HISTEX PDX DROPS- triprolidine hydrochloride syrup Allegis Pharmaceuticals, LLC

HISTEX™ PDX Drops

Drug Facts

Active ingredient (in each 1 mL dropperful)

Triprolidine HCl 1.25 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 the child has

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma

Ask a doctor before use if the child is taking sedatives or tranquilizers

When using this product

- excitability may occur, especially in children
- may cause drowsiness
- sedatives and tranquilizers may increase the drowsiness effect

Stop use and ask a doctor if

new symptoms occur

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.

- use only the enclosed dropper.
- do not use enclosed dropper for any other drug products.

AGE	DOSE
Adutls & Children 12 years of age or older:	2 mL (2.5 mg) every 4 to 6 hours, not to exceed 8 mL (10 mg) in 24 hours or as directed by a doctor.
Children 6 to under12 years of age:	1 mL (1.25 mg) every 4 to 6 hours, not to exceed 4 mL (5 mg) in 24 hours or as directed by a doctor.
Children under 6 years of age:	Consult a docotor.

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.

Inactive ingredients

bubble gum flavor, citric acid, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose.

Questions? Comments?

Call 1-866-633-9033.

PRINCIPAL DISPLAY PANEL - 30 mL Bottle Carton

NDC 28595-808-30

HISTEX™ PDX

Drops

Antihistamine

Each dropperful (1 mL)

contains:

Triprolidine HCI

1.25 mg

Sugar-Free • Dye Free

Alcohol Free

Bubble Gum Flavor

1 fl oz (30 mL) Bottle

Drug Facts Active ingredient Purpose (in each 1 mL dropperful) Triprolidine HCI 1.25 mg ... 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 the child has ■ a breathing problem such as emphysema or chronic bronchitis glaucoma Ask a doctor before use if the child is taking sedatives or tranquilizers When using this product ■ excitability may occur, especially in children ■ may cause drowsiness ■ sedatives and tranquilizers may increase the drowsiness effect Stop use and ask a doctor if ■ new symptoms occur Keep out of the reach of children. In case of overdose, get medical help or contact a Poison Control Directions Do not exceed recommended dosage. use only with enclosed dropper do not use enclosed dropper for any other drug product ΔGF DOSE Adults & Children 12 2 mL (2.5 mg) every 4 to 6 hours, not to exceed years of age or older: 8 mL (10 mg) in 24 hours, or as directed by a doctor. Children 6 to under 1 mL (1.25 mg) every 4 to 6 hours, not to exceed 12 years of age: 4 mL (5 mg) in 24 hours, or as directed by a doctor. Children under Consult a doctor. 6 years of age:



HISTEX PDX DROPS

triprolidine hydrochloride syrup

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:28595-808
Route of Administration	ORAL		

Other Information

broken.

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
	TRIPROLIDINE HYDROCHLORIDE	1.25 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
SUCRALOSE (UNII: 96K6UQ3ZD4)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
GLYCERIN (UNII: PDC6A3C0OX)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			

SORBITOL (UNII: 506T60A25R)

Product Characteristics

Color		Score	
Shape		Size	
Flavor	BUBBLE GUM	Imprint Code	
Contains			

Packaging

- actualing				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:28595-808- 30	1 in 1 CARTON	01/16/2023	
1		30 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
OTC Monograph Drug	M012	01/16/2023	

Labeler - Allegis Pharmaceuticals, LLC (792272861)

Revised: 1/2024 Allegis Pharmaceuticals, LLC