CHILDRENS ALLERGY- fexofenadine hcl suspension
H E B

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 to 77ºF)
- before using any medication, read all label directions. Keep carton, it contain 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 toll free: 1-**877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

Compare to Children's Allegra® Allergy active ingredient**

Childrens

ALLERGY

Fexofenadine HCl

Oral Suspension 30 mg/5mL

Antihistamine

Allergy Relief

Alcohol Free

Dye Free

Non-Drowsy

For Ages 2 years and Over

12 HOUR

Indoor/Outdoor Relief of:

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

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.

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

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

MADE WITH PRIDE AND CARE FOR H-E-B®

SAN ANTONIO, TX 78204

Product Label

H-E-B Children's Allergy Berry Flavor

CHILDRENS ALLERGY

fexofenadine hcl suspension

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-623
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:37808-623-04	1 in 1 CARTON	04/30/2021	
1		118 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	04/30/2021	

Labeler - H E B (007924756)

Revised: 6/2023 H E B