CHILDRENS FEXOFENADINE HYDROCHLORIDE ALLERGY- fexofenadine hydrochloride suspension Walgreens Company

Children's Fexofenadine HCl Allergy

Drug Facts

Active ingredient (in each 5 mL)

Fexofenadine HCl 30 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.

Directions

- shake well before using
- use only with enclosed dosing cup

adults and children 12 years of age and	take 10 mL every 12 hours; do not take more than 20 mL in	
over	24 hours	
children 2 to under 12 years of age	take 5 mL every 12 hours; do not take more than 10 mL in 24 hours	
children under 2 years of age	ask a doctor	
adults 65 years of age and older	ask a doctor	
consumers with kidney disease	ask a doctor	

Note: mL = milliliters

Other information

- each 5 mL contains: **sodium 14 mg**
- safety sealed: do not use if carton is opened or if foil inner seal on bottle is torn or missing
- store between 20° and 25°C (68° and 77°F)

Inactive ingredients

artificial raspberry flavor, edetate disodium, maltitol solution, poloxamer 407, potassium sorbate, propylene glycol, purified water, sodium benzoate, sodium phosphate dibasic heptahydrate, sodium phosphate monobasic monohydrate, sucralose powder, titanium dioxide and xanthan gum

Questions?

Call **1-866-923-4914**

DISTRIBUTED BY: WALGREEN CO. DEERFIELD, IL 60015

PRINCIPAL DISPLAY PANEL - 120 mL Bottle Carton

NDC 0363-2141-08

Children's Allergy Relief Fexofenadine HCI Oral Suspension, 30 mg/ 5 mL

4 FL OZ



CHILDRENS FEXOFENADINE HYDROCHLORIDE ALLERGY

fexofenadine hydrochloride suspension

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-2141
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII: E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	30 mg in 5 mL		

Inactive Ingredients				
Ingredient Name	Strength			
EDETATE DISODIUM (UNII: 7FLD91C86K)				
MALTITOL (UNII: D65DG142WK)				
POLOXAMER 407 (UNII: TUF2IVW3M2)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE (UNII: 70WT22SF4B)				
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
XANTHAN GUM (UNII: TTV12P4NEE)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor	RASPBERRY	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-2141- 08	1 in 1 CARTON	12/06/2024	
1		120 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0363-2141- 01	1 in 1 CARTON	12/06/2024	
2		240 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA208123	12/06/2024	

Establishment				
Name	Address	ID/FEI	Business Operations	
Taro Pharmaceutical Industries Ltd.		600072078	manufacture(0363-2141)	

Revised: 6/2025 Walgreens Company