

**FEXOFENADINE HYDROCHLORIDE- fexofenadine hcl tablet, film coated
Proficient Rx LP**

Perrigo Fexofenadine Hydrochloride Tablets, 60 mg Drug Facts

Active ingredient (in each tablet)

Fexofenadine HCl 60 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- itchy, watery eyes
- sneezing
- 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.

Ask a doctor before use if you have

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)

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.

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

adults and children 12 years of age	take one 60 mg tablet with water every 12 hours; do not take
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and over	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

Other information

- do not use if printed foil under cap is broken or missing
- store at 20°-25°C (68°-77°F)
- protect from excessive moisture
- this product meets the requirements of USP Dissolution Test 3

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, titanium dioxide

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

Compare to Allegra® Allergy active ingredient

Fexofenadine Hydrochloride Tablets, 60 mg

Antihistamine

Non-Drowsy

Relief of:

Sneezing

Runny Nose

Itchy, Watery Eyes

Itchy Nose or Throat

Original Prescription Strength

12 Hour

60 mg Each

Actual Size

Indoor & Outdoor Allergies

Allergy

Repackaged by:

Proficient Rx LP

Thousand Oaks, CA 91320



NDC 63187-629-60

Lot #:00000
Exp. 00/00/00
SN# MASTER

Fexofenadine HCl 60mg

#60 TabletsEach tablet contains: Fexofenadine HCl 60mg
Antihistamine*Round, orange / peach colored, unscored tablet debossed with "93" on one side and "7252" on the other side.*

Product ID: PF062960

Dist. By: Perrigo Allegan, MI 49010 Made in Israel

Store at 20°-25°C (68°-77°F)

Keep medication out of the reach of children

Fexofenadine HCl 60mg
#60 Tablets
Lot #:00000 SN# MASTER
NDC 63187-629-60 Exp:00/00/00Fexofenadine HCl 60mg
#60 Tablets
Lot #:00000 SN# MASTER
NDC 63187-629-60 Exp:00/00/00Fexofenadine HCl 60mg
#60 Tablets
Lot #:00000 SN# MASTER
NDC 63187-629-60 Exp:00/00/00Packaged By: Proficient Rx LP
Thousand Oaks, CA 91320

Fexofenadine Hydrochloride Tablets, 60 mg Carton

FEXOFENADINE HYDROCHLORIDE

fexofenadine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63187-629(NDC:45802-425)
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

Inactive Ingredients

Ingredient Name	Strength
SILICON DIO XIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TITANIUM DIO XIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	ORANGE (Peach)	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	93;7252
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63187-629-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2018	
2	NDC:63187-629-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2018	
3	NDC:63187-629-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076447	08/08/2011	

Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(63187-629) , RELABEL(63187-629)

Revised: 11/2019

Proficient Rx LP