ALL DAY ALLERGY-D- cetirizine hydrochloride and pseudoephedrine hydrochloride tablet
Strategic Sourcing Services

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all day allergy-D

Drug Facts

Active ingredients (in each extended-release tablet)	***Purpose***
Cetirizine HCl, USP 5 mg	Antihistamine
Pseudoephedrine HCl, USP 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 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) (cer tain 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 at **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 carton is opened or if the blister unit is broken
- see side panel for batch number and expiration date

Inactive ingredients

hydroxyethyl cellulose, hydroxypropyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, stearic acid, titanium dioxide

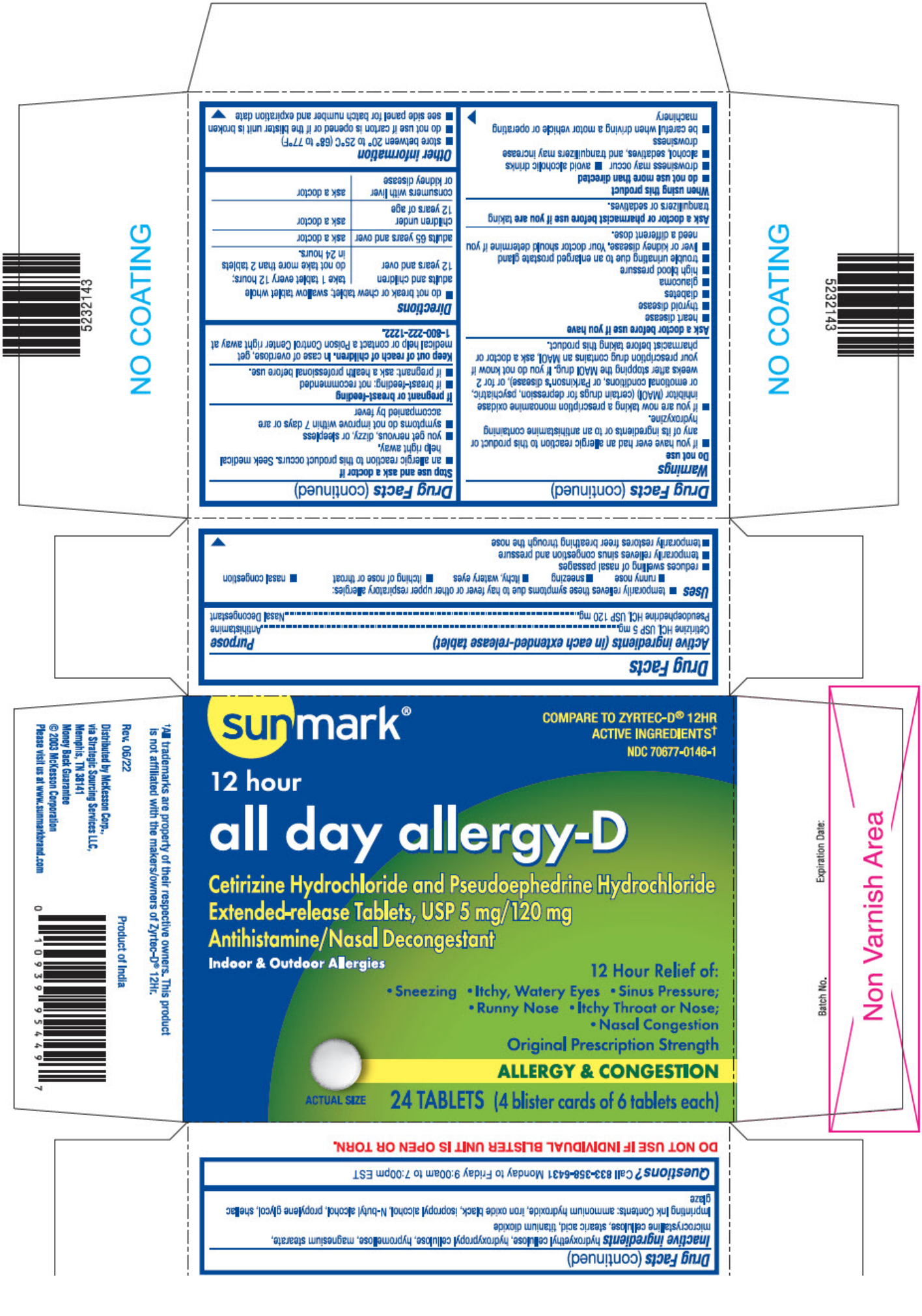
Imprinting Ink Contents: ammonium hydroxide, iron oxide black, isopropyl alcohol, N-butyl alcohol, propylene glycol, shellac glaze

Questions?

Call **833-358-6431** Monday to Friday 9:00am to 7:00pm EST

Distributed by McKesson Corp.,
via Strategic Sourcing Services LLC,
Memphis, TN 38141

PRINCIPAL DISPLAY PANEL - 24 Tablet Blister Card Carton


NO COATING
5232143
Drug Facts (continued)
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Imprinting Ink Contents: ammonium hydroxide, iron oxide black, isopropyl alcohol, N-butyl alcohol, propylene glycol, shellac glaze
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DO NOT USE IF INDIVIDUAL BLISTER UNIT IS OPEN OR TORN.
sunmark®
12 hour
all day allergy-D
Cetirizine Hydrochloride and Pseudoephedrine Hydrochloride Extended-release Tablets, USP 5 mg/120 mg
Antihistamine/Nasal Decongestant
Indoor & Outdoor Allergies
COMPARE TO ZYRTEC-D® 12HR ACTIVE INGREDIENTS†
NDC 70677-0146-1
12 Hour Relief of:
• Sneezing • Itchy, Watery Eyes • Sinus Pressure;
• Runny Nose • Itchy Throat or Nose;
• Nasal Congestion
Original Prescription Strength
ALLERGY & CONGESTION
ACTUAL SIZE
24 TABLETS (4 blister cards of 6 tablets each)
†All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Zyrtec-D® 12Hr.
Rev. 06/22
Product of India
Distributed by McKesson Corp., via Strategic Sourcing Services LLC, Memphis, TN 38141
Money Back Guarantee
© 2003 McKesson Corporation
Please visit us at www.sunmarkbrand.com
0 10939 95449 7
Expiration Date:
Batch No.
Non Varnish Area
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ALL DAY ALLERGY-D

cetirizine hydrochloride and pseudoephedrine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70677-0146
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
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
AMMONIA (UNII: 5138Q19F1X)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
BUTYL ALCOHOL (UNII: 8PJ61P6TS3)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
HYDROXYETHYL CELLULOSE (4000 MPA.S AT 1%) (UNII: ZYD53NBL45)	

Product Characteristics

Color	WHITE	**Score**	no score
Shape	ROUND (circular)	**Size**	9mm
Flavor		**Imprint Code**	915
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70677-0146-1	4 in 1 CARTON	11/01/2018	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination		

1		Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090922	11/01/2018	

Labeler - Strategic Sourcing Services (116956644)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharmaceutical Industries Limited		650445203	MANUFACTURE(70677-0146)

Revised: 6/2022 Strategic Sourcing Services