CHILDREN CLARITIN ALLERGY- loratadine solution
Bayer HealthCare LLC.

Children's Claritin Allergy (Grape Flavor)

Drug Facts

Active ingredient (in each 5 mL teaspoonful)

Loratadine 5 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 liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

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

Use only with enclosed dosing cup

- adults and children 6 years and over: 2 teaspoonfuls (tsp) daily; do not take more than 2 teaspoonfuls (tsp) in 24 hours

- children 2 to under 6 years of age: 1 teaspoonful (tsp) daily; do not take more than 1 teaspoonful (tsp) in 24 hours
- children under 2 years of age: ask a doctor
- consumers with liver or kidney disease: ask a doctor

- **each teaspoonful contains**: sodium 6 mg
- do not use if tape imprinted with “SEALED FOR YOUR PROTECTION” on top and bottom flaps of carton is not intact.
- store between 20° and 25°C (68° and 77°F)

edetate disodium, flavor, glycerin, maltitol, monobasic sodium phosphate, phosphoric acid, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose

Questions or comments?

1-800-CLARITIN (1-800-252-7484) or www.claritin.com

CHILDREN CLARITIN ALLERGY

loratadine solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-4360
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
SORBITOL (UNII: 506T60A25R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
MALTITOL (UNII: D65DG142WK)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
WATER (UNII: 059QF0KO0R)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-4360-5	1 in 1 CARTON	05/26/2015	
1		240 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11523-4360-2	1 in 1 CARTON	09/25/2015	
2		120 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:11523-4360-6	1 in 1 CARTON	10/05/2015	
3		180 mL in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:11523-4360-4	1 in 1 CARTON	11/01/2015	
4		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:11523-4360-7	2 in 1 CARTON	03/01/2016	
5		180 mL in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:11523-4360-3	1 in 1 CARTON	01/01/2016	
6		150 mL in 1 BOTTLE; Type 0: Not a Combination Product		

7	NDC:11523-4360-1	1 in 1 CARTON	05/26/2015	
7		60 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
NDA	NDA020641	05/26/2015	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 11/2023 Bayer HealthCare LLC.