HISTEX- triprolidine hydrochloride syrup Allegis Pharmaceuticals, LLC

HISTEX[™] Syrup

Drug Facts

Active ingredients (in each 5 mL teaspoonful)

Triprolidine HCl 2.5 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

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

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

When using this product

- excitability may occur, especially in children
- may cause drowsiness
- 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 the reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

Do not exceed recommended dosage.

AGE	DOSE				
Adults and Childen 12 years of age and older:	1 teaspoonful (5 mL) every 4 to 6 hours, not to exceed 4 teaspoonfuls (20 mL) in 24 hours, or as directed by a doctor				
Childen 6 to under 12 years of age	¹ / ₂ teaspoonful (2.5 mL) every 4 to 6 hours, not to exceed 2 teaspoonfuls (10 mL) in 24 hours, or as directed by a doctor.				
Children under 6 years of age	Consult a doctor				

Other Information

Store at 15°-30° C (59°-86° F).

Tamper evident by foil seal under cap. Do not use if foil seal is missing or broken.

Dispense in a tight, light-resistant container with a child-resistant cap.

Inactive ingredients

Citric Acid, Glycerin, Propylene Glycol, Purified Water, Sodium Citrate, Sodium Saccharin, Sorbitol, Bubble Gum Flavor.

Questions? Comments?

Call 1-866-633-9033.

PRINCIPAL DISPLAY PANEL - 237 mL Bottle Label

NDC 28595-802-08

Antihistamine

HISTEX™

Syrup

contains:

Triprolidine HCl 2.5 mg

Sugar-Free • Dye Free

Alcohol Free

Bubble Gum Flavor

8 fl oz (237 mL)



Drug Facts (continued)		Drug Facts (continued)			
<i>Directions</i> Do not exceed recommended dosage.		Tamper evident by foil seal under cap. Do not use if foil seal is missing or broken.			
AGE	DOSE	Inostivo ingradianto			
Adults and Children 12 years of age and older:	1 teaspoonful (5 mL) every 4 to 6 hours, not to exceed 4 teaspoonfuls (20 mL) in 24 hours, or as directed by a doctor	Inactive ingredients Bubble gum flavor, citric acid, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sodium saccharin, sorbitol solution			
Children 6 to under 12 years	1/2 teaspoonful (2.5 mL) every 4 to 6 hours, not				
of age	to exceed 2 teaspoonfuls (10 mL) in 24 hours, or as directed by a doctor.	Questions? Comments? Call 1-866-633-9033.			
Children under 6 years of age	Consult a doctor	Manufactured for: Allegis Pharmaceuticals, LLC			
Other information Store at 15°-30° C (59°-86° F).		Canton, MS 39046 www.allegispharma.com Rev. 03/22			

triprolidine hydrochloride syrup

triproliaine hydroci	nioride syru	p							
Product Inform	nation								
Product Type		HUMAN OTC DRUG	ltem (Code (Source)		NDC:28	8595-802		
Route of Adminis	tration	ORAL					100.20333 002		
Route of Adminis	tration	ONAL							
Active Ingredie	nt/Active	Moiety							
Ingredient Name					Basis of Strength Stren				
	TRIPROLIDINE HYDROCHLORIDE (UNII: YAN7R5L890) (TRIPROLIDINE		OLIDINE	-	TRIPROLIDINE 2.5		2.5 mg		
UNII:2L8T9S52QM)					HYDROCHLORIDE		in 5 mL		
Inactive Ingred	lients								
Ingredient Name							Strength		
SODIUM BENZOATE	: (UNII: 0 245F	-							
CITRIC ACID MONO		•							
GLYCERIN (UNII: PDC6A3C0OX)									
PROPYLENE GLYCO	L (UNII: 6DC90	Q167V3)							
WATER (UNII: 059QF	0KO0R)								
SODIUM CITRATE, U	JNSPECIFIED	FORM (UNII: 1Q73Q2JULR)							
SACCHARIN SODIUM	4 (UNII: SB8Z	UX40TY)							
SORBITOL (UNII: 506T60A25R)									
Product Charac	cteristics								
Color		Score							
Shape			Size						
Flavor	BUB	BLE GUM	Imprint Code						
Contains									
Dackaging									
Packaging									
# Item Code	Pa	ckage Description		Mar	keting Start Date	Mark	ceting End Date		
	37 mL in 1 BC roduct	n 1 BOTTLE; Type 0: Not a Combination		03/04/2014					
Marketing I	nformat	ion							
Marketing Category		tion Number or Monog Citation	Iraph	Ма	rketing Start Date	Mar	keting End Date		
OTC Monograph Drug	M012			03/04	/2014				
5 5									