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?

call **1-800-719-9260**

Package/Label Principal Display Panel

Compare to Zyrtec-D® active ingredients

ORIGINAL PRESCRIPTION STRENGTH

ALL DAY allergy-D

Cetirizine Hydrochloride and Pseudoephedrine Hydrochloride Extended Release tablets, 5 mg/120 mg

Antihistamine/Nasal Decongestant

ALLERGY + SINUS

INDOOR + OUTDOOR ALLERGIES

12 hour

12 HOUR RELIEF

Sneezing

Itchy, Watery Eyes

Runny Nose

Itchy Nose or Throat

Actual Size

NASAL CONGESTION + SINUS PRESSURE

24 EXTENDED RELEASE TABLETS


CONVENIENT RECLOSING TAB
Distributed By Perrigo® Allegan, MI 49010 perrigo.com
OPEN OTHER END
14762 55 C3
Important: Read all product information before using. Keep this box for important information.
Drug Facts
Active ingredients (in each extended release tablet)
Purpose
Cetirizine HCl 5 mg
Antihistamine
Pseudoephedrine HCl 120 mg
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.
Drug Facts (continued)
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 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
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Drug Facts (continued)
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? call 1-800-719-9260
3 70030 16513 1
ORIGINAL PRESCRIPTION STRENGTH
ALL DAY allergy-D
Compare to Zyrtec-D® active ingredients
NDC 0113-2147-62
ORIGINAL PRESCRIPTION STRENGTH
ALL DAY allergy-D
Cetirizine Hydrochloride and Pseudoephedrine Hydrochloride
Extended Release Tablets, 5 mg/120 mg
Antihistamine/Nasal Decongestant
ALLERGY + SINUS
INDOOR + OUTDOOR ALLERGIES
12 hour
12 HOUR RELIEF
Sneezing
Runny Nose
Itchy, Watery Eyes
Itchy Nose or Throat
NASAL CONGESTION + SINUS PRESSURE
Actual Size
24 EXTENDED RELEASE TABLETS

ALL DAY ALLERGY D

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

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0113-2147
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CETIRIZINE HYDROCHLORIDE (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	5 mg
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	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: 3WQ0SDW1A)	
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:0113-2147-62	24 in 1 CARTON	10/02/2025	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
ANDA	ANDA210719	10/02/2025	

Labeler - L. Perrigo Company (006013346)

Revised: 10/2025 L. Perrigo Company