

**FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE-
fexofenadine hydrochloride and pseudoephedrine hydrochloride tablet,
extended release
Strategic Sourcing Services, LLC**

**Fexofenadine Hydrochloride and Pseudoephedrine Hydrochloride Extended-
Release Tablets USP, 60 mg/120 mg**

Active ingredient(s)

Fexofenadine HCl USP, 60 mg

Pseudoephedrine HCl USP, 120 mg

Purpose

Antihistamine

Nasal decongestant

Use(s)

- 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
- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies
- 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
- 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.
- if you have difficulty swallowing

Ask a doctor before use if you have

- heart disease
- thyroid disease

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

When using this product

- **do not take more than directed**
- do not take at the same time as aluminum or magnesium antacids
- do not take with fruit juices (see Directions)
- the tablet coating may be seen in the stool (this is normal). Continue to take as directed (see Directions).

Stop use and ask doctor if

- an allergic reaction to this product occurs. Seek medical help right away.
- symptoms do not improve within 7 days or are accompanied by a fever
- you get nervous, dizzy, or sleepless

If pregnant or breast-feeding

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.

Directions

- do not divide, crush, chew or dissolve the tablet; swallow tablet whole

adults and children 12 years of age and over	take 1 tablet with a glass of water every 12 hours on an empty stomach; do not take more than 2 tablets in 24 hours
children under 12 years of age	do not use
adults 65 years of age and older	ask a doctor
consumers with kidney disease	ask a doctor

Other information

- safety sealed: do not use if carton is opened or if individual blister units are torn or opened
- store between 20° to 25°C (68° to 77°F) store between 20° to 25°C (68° to 77°F)
- FDA approved dissolution test specifications differ from USP.
- FDA approved organic impurities test procedure differs from USP Procedure 1.

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, ferric oxide (iron oxide yellow), HPMC 2910 / hypromellose (6 Cps), hydroxypropyl cellulose, hypromellose (methocel K100M DC2), macrogol (polyethylene glycol MW 400), macrogol (polyethylene glycol MW 8000), magnesium stearate, microcrystalline cellulose (avicel PH 101), microcrystalline cellulose (avicel PH102), pregelatinized starch.

Questions?

Call **1-888-375-3784**

Principal Display Panel

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Product of India
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Questions or comments? Call 888-20-8449 Monday through Friday 9 am to 5:30 pm EST.

Inactive ingredients: Cellulose, croscarmellose sodium, hydroxypropyl methylcellulose, polyethylene glycol, polyethylene glycol 400, polyethylene glycol 600, polyethylene glycol 800, polyethylene glycol 1000, polyethylene glycol 1500, polyethylene glycol 2000, polyethylene glycol 3000, polyethylene glycol 4000, polyethylene glycol 6000, polyethylene glycol 8000, polyethylene glycol 10000, polyethylene glycol 15000, polyethylene glycol 20000, polyethylene glycol 30000, polyethylene glycol 40000, polyethylene glycol 60000, polyethylene glycol 80000, polyethylene glycol 100000, polyethylene glycol 150000, polyethylene glycol 200000, polyethylene glycol 300000, polyethylene glycol 400000, polyethylene glycol 600000, polyethylene glycol 800000, polyethylene glycol 1000000.

Other information: Safety: Read the directions and warnings carefully. Do not use if the seal is broken or if the tablets are discolored. Do not use if the tablets are broken or if the tablets are discolored. Do not use if the tablets are broken or if the tablets are discolored.

Directions: Do not drink alcohol while taking this medicine. Do not drink alcohol while taking this medicine. Do not drink alcohol while taking this medicine.

Drug Facts (continued): When using this product, do not take more than directed. Do not take more than directed. Do not take more than directed.

IMPORTANT: READ THE DIRECTIONS AND WARNINGS BEFORE USE. KEEP THE CAPTION, IT CONTAINS IMPORTANT INFORMATION.

Drug Facts (continued): Ask a doctor before use if you have heart disease, high blood pressure, or if you are taking any other medicine. Ask a doctor before use if you have heart disease, high blood pressure, or if you are taking any other medicine.

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Drug Facts (continued): When using this product, do not take more than directed. Do not take more than directed. Do not take more than directed.

LOT
EXP



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NDC 70577-1010-1

Foster & Thrive™

12 HOUR

Allergy & Congestion-D

Fexofenadine Hydrochloride & Pseudoephedrine Hydrochloride Extended-Release Tablets, USP 60 mg/120 mg ANTIHISTAMINE & NASAL DECONGESTANT

Non-Drowsy INDOOR/OUTDOOR RELIEF OF

- Nasal and Sinus Congestion Due to Colds or Allergies
- Sneezing; Runny Nose; Itchy, Watery Eyes and Itchy Nose or Throat Due to Allergies



ACTUAL SIZE

20 EXTENDED-RELEASE TABLETS

COMPARE TO THE ACTIVE INGREDIENTS IN ALLEGRA-D® 12 HOUR ALLERGY & CONGESTION TABLETS*

Foster & Thrive™

12 HOUR

Allergy & Congestion-D

Fexofenadine Hydrochloride & Pseudoephedrine Hydrochloride Extended-Release Tablets, USP 60 mg/120 mg ANTIHISTAMINE & NASAL DECONGESTANT

FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

fexofenadine hydrochloride and pseudoephedrine hydrochloride tablet, extended release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-1010
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	60 mg
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	120 mg

Inactive Ingredients

Ingredient Name	Strength
MICROCRYSTALLINE CELLULOSE 101 (UNII: 7T9FYH5QMK)	
Croscarmellose Sodium (UNII: M28OL1HH48)	
Ferric Oxide Yellow (UNII: EX438O2MRT)	
Silicon Dioxide (UNII: ETJ7Z6XBU4)	
Magnesium Stearate (UNII: 70097M6I30)	
Polyethylene Glycol 400 (UNII: B697894SGQ)	
Hypromellose 2208 (100000 Mpa.S) (UNII: VM7F0B23ZI)	
HYDROXYPROPYL CELLULOSE (45000 WAMW) (UNII: 8VAB711C5E)	
Hypromellose 2910 (6 Mpa.S) (UNII: 0WZ8WG20P6)	
Polyethylene Glycol 8000 (UNII: Q662QK8M3B)	
Starch, Corn (UNII: O8232NY3SJ)	
MICROCRYSTALLINE CELLULOSE 102 (UNII: PNR0YF693Y)	

Product Characteristics

Color	WHITE (one white to off-white color layer and other light yellow to yellow color)	Score	no score
Shape	CAPSULE	Size	16mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70677-1010-1	4 in 1 CARTON	05/05/2023	
1		5 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	ANDA215434	03/31/2022	

Labeler - Strategic Sourcing Services, LLC (116956644)

Revised: 4/2023

Strategic Sourcing Services, LLC