FEXOFENADINE HCL- fexofenadine hcl tablet
CVS

Fexofenadine HCL Tablets USP, 180mg

Antihistamine

Indoor/Outdoor Allergy Relief

- **Sneezing**
- **Runny nose**
- **Itchy, watery eyes**
- **Itchy nose or throat**

Non-Drowsy

Active ingredient (in each tablet)

Fexofenadine HCl USP, 180 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

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.

Directions

adults and children 12 years of age and take one 180 mg tablet with water once a day; do not take more than 1 tablet in 24 hours

Other information

■ safety sealed: do not use if imprinted safety seal under cap is broken or missing
■ store between 20° and 25°C (68° and 77°F)
■ protect from excessive moisture

Inactive ingredients

ammonium hydroxide, colloidal silicon dioxide, croscarmellose sodium, D & C Red 33, ferrosoferric oxide, FD & C blue 1, FD & C yellow 6, gelatin, hydroxypropyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, pregelatinized starch, propylene glycol, shellac, titanium dioxide.

Questions or comments?

contact **1-877-770-3183** Mon-Fri 8:00 AM EST to 5:00 PM PST

PDP

FEXOFENADINE HCL

fexofenadine hcl tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51316-768
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	180 mg

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
AMMONIA (UNII: 5138Q19F1X)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL 8000 (UNII: Q662QK8M3B)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
GELATIN (UNII: 2G86QN327L)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

HYPROMELLOSE 2910 (3 MPA.S) (UNII: 0VUT3PMY82)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
HYDROXYPROPYL CELLULOSE (90000 WAMW) (UNII: UKE75GEA7F)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	gray ((White band in the middle))	**Score**	no score
Shape	OVAL ((capsule shaped tablet))	**Size**	20mm
Flavor		**Imprint Code**	G18
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-768-06	60 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211075	01/01/2024	

Labeler - CVS (062312574)

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