UP AND UP ULTRA ITCHY RELIEF- tetrahydrozoline hci and zinc sulfate solution/ drops
Target Corporation

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.

Up & Up Ultra Itch Relief Eye Drops 0.5 oz NBE Visine AC 2019

Active ingredients Purposes

Tetrahydrozoline HCI 0.05%.....Eye redness reliever

Zinc Sulfate 0.25%.....Eye astringent

Uses

- for temporary relief of discomfort and redness of the eye due to minor eye irritations

Warnings

For external use only

Ask a doctor before use if you have narrow angle glaucoma.

When using this product

- pupils may become enlarged temporarily
- overuse may cause more eye redness
- remove contact lenses before using
- do not use if this solution changes color or becomes cloudy
- do not touch tip of container to any surface to avoid contamination
- replace cap after each use

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur
- redness or irritation of the eye lasts
- condition worsens or lasts more than 72 hours

If pregnant or breast-feeding, ask a health professional before use.

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

Directions

- to open bottle, push cap down and twist counterclockwise
- put 1 to 2 drops in the affected eye(s) up to 4 times daily
- children under 6 years of age: ask a doctor

Other information

- some users may experience a brief tingling sensation
- store between 15-25°C (59-77°F)

Inactive ingredients

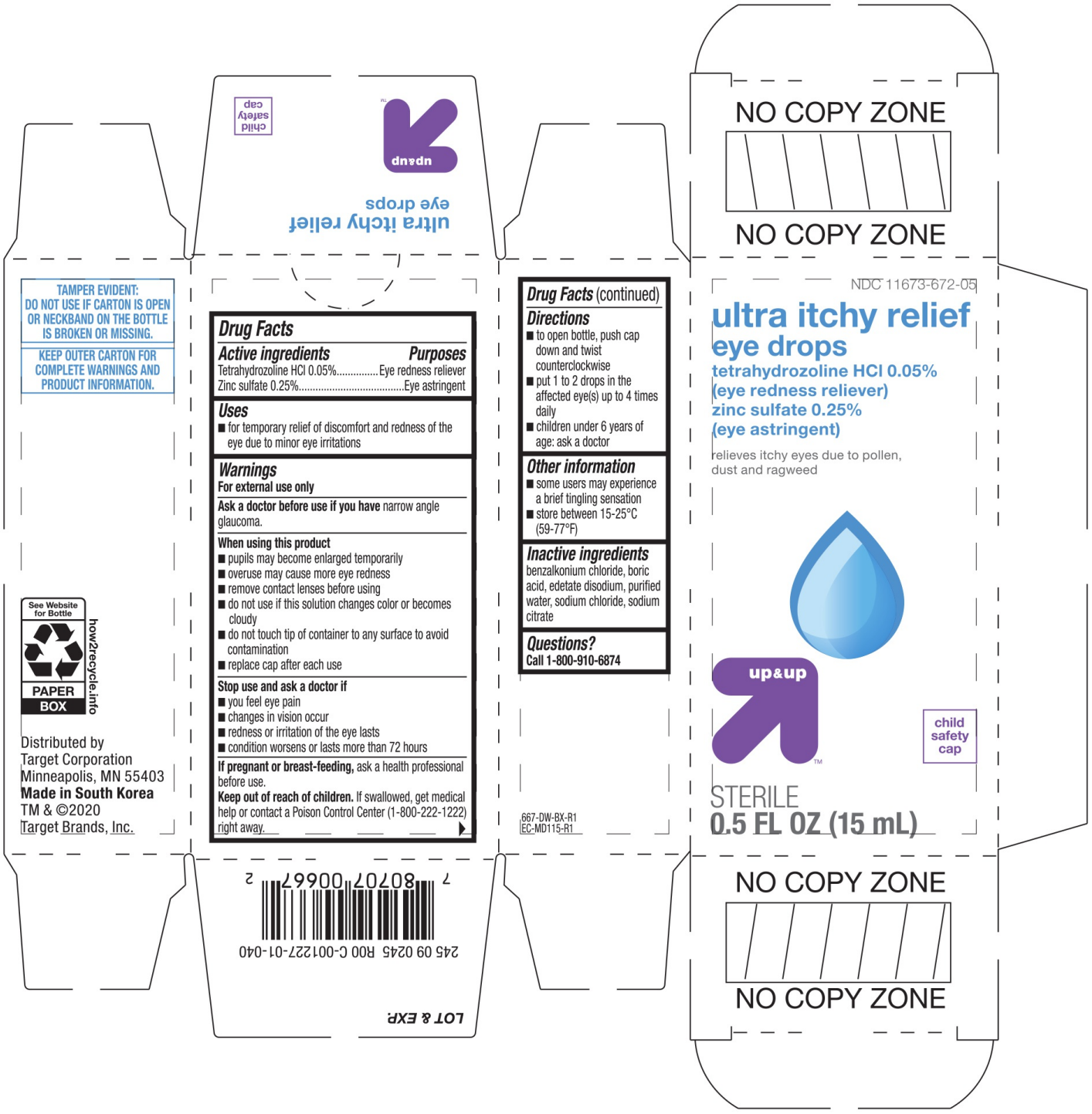
benzalkonium chloride, boric acid, edetate disodium, purified water, sodium chloride, sodium citrate

Distributed by

Target Corporation

Minneapolis, MN 55403

Made in Korea



UP AND UP ULTRA ITCHY RELIEF

tetrahydrozoline hci and zinc sulfate solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-672
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.5 mg in 1 mL
ZINC SULFATE (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC SULFATE	2.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-672-05	1 in 1 CARTON	09/20/2019	
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	09/20/2019	

Labeler - Target Corporation (006961700)

Revised: 11/2019 Target Corporation