ALLERGY RELIEF-D- cetirizine hydrochloride and pseudoephedrine hydrochloride tablet, film coated, extended release
STRATEGIC SOURCING SERVICES LLC

Allergy relief-D

Drug Facts

Active ingredients (in each extended-release tablet)	***Purposes***
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) (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 aretaking 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 9:00 am to 7:00 pm EST

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

PRINCIPAL DISPLAY PANEL - 24 Tablet Blister Card Carton

NDC 70677-1020-1

COMPARE TO ZYRTEC-D ®
12 HR ACTIVE INGREDIENTS*

Foster & Thrive™

ORIGINAL PRESCRIPTION STRENGTH

All Day Allergy-D

Cetirizine Hydrochloride and Pseudoephedrine Hydrochloride Extended-release Tablets, USP 5 mg/120 mg
ANTIHISTAMINE/NASAL DECONGESTANT

Indoor & Outdoor Allergies | Allergy & Sinus

12 hour relief of:

- Sneezing
- Itchy, Watery Eyes
- Sinus Pressure
- Runny Nose
- Itchy Throat or Nose
- Nasal Congestion

ACTUAL SIZE

24 TABLETS (4 blister cards of 6 tablets each)

NDC 70677-1020-1

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ACTUAL SIZE

24 TABLETS (4 blister cards of 6 tablets each)

NON VARNISH
NO INK AREA

*All trademarks are property of their respective owners. This product is not affiliated with the owners/makers of Zyrtec-D®.

Distributed by: McKesson Corp., via Strategic Sourcing Services LLC, Memphis, TN 38141
Money Back Guarantee
www.fosterandthrive.com
PLD-A604A LB00880 Rev 08/24
Product of India
DNH/DRUGS/NH/138

0 10939 95538 8

DO NOT USE IF INDIVIDUAL BLISTER UNIT IS OPEN OR TORN.

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

Inactive ingredients hydroxypropyl cellulose, hypromellose, 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

5254313

4x6 Tablets

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Drug Facts (continued)

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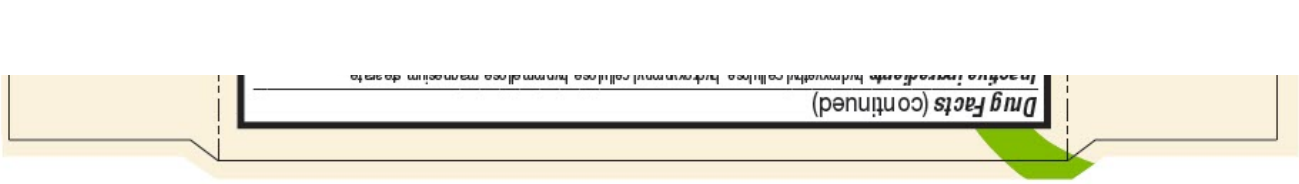
- see side panel for batch number and expiration date

NO COATING

NVZ

NO COATING

NVZ



ALLERGY RELIEF-D

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

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70677-1020
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:70677-1020			

1	NDC:70677-1020-1	4 in 1 CARTON	08/11/2023	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:70677-1020-2	2 in 1 CARTON	10/04/2024	
2		6 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	ANDA090922	08/11/2023	

Labeler - STRATEGIC SOURCING SERVICES LLC (116956644)

Registrant - Sun Pharmaceutical Industries Limited (650172430)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharmaceutical Industries Limited		650445203	manufacture(70677-1020)

Revised: 10/2024 STRATEGIC SOURCING SERVICES LLC