

**CHILDREN CLARITIN ALLERGY- loratadine solution**  
**Bayer HealthCare LLC.**

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**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: 10 mL daily; do not take more than 10 mL in 24 hours

- children 2 to under 6 years of age: 5 mL daily; do not take more than 5 mL in 24 hours
- children under 2 years of age: ask a doctor
- consumers with liver or kidney disease: ask a doctor
  
- **each 5 mL 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](http://www.claritin.com)



## CHILDREN CLARITIN ALLERGY

loratadine solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11523-4360
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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<b>LORATADINE</b> (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	5 mg in 5 mL
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<b>Inactive Ingredients</b>	
<b>Ingredient Name</b>	<b>Strength</b>
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>MALTITOL</b> (UNII: D65DG142WK)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PHOSPHORIC ACID</b> (UNII: E4GA8884NN)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM PHOSPHATE, MONOBASIC</b> (UNII: 3980JH2SW)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	

<b>Product Characteristics</b>			
<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	GRAPE	<b>Imprint Code</b>	
<b>Contains</b>			

<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
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	
		60 mL in 1 BOTTLE; Type 0: Not a Combination Product		

7	60 mL in 1 BOTTLE; Type O: Not a Combination Product		
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**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: 5/2025

Bayer HealthCare LLC.