EXTREME RELIEF LUBRICANT- polyethylene glycol 400, and propylene glycol 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.

Active ingredients Purpose

Polyethylene glycol 400 0.4%..... Lubricant

Propylene glycol 0.3%..... Lubricant

Uses

For the temporary relief of burning and irritation due to eye dryness

Warnings

For external use only

retain carton for full drug facts

Do not use
- if solution changes color or becomes cloudy
- if you are sensitive to any ingredient in this product

When using this product
- 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(s) gets worse, persists or last more than 72 hours

Keep this and all drugs out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Directions
- Shake well before use
- Instill 1 or 2 drops in the affected eye(s) as needed

Other information
- Store at room temperature

Inactive ingredients

boric acid, calcium chloride, 20% chlorhexidine, glyconate, hydrochloric acid, hypromellose 2910, magnesium chloride, potassium chloride, purified water, sodium chloride, sodium hydroxide, zinc chloride

DISTRIBUTED BY:

TARGET CORP.

MINNEAPOLIS, MN 55403

MADE IN KOREA

extreme relief

lubricant eye drops

**Compare to Systane®
Ultra Eye Drops***

helps relieve minor eye irritations
lubricates and refreshes
fast-acting

STERILE
0.5 FL OZ (15 mL)

extreme relief
lubricant eye drops

up&up

DO NOT USE IF CARTON IS OPEN OR GRAY NECKBAND ON THE BOTTLE IS BROKEN OR MISSING.

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Drug Facts (continued)

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Questions? Call
1-800-910-6874

*This product is not manufactured or distriuted by Alcon, owner of the registered trademark Systane® Ultra Eye Drops.

Dist. by Target Corp.
Minneapolis, MN 55403
Made in Korea

LOT NO.:
EXP. DATE:

245 09 0280 R00
IDC-000021-01-094

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EXTREME RELIEF LUBRICANT

polyethylene glycol 400, and propylene glycol solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-576
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	4 mg in 1 mL

PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
MAGNESIUM CHLORATE (UNII: M536P01U3N)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ZINC CHLORIDE (UNII: 86Q357L16B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-576-15	1 in 1 BOX		
1		15 mL in 1 BOTTLE, DROPPER		
2	NDC:11673-576-30	2 in 1 BOX		
2		15 mL in 1 BOTTLE, DROPPER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	03/04/2015	

Labeler - Target Corporation (006961700)

Revised: 3/2015 Target Corporation