TRIPROLIDINE HYDROCHLORIDE- triprolidine hydrochloride syrup
Westminster Pharmaceuticals, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Triprolidine HCl

Drug Facts

Active ingredient (in each 1 mL dropperful)

Triprolidine HCl 0.938 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever (allergic rhinitis) or other upper respiratory allergies:

- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes

Warnings

Do not exceed recommended dosage.

Ask a doctor before use if you have

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged prostate gland

Ask a doctor before use if you are taking sedatives or tranquilizers

When using this product

- may cause drowsiness
- excitability may occur, especially in children
- alcohol, sedatives and tranquilizers may increase the drowsiness effect
- avoid alcoholic beverages
- use caution when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

- new symptoms occur

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

- **do not exceed recommended dosage.**
- use only the enclosed dropper.
- do not use enclosed dropper for any other drug product.

AGE	*DOSE*
Adults & Children 12 years of age or older:	2.67 mL (2.5 milligrams) every 4 to 6 hours, not to exceed 10.67 mL (10 milligrams) in 24 hours, or as directed by a doctor.
Children 6 to under 12 years of age:	1.33 mL (1.25 milligrams) every 4 to 6 hours, not to exceed 5.33 mL (5 milligrams) in 24 hours, or as directed by a doctor.
Children under 6 years of age:	consult a doctor.

Other Information

- This packaging is child-resistant.
- Store at room temperature 15°-30° C (59°-86° F).
- Tamper evident by foil seal under cap. Do not use if foil seal is missing or broken.

Inactive ingredients

bubble gum flavor, citric acid, glycerin, methylparaben, monoammonium glycyrrhizinate, potassium citrate, potassium sorbate, propylene glycol, propylparaben, purified water, sucralose.

Questions? Comments?

Call 1-844-221-7294.

PRINCIPAL DISPLAY PANEL - 30 mL Bottle Carton

NDC 69367-253-30

Triprolidine HCl
Antihistamine
Each dropperful (1 mL) contains:
Triprolidine HCl 0.938 mg

Sugar-Free • Dye Free
Alcohol Free

Tamper evident by foil seal under cap.
Do no use if foil seal is broken or missing.

Bubble Gum Flavor
1 fl oz (30 mL)

Westminster
Pharmaceuticals

TRIPROLIDINE HYDROCHLORIDE

triprolidine hydrochloride syrup

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69367-253
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRIPROLIDINE HYDROCHLORIDE (UNII: YAN7R5L890) (Triprolidine - UNII:2L8T9S52QM)	TRIPROLIDINE HYDROCHLORIDE	0.938 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
AMMONIUM GLYCYRRHIZATE (UNII: 3VRD35U26C)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics			
Color		**Score**	
Shape		**Size**	
Flavor	BUBBLE GUM	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69367-253-30	1 in 1 CARTON	03/09/2020	
1		30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part341	03/09/2020	

Labeler - Westminster Pharmaceuticals, LLC (079516651)

Revised: 5/2020 Westminster Pharmaceuticals, LLC