PENTREXCILINA DAY TIME- cetirizine hydrochloride tablet OPMX LLC

Pentrexcilina Day time

Drug Facts

Drug Facts

Active Ingredient

Active Ingredient (in each tablet)	Purpose
Cetirizine HCl 10 mg	Antihistamine

Purpose

Antihistamine

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

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

Ask a doctor before use if you have

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 sedatives or tranquilizers.

When using this product

- 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.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

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OVERDOSE WARNING:

In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as children even if you do not notice any signs or symptoms.

Directions

Adults & Children 6 years and over One 10mg tablet once daily; do not take more than one 10mg tablet in 24 hours. A 5 mg product may be appropriate for less severe symptoms

Adults 65 years and over ask a doctor

Children under 6 years ask a doctor

Consumers with liver or kidney disease ask a doctor

Other information

tore at room temperature between 20-25°C (68-77°F).

close cap tightly after use.

contains no ingredients made from a gluten-containing grain (wheat, barley or rye).

TAMPER EVIDENT: Do not use if imprinted safety seal is broken or missing

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide.

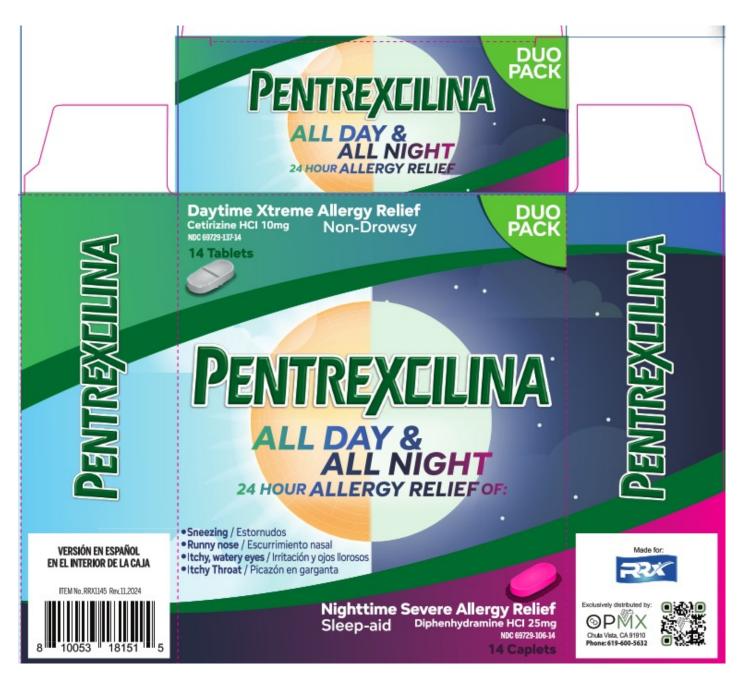
Questions or comments?

Call (619) 600-5632 Monday through Friday 9 AM- 5 PM EST

ALL DAY

Drug Facts

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Active Ingredient (in each tablet) Purpos Cetirizine HCl 10 mgAntihistami	_
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	r.
Warnings Do not use if you have ever had an allergic reaction to this produ or any of its ingredients or to an antihistamine containing hydroxyzine.	ct
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If pregnant or breast-feeding, ask a health professional before use.	-
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Daytime Xtreme Allergy Relief

Cetirizine HCI 10mg

Non-Drowsy

NDC 69729-137-14

14 Tablets

DUO PACK

PENTREXCILINA

ALL DAY & ALL NIGHT

24 HOUR ALLERGY RELIEF OF:

- Sneezing/ Estornudos
- Runny nose/ Escurrimiento nasal
- Itchy, watery eyes/ Irritación y ojos llorosos

• Itchy throat/ Picazón en garganta

Nighttime Severe Allergy Relief

Sleep-aid

Diphenhydramine HCl 25mg

NDC 69729-106-14

14 Caplets

Made for RRX

Exclusively distributed by:

OPMX

Chula Vista CA 91910

Phone: 619-600-5632

PENTREXCILINA DAY							
cetirizine hydrochloride tablet							
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Product Information							
Product Type	HUMAN OTC DRUG		Item Code (Source) ND			DC:69729-137	
Route of Administration	ORAL						
Active Ingredient/Active	Moiety						
Ingre	dient Name			Basis of St	rength	Strength	
CETIRIZINE HYDROCHLORIDE (U UNII:YO7261ME24)	TIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE - CETIRIZINE				E	10 mg	
Inactive Ingredients							
	Ingredient N	lame			St	Strength	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)							
POLYETHYLENE GLYCOL, UNSPE	CIFIED (UNII: 3WJQ	0SDW1A	A)				
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)							
CROSCARMELLOSE SODIUM (UN							
HYPROMELLOSE, UNSPECIFIED)					
TITANIUM DIOXIDE (UNII: 15FIX9V	•						
SILICON DIOXIDE (UNII: ETJ7Z6XB							
MAGNESIUM STEARATE (UNII: 700	097M6I30)						
Product Characteristics							
Color white	2	Score			2 pieces		
Shape CAPS	ULE	Size			9mm		
Flavor		Imprin	t Code		G;4		

Co	ontains			
n -				
۲a	ackaging			
#	ltem Code	CodePackage DescriptionMarketing StartModelDate		Marketing End Date
	NDC:69729-137- 14 in 1 BOTTLE; Type 0: Not a Combination Product		11/15/2024	
Л	arkating	nformation		
	arketing			
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
AN	DA	ANDA209274	11/15/2024	

Labeler - OPMX LLC (029918743)

Establishment								
Name	Address	ID/FEI	Business Operations					
Granules India Limited		918609236	manufacture(69729-137)					

Revised: 11/2024

OPMX LLC