SOUNDBODY ADVANCED RELIEF- dextarn 70, polyethylene glycol 400, povidone, tetrahydrozoline hci 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

Dextran 70 0.1%..... Lubricant

Polyethylene glycol 400 1%..... Lubricant

Povidone 1%..... Lubricant

Tetrahydrozoline HCI 0.05%..... Redness reliever

Uses

- for the relief of redness of the eyes due to minor eye irritations
- for protection against further irritation

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
- 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 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 Poision Control Center right away

Directions

- to open bottle, push cap down and twist counterclockwise. To close bottle, twist clockwise until it stops turning.
- 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 at 15°-25°C (59°-77°F)

Inactive ingredients

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

DISTRIBUTED BY:

UNITED EXCHANGE CORP.

17211 VALLEY VIEW AVE.

CERRITOS, CA 90703 USA

Made in Korea

SOUNDBODY ADVANCED RELIEF

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65923-637

Route of Administration	OPHTHALMIC

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DEXTRAN 70 (UNII: 7SA290YK68) (DEXTRAN 70 - UNII:7SA290YK68)	DEXTRAN 70	1 mg in 1 mL
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	10 mg in 1 mL
POVIDONE (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE	10 mg in 1 mL
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65923-637-15	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	11/25/2014	

Labeler - United Exchange Corp. (840130579)

Revised: 11/2014 United Exchange Corp.