

TRIPROLIDINE HCL DROPS- triprolidine hydrochloride liquid
Brandywine Pharmaceuticals, LLC

Triprolidine HCl Drops

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NDC 71321-701-50

Active ingredients (in each 2 mL)

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 ■ itching of the nose or throat
- sneezing ■ 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
- trouble urinating due to an enlarged prostate

Ask a doctor before use if the child is

taking sedatives or tranquilizers

When using this product

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

Stop use and ask a doctor if

- new symptoms occur

Keep out of reach of children.

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

Directions

- use only with enclosed dropper ■ mL= milliliter
- do not use dropper for any other drug product

children 6 to under 12 years: 2 mL every 6 hours, not to exceed 4 doses in 24 hours, or as directed by a doctor.

children under 6: ask a doctor.

Other information

- read all product information before using
- this packaging is child-resistant.
- store at room temperature 20-30°C (68-86°F)

Inactive ingredients

citric acid anhydrous, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sodium saccharin, sorbitol solution

Questions or comments?

Call 610-314-7943 9 a.m. - 5 p.m. EST.

PRINCIPAL DISPLAY PANEL

NDC 71321-701-50

Triprolidine HCl Drops

1.69 fl oz (50 mL)

**TAMPER EVIDENT:
DO NOT USE IF FOIL SEAL
UNDER CAP IS BROKEN
OR MISSING**

71321-701-50

Triprolidine HCl Drops

Antihistamine

Each dropperful (2 mL), for oral administration, contains:
Triprolidine HCl 1.25 mg

Gluten Free • Dye Free
Sugar Free • Alcohol Free

Manufactured for: Brandywine Pharmaceuticals, LLC
West Chester, PA, 19382

RELIEVES:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Nose
- Itchy Throat



Rev 1/2023

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PEEL
HERE

<p>Drug Facts</p> <table border="1"> <thead> <tr> <th>Active ingredients (in each 2 mL dropperful)</th> <th>Purpose</th> </tr> </thead> <tbody> <tr> <td>Triprolidine HCl 1.25 mg.....</td> <td>Antihistamine</td> </tr> </tbody> </table> <p>Uses</p> <ul style="list-style-type: none"> temporarily relieves these symptoms due to hay fever (allergic rhinitis) or other upper respiratory allergies: <ul style="list-style-type: none"> runny nose sneezing itching of the nose or throat itchy, watery eyes <p>Warnings</p> <p>Do not exceed recommended dosage.</p> <p>Ask a doctor before use if the child has</p> <ul style="list-style-type: none"> a breathing problem such as emphysema or chronic bronchitis glaucoma trouble urinating due to an enlarged prostate gland <p>Ask a doctor before use if the child is taking sedatives or tranquilizers</p> <p>When using this product</p> <ul style="list-style-type: none"> excitability may occur, especially in children marked drowsiness may occur sedatives and tranquilizers may increase the drowsiness effect <p>Stop use and ask a doctor if new symptoms occur</p> <p>Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.</p> <p>Directions</p> <ul style="list-style-type: none"> use only with enclosed dropper do not use dropper for any other drug product <table border="1"> <thead> <tr> <th>Age</th> <th>Dose</th> </tr> </thead> <tbody> <tr> <td>Children 6 to under 12 years:</td> <td>2 mL every 6 hours, not to exceed 4 doses in 24 hours, or as directed by a doctor.</td> </tr> <tr> <td>children under 6:</td> <td>ask a doctor.</td> </tr> </tbody> </table> <p>Other information</p> <ul style="list-style-type: none"> read all product information before using this packaging is child-resistant store at room temperature 20-30°C (68-86°F) <p>Inactive ingredients</p> <p>citric acid anhydrous, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sodium saccharin, sorbitol solution</p> <p>Questions or comments? Call 610-314-7943 9 a.m. - 5 p.m. EST</p>	Active ingredients (in each 2 mL dropperful)	Purpose	Triprolidine HCl 1.25 mg.....	Antihistamine	Age	Dose	Children 6 to under 12 years:	2 mL every 6 hours, not to exceed 4 doses in 24 hours, or as directed by a doctor.	children under 6:	ask a doctor.	<p>TEMPORARILY RELIEVES:</p> <ul style="list-style-type: none"> Sneezing Runny Nose Itchy, Watery Eyes Itchy Nose Itchy Throat <p>Gluten Free Dye Free Sugar Free Alcohol Free</p> <p>Manufactured for: Brandywine Pharmaceuticals, LLC West Chester, PA, 19382</p> <p>Rev 1/2023</p>	<p>71321-701-50</p> <h2>Triprolidine HCl Drops</h2> <p>Antihistamine</p> <p>Each dropperful (2 mL), for oral administration, contains: Triprolidine HCl 1.25 mg</p> <p>Sneezing • Runny Nose Itchy, Watery Eyes Itchy Nose • Itchy Throat</p> <p>Gluten Free • Dye Free Sugar Free • Alcohol Free</p> <p>BRANDYWINE PHARMACEUTICALS, LLC</p> <p>1.69 fl oz (50 mL)</p>	<p>WARNING: KEEP OUT OF REACH OF CHILDREN</p> <p>TAMPER EVIDENT: DO NOT USE IF FOIL SEAL UNDER CAP IS BROKEN OR MISSING</p> <p>Use with enclosed calibrated dropper</p>
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TRIPROLIDINE HCL DROPS

triprolidine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71321-701
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 2 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71321-701-50	50 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	03/08/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/08/2023	

Labeler - Brandywine Pharmaceuticals, LLC (080581956)

Revised: 1/2024

Brandywine Pharmaceuticals, LLC