DYE FREE WAL FEX CHILDRENS ALLERGY- fexofenadine hcl suspension Walgreens

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DRUG FACTS

Active ingredient (in each 5 mL)

Fexofenadine HCL, USP 30 mg

Purpose

Antihistamine

Uses

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

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 (1-800-222-

1222) right away.

Direction

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

adults and children 12 years of age and over	take 10 mL every 12 hours; do not take more than 20 mL in 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 18 mg
- store between 20º to 25ºC (68º and 77ºF)
- before using any medication, read all label directions. Keep carton, it contains important information.

Inactive ingredients

artificial raspberry flavor, butylparaben, edetate disodium, poloxamer 407, propylene glycol, propylparaben, purified water, sodium phosphate dibasic, sodium phosphate monobasic, sucrose, titanium dioxide, xanthan gum, xylitol

Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

*Compare to the active ingredient in Children's Allegra® Allergy††

NON-DROWSY

2 YEARS & OLDER

Children's Allergy Relief

FEXOFENADINE HCl ORAL SUSPENSION

30 mg /PER 5 mL / ANTIHISTAMINE

- DYE FREE
- NON-DROWSY
- 12 - HOUR RELIEF

- ALCOHOL FREE
- Indoor/outdoor allergy relief of sneezing, runny nose itchy nose or throat & itchy, watery eyes

AGES 2 YEARS & OLDER

Berry Flavor

FL OZ (mL)

USE ONLY WITH ENCLOSED DOSING CUP.

WASH AND LET AIR DRY AFTER EACH USE

††This product is not manufactured or distributed by Chattem Inc., distributor of Children's Allegra® Allergy.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

TAMPER EVIDENT: DO NOT USE IF CARTON, UNPRINTED FOIL INNER SEAL OR NECKBAND PRINTED WITH "SEAL FOR YOUR PROTECTION" IS OPENED, TORN OR MISSING

DISTRIBUTED BY: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015

Product Label

WALGREENS Children's Allergy Relief Berry Flavor

DYE FREE WAL FEX CHILDRENS ALLERGY

fexofenadine hcl suspension

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-9230
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
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPUU)	
SUCROSE (UNII: C151H8M554)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
XANTHAN GUM (UNII: TTV12P4NEE)	
XYLITOL (UNII: VCQ006KQ1E)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	BERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-9230-04	1 in 1 BOX	06/01/2020	
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:0363-9230-08	1 in 1 BOX	06/01/2020	
2		237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA203330	06/01/2020	

Labeler - Walgreens (008965063)

Revised: 6/2022 Walgreens