

HISTEX PD DROPS NEW FORMULATION- triprolidine hydrochloride syrup
Allegis Pharmaceuticals, LLC

HISTEX™ PD New Formulation

Drug Facts

Active ingredient (in each 1 mL dropperful)

Tripolidine 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
Adults & 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 under 12 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 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.

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 28595-809-30

HISTEX™ PD New Formulation

Drops

Antihistamine

Each dropperful (1 mL)

contains:

Triprolidine HCl

1.25 mg

Sugar-Free • Dye Free

Alcohol Free

Bubble Gum Flavor

1 fl oz (30 mL) Bottle

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 with enclosed dropper ■ do not use enclosed dropper for any other drug product	
AGE	DOSE
Adults & 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 under 12 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 doctor.

Drug Facts (continued)
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
Questions? Comments? Call 1-866-633-9033.

NDC 28595-809-30

NEW FORMULATION

HISTEXTM PD Drops

Antihistamine

Each dropperful (1 mL) contains:
Triprolidine HCl 1.25 mg

Sugar-Free • Dye Free
Alcohol Free

Bubble Gum Flavor

Tamper Evident: Do not use if seal is broken or missing

1 fl oz (30 mL)

KEEP OUT OF REACH OF CHILDREN.

Manufactured for:
Allegis Pharmaceuticals, LLC
Canton, MS 39046
www.allegispharma.com

Rev. Date: 12/2023

CALIBRATED DROPPER ENCLOSED

HISTEX PD DROPS NEW FORMULATION

triprolidine hydrochloride syrup

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:28595-809
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRIPROLIDINE HYDROCHLORIDE (UNII: YAN7R5L890) (TRIPROLIDINE - UNII:2L8T9S52QM)	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)	

SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)

SORBITOL (UNII: 506T60A25R)

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:28595-809-30	1 in 1 CARTON	12/13/2023	
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	12/13/2023	

Labeler - Allegis Pharmaceuticals, LLC (792272861)

Revised: 1/2025

Allegis Pharmaceuticals, LLC