-9260

Package/Label Principal Display Panel

Compare to $Zyrtec-D^{\mathbb{R}}$ active ingredients Original Prescription Strength

All Day Allergy-D

ALLERGY & CONGESTION

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

actual size

24 Extended Release Tablets



Original Prescription Strength

ALLERGY & CONGESTION

Day Allergy-D

cetirizine hydrochloride and pseudoephedrine hydrochloride extended rélease tablets, 5 mg/120 mg antihistamine/nasal decongestant



Compare to Zyrtec-D° active ingredients'

NDC 46122-626-62

CONGESTION

ALLERGY &

Original Prescription Strength

All Day Allergy-I

cetirizine hydrochloride and pseudoephedrine hydrochloride extended release tablets, 5 mg/120 mg antihistamine/nasal decongestant

Indoor & Outdoor Allergies

12 Hour Relief of:

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Good Neighbor Pharmacy® All Day Allergy-D is not manufactured or distributed by Johnson & Johnson Consumer Inc., distributor of Zyrtec-D®

GLUTEN FREE

Distributed By AmerisourceBergen 1300 Morris Drive Chesterbrook, PA19087 Questions or Concerns? www.mygnp.com



Drug Facts (continued)

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Inactive in gred ients colloidal silicon dioxide, hypromellose, lactose m on o hydrate, low-subs tituted hydroxypropyl cellul ose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvin ylalcohol, talc, titanium dioxide

Questions or comments? 1 -800-7 19-92 60

AB C#:1 02 2965 0



GOOD NEIGHBOR PHARMACY ALL DAY ALLERGY D

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

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46122-626
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
CETIRIZINE HYDRO CHLO RIDE (UNII: 640047KTOA) (CETIRIZINE - UNII: YO7261ME24)	CETIRIZINE HYDROCHLORIDE	5 mg		
PSEUDO EPHEDRINE HYDRO CHLO RIDE (UNII: 6 V9 V2RYJ8 N) (PSEUDO EPHEDRINE - UNII:7CUC9 DDI9 F)	PSEUDOEPHEDRINE HYDROCHLORIDE	120 mg		

Inactive Ingredients		
Ingredient Name	Strength	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29 V3WO)		
LACTO SE MONO HYDRATE (UNII: EWQ57Q8I5X)		
LOW-SUBSTITUTED HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 2165RE0K14)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)		
TALC (UNII: 7SEV7J4R1U)		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)		

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

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-626-62	24 in 1 CARTON	03/30/2020	
1		$1\ \text{in}\ 1\ \text{BLISTER}$ PACK; Type $\ 0\ :$ Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210719	03/30/2020	

Labeler - Amerisource Bergen (007914906)

Revised: 7/2020 Amerisource Bergen