SOOTHE LONG LASTING- povidone solution
Bausch & Lomb Incorporated

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.

Drug Facts

Active ingredient

Povidone (2.0%)

Purpose

Lubricant

Uses

■ relieves dryness of the eye

■ prevents further irritation

Warnings

For external use only.

Do not use

■ if solution changes color or becomes cloudy

When using this product

■ do not touch tip of container to any surface to avoid contamination

■ remove contact lenses before using

■ replace cap after using

Stop use and ask a doctor if you experience

■ eye pain

■ change in vision

■ continued redness or irritation of the eye

■ or if condition worsens or lasts more than 72 hours

If pregnant or breastfeeding

ask a health professional before use.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control

Center right away.

Directions

■ Instill 1 or 2 drop(s) in the affected eye(s) as needed or as directed by your doctor

Other information

■ Storage: 15°-25°C (59°-77°F)

■ Use before expiration date marked on the carton or bottle

Inactive ingredients

Benzalkonium chloride (0.005%), boric acid, edetate disodium dihydrate, purified water, sodium borate, sodium chloride, tyloxapol. Hydrochloric acid and/or sodium hydroxide may be used to adjust pH.

Questions?

Call:1-800-553-5340

Package/Label Principal Display Panel

Bausch & Lomb

Soothe

Lubricant Eye Drops

LONG LASTING

Dry Eye Therapy

• Instant and Long Lasting Relief

• Moisturizes and Protects

• For All Types of Dry Eye

Sterile

0.5 FL OZ (15mL)

SOOTHE LONG LASTING

povidone solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24208-544
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E) (POVIDONE, UNSPECIFIED - UNII:FZ989GH94E)	POVIDONE, UNSPECIFIED	20 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)	
TYLOXAPOL (UNII: Y27PUL9H56)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:24208-544-01	1 in 1 CARTON	06/01/2013	
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	06/01/2013	

Labeler - Bausch & Lomb Incorporated (196603781)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		079587625	MANUFACTURE(24208-544)

Revised: 12/2017 Bausch & Lomb Incorporated