## EQUATE EYE DROPS- tetrahyrozoline hydrochloride liquid Prestige Brands Holdings, Inc.

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.

-----

#### **Equate Eye Drops**

#### **Drug Facts**

#### Active ingredient

Tetrahydrozoline Hydrochloride 0.05%

#### **Purpose**

Redness reliever

#### Uses

• relieves redness of the eye due to minor eye irritations

#### **Warnings**

#### For external use only

#### Do not use

if solution changes color or becomes cloudy

#### Ask a doctor before use if you have

• narrow angle glaucoma

#### When using this product

- to avoid contamination, do not touch tip of container to any surface
- replace cap after using
- overuse may produce increased redness of the eye
- pupils may become enlarged temporarily

#### Stop use and ask a doctor if

- you experience eye pain
- you experience changes in vision
- you experience continued redness or irritation of the eye
- the condition worsens or persists for more than 72 hours

#### Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

#### Directions

• instill 1 to 2 drops in the affected eye(s) up to four times daily

## Other information

- store at room temperature
- remove contact lenses before using

## **Inactive ingredients**

benzalkonium chloride, boric acid, edetate disodium, purified water, sodium borate

### Questions?

1-888-287-1915

## **Principal Display Panel**

Equate<sup>®</sup>
Original Redness Reliever
Tetrahydrozoline HCI
Sterile Eye Drops
1 FL OZ (30 mL)



## **Principal Display Panel**

Equate<sup>®</sup>
Original Redness
Reliever
Tetrahydrozoline HCI
Sterile Eye Drops
0.5 FL OZ (15 mL)



## **EQUATE EYE DROPS**

tetrahyrozoline hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67172-282
Route of Administration	ОРНГНАЬМІС		

Active Ingredient/Active Moiety				
Basis of Strength	Strength			
ΓΕΤRAHYDROZOLINE HYDROCHLORIDE	0.5 mg in 1 mL			
	TETRAHYDROZOLINE			

#### **Inactive Ingredients**

Ingredient Name	Strength
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	

	Packaging				
;	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:67172-282- 01	1 in 1 BOX	02/11/2013		
	1	15 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	part349	0 2/11/20 13		

## **EQUATE EYE DROPS**

tetrahyrozoline hydrochloride liquid

Product Type HUMAN OTC DRUG Item Code (Source) NDC:67172-281

Route of Administration OPHTHALMIC

# Active Ingredient/Active Moiety Ingredient Name Basis of Strength TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0 YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077) TETRAHYDROZOLINE hYDROCHLORIDE 0.5 mg in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
BORIC ACID (UNII: R57ZHV85D4)				
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)				
WATER (UNII: 059QF0KO0R)				
SODIUM BORATE (UNII: 91MBZ8H3QO)				

Packaging			
# Itam Cada	Doologo Doogrintion	Marketing Start	Marketing End

	#	Item Code		rackage Description	Date	Date
	1	NDC:67172-281- 01	1 in 1 E	3OX	0 2/11/20 13	
	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product			, , , , , , , , , , , , , , , , , , , ,		
Marketing Information						
Marketing Category		gory	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH FINAL		FINAL	part349	02/11/2013		

## Labeler - Prestige Brands Holdings, Inc. (159655021)

Revised: 1/2020 Prestige Brands Holdings, Inc.