

SOUNDBODY LONG LASTING LUBRICANT- polyethylene glycol 400, propylene glycol solution/ drops
United Exchange Corp.

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

Do not use

- if solution changes color or becomes cloudy
- if you are sensitive or allergic to any ingredient in this product

When using this product

- to avoid contamination, do not touch tip of container to any surface
- replace cap after each use

Stop use and ask a doctor if

- you experience eye pain
- changes in vision occur
- redness or irritation of the eye(s) gets worse or lasts more than 72 hours

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center immediately

Directions

- Put 1 or 2 drops in the affected eye(s) as needed

Other information

- Store at 15° to 30°C (59° to 86°F)

Inactive ingredients

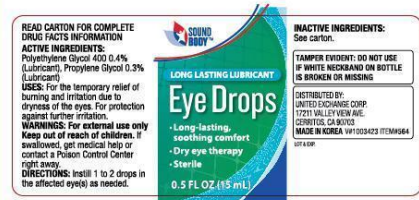
boric acid, calcium chloride, chlorhexidine gluconate, hydrochloric acid, hypromellose 2910, magnesium chloride, potassium chloride, purified water, sodium chloride, sodium hydroxide, zinc chloride

DISTRIBUTED BY:

UNITED EXCHANGE CORP.

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SOUNDBODY LONG LASTING LUBRICANT

polyethylene glycol 400, propylene glycol solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-644
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: R57ZH85D4)	
CALCIUM CHLORIDE (UNII: M410D6VV5M)	
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

HYPROMELLOSE 2910 (15 MPA.S) (UNII: 36SFW2JZ0W)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0K00R)	
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:65923-644-15	1 in 1 BOX		
1		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	02/26/2014	

Labeler - United Exchange Corp. (840130579)

Revised: 2/2014

United Exchange Corp.