UP AND UP CHILDRENS ALLERGY RELIEF- loratadine solution Target Corporation

Target Corporation Children's Allergy Relief 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
- 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

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 (1-800-222-1222).

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 |

Other information

- do not use if carton is opened, or if printed neckband is broken or missing
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

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

Questions?

Call 1-888-547-7400

Package/Label Principal Display Panel

Compare to active ingredient in Children's Claritin® dye-free children's allergy relief loratadine oral solution 5 mg/5 mL antihistamine indoor and outdoor allergies non-drowsy* 24-hour relief of: sneezing runny nose itchy, watery eyes itchy throat or nose sugar-free alcohol-free dosing cup included *When taken as directed. See drug facts panel 24 HOUR RELIEF **GRAPE FLAVOR**

AGES 2+ YEARS 4 FL OZ (120 mL)



UP AND UP CHILDRENS ALLERGY RELIEF

loratadine solution

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:11673-558

| Active Ingredient/Active Moiety | | | | | | | | |
|--|--|---------------------|----------------------|----------------------|-----------|----------|--|--|
| Ingredient Name | | | Basis of Stre | ength S | trength | | | |
| LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN) | | | LORATADINE | 5 mg | g in 5 mL | | | |
| | | | | | | | | |
| | | | | | | | | |
| Inactive Ingredients | | | | | | | | |
| Ingredient Name | | | | | | ength | | |
| EDETATE DISO DIUM (UNII: 7FLD91C86K) | | | | | | | | |
| GLYCERIN (UNII: PDC6 | | | | | | | | |
| MALTITOL (UNII: D65D | | | | | | | | |
| SO DIUM PHO SPHATE, MO NO BASIC (UNII: 3980 JIH2SW) | | | | | | | | |
| PHOSPHORIC ACID (UNII: E4GA8884NN) | | | | | | | | |
| | PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | | | | | | | |
| WATER (UNII: 059QF0K | | | | | | | | |
| SODIUM BENZOATE (U | | 25EU) | | | | | | |
| SORBITOL (UNII: 506T60A25R) | | | | | | | | |
| SUCRALOSE (UNII: 96K | (6UQ3ZD4) | | | | | | | |
| | | | | | | | | |
| Product Characteristics | | | | | | | | |
| Color | | | Score | | | | | |
| Shape | Size | | Size | | | | | |
| Flavor | vor GRAPE Imprint Code | | e | | | | | |
| Contains | | | | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| Packaging | | | | | | | | |
| # Item Code | | Package Description | | Marketing Start Date | Marketing | End Date | | |
| 1 NDC:11673-558-26 1 i | in 1 CARTON | ARTON | | 06/05/2018 | | | | |
| 120 mL in 1 BOTTLE; Type 0: Not a Combination Product | | | | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| Marketing Information | | | | | | | | |
| Marketing Category | ting Category Application Number or Monograph Citation | | Marketing Start Date | Marketing End Date | | | | |
| ANDA | ANDA07572 | 28 | | 06/05/2018 | 18 | | | |
| | | | | | | | | |

Labeler - Target Corporation (006961700)

Revised: 6/2018

Target Corporation