

FEXOFENADINE HCL AND PSEUDOEPHEDRINE HCl- fexofenadine hcl and pseudoephedrine hci tablet, extended release
Dr. Reddy's Laboratories Inc.

Fexofenadine HCl 60 mg and Pseudoephedrine HCl 120 mg ER Tablets USP

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)
- this product meets the requirements of USP dissolution test 3.

Inactive ingredients

corn starch, croscarmellose sodium, colloidal silicon dioxide, ferric oxide, hypromellose, kollidon SR, magnesium stearate, mannitol, powder cellulose and triethyl citrate.

Questions?

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

Distributed by:

Dr. Reddy's Laboratories, Inc.

Princeton, NJ 08540

Made in India

Principal Display Panel

Blister carton

TAMPERS EVIDENT: NON-DISBURSTERS UNLESS SEAL OR MORE OF DM.
DO NOT USE IF INDIVIDUAL BLISTER SHOWS SIGNS OF TAMPERING.
IMPORTANT: Read the directions and warnings on the top of the carton. Do not take more than the amount

Drug Facts (continued)

Warnings

- Do not use if you have ever had an allergic reaction to this product.
- Do not use if you are taking any other medicine for allergies, such as antihistamines, decongestants, or pseudoephedrine.
- Do not use if you are taking any other medicine for allergies, such as antihistamines, decongestants, or pseudoephedrine.

Directions

Take 1 tablet twice a day with meals. Do not take more than 2 tablets in 24 hours.

Other Information

Keep out of the reach of children. In case of overdose, get medical help or contact Poison Control Center right away.

Drug Facts (continued)

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LOT
EXP



Dr.Reddy's NDC 43535-223-14
Compare to the active ingredients in Allegra-C®
12 Hour Allergy & Congestion Tablets*

Non-Drowsy • Original Prescription Strength
Fexofenadine HCl 60 mg & Pseudoephedrine HCl 120 mg

Extended-Release Tablets USP

ALLERGY & CONGESTION

Fexofenadine HCl 60 mg / antihistamine
Pseudoephedrine HCl 120 mg / nasal decongestant

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



20 Tablets

Dr.Reddy's Non-Drowsy • Original Prescription Strength

Fexofenadine HCl 60 mg & Pseudoephedrine HCl 120 mg

Extended-Release Tablets USP

ALLERGY & CONGESTION

Fexofenadine HCl 60 mg / antihistamine
Pseudoephedrine HCl 120 mg / nasal decongestant

Made in India

*This product is not manufactured or distributed by Chiesi, Inc. (part of the Bristol-Myers Squibb company) or Allegra-C® 12 Hour Allergy & Congestion Tablets. Allegra-C® is a registered trademark of Aventisub LLC.

FEXOFENADINE HCL AND PSEUDOEPHEDRINE HCl

fexofenadine hcl and pseudoephedrine hci tablet, extended release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43598-823
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
MANNITOL (UNII: 3OWL53L36A)	
PO WDERED CELLULOSE (UNII: SMD1X3XO9M)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
SILICON DIO XIDE (UNII: ETJ7Z6XBU4)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	

Product Characteristics

Color	WHITE (buff white to pale yellow color and other layer light red to red color)	Score	no score
Shape	CAPSULE	Size	16mm
Flavor		Imprint Code	R;195
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43598-823-14	4 in 1 CARTON	07/18/2019	
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:43598-823-31	6 in 1 CARTON	07/18/2019	
2		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:43598-823-35	2 in 1 CARTON	10/31/2019	
3		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	ANDA076667	07/18/2019	

Labeler - Dr. Reddy's Laboratories Inc. (802315887)

Revised: 9/2019

Dr. Reddy's Laboratories Inc.