GOOD SENSE ALL DAY ALLERGY D- cetirizine hydrochloride, pseudoephedrine hydrochloride tablet, film coated, extended release L. Perrigo Company

Perrigo 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

Allergy & Congestion

Original Prescription Strength

All Day Allergy-D

Actual Size

Cetirizine Hydrochloride and Pseudoephedrine Hydrochloride

Extended Release Tablets, 5 mg/120 mg

Antihistamine/Nasal Decongestant

12 Hour Relief of:

Indoor & Outdoor Allergies

Sneezing

Itchy, Watery Eyes

Runny Nose

Itchy Throat or Nose

Sinus Pressure

Nasal Congestion

Compare to active ingredients of Zyrtec-D®

100% SATISFACTION GUARANTEED

12 Extended Release Tablets



Purp ose

OPEN OTHER END

4753 C2 C2

Drug Facts

Active ingredients

(in each extended release tablet)

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Ask adoctor before use if you have

- heart disease thyroid disease diabetes

- glaucoma high blood pressure trouble urinating due to an enlarged prostate gland liveror kidney disease. Your doctor should determine if you need a different dose.

Drug Facts (continued)

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When using this product ■donotuse more than directed

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- right away. you get nervous, dizzy, ors leepless
- symptoms don ot improve within 7 days or are accompanied

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SONVENIENT RECLOSING TAB

Drug Facts (continued)

Other information

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 donotuse if blister unit is broken ortorn
 see side panel for lot numberand expiration date meets USP *Dissolut on Test 2*

Inactive in gred ients colloidal silicon dioxide, hypromellose, lactose monohydrate, low-subs futed hydroxypropyl cellul ose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvin yl alcohol, talc, titanium dioxide

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



Gluten Free

GOOD SENSE ALL DAY ALLERGY D

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

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0113-0147
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)		
LACTOSE MONOHYDRATE (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:0113-0147-53	12 in 1 CARTON	01/29/2021	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0113-0147-62	24 in 1 CARTON	01/27/2021	
2		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 12/27/2020

Labeler - L. Perrigo Company (006013346)

Revised: 1/2021 L. Perrigo Company