HYDRATE LUBRICANT- polyethylene glycol 400 and propylene glycol solution/ drops
Sproose Products Inc.

HYDRATE Lubricant

Drug Facts

Active Ingredients	Purposes
Polyethylene glycol 400 0.04%	Lubricant
Propylene glycol 0.3%	Lubricant

Uses
- for the temporary relief of burning and irritation due to dryness of the eye

Warnings

For external use only

Do not use
- if this product changes color or becomes cloudy
- if you are sensitive 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
- remove contacts before using

Stop use and ask a doctor if you experience any of the following
- eye pain
- changes in vision
- continued redness or irritation of the eye
- the condition worsens or persists for more than 72 hours

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

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

Other information

store at 15°-30°C (59°-86°F)

Inactive Ingredients

Aminomethylpropanol, benzalkonium chloride as preservative, boric acid, hypromellose, potassium chloride, purified water, sodium chloride, sorbitol. May contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Questions?

866-778-0043 Alchemy43.com

Distributed by
Sproose Products Inc.
West Hollywood, CA

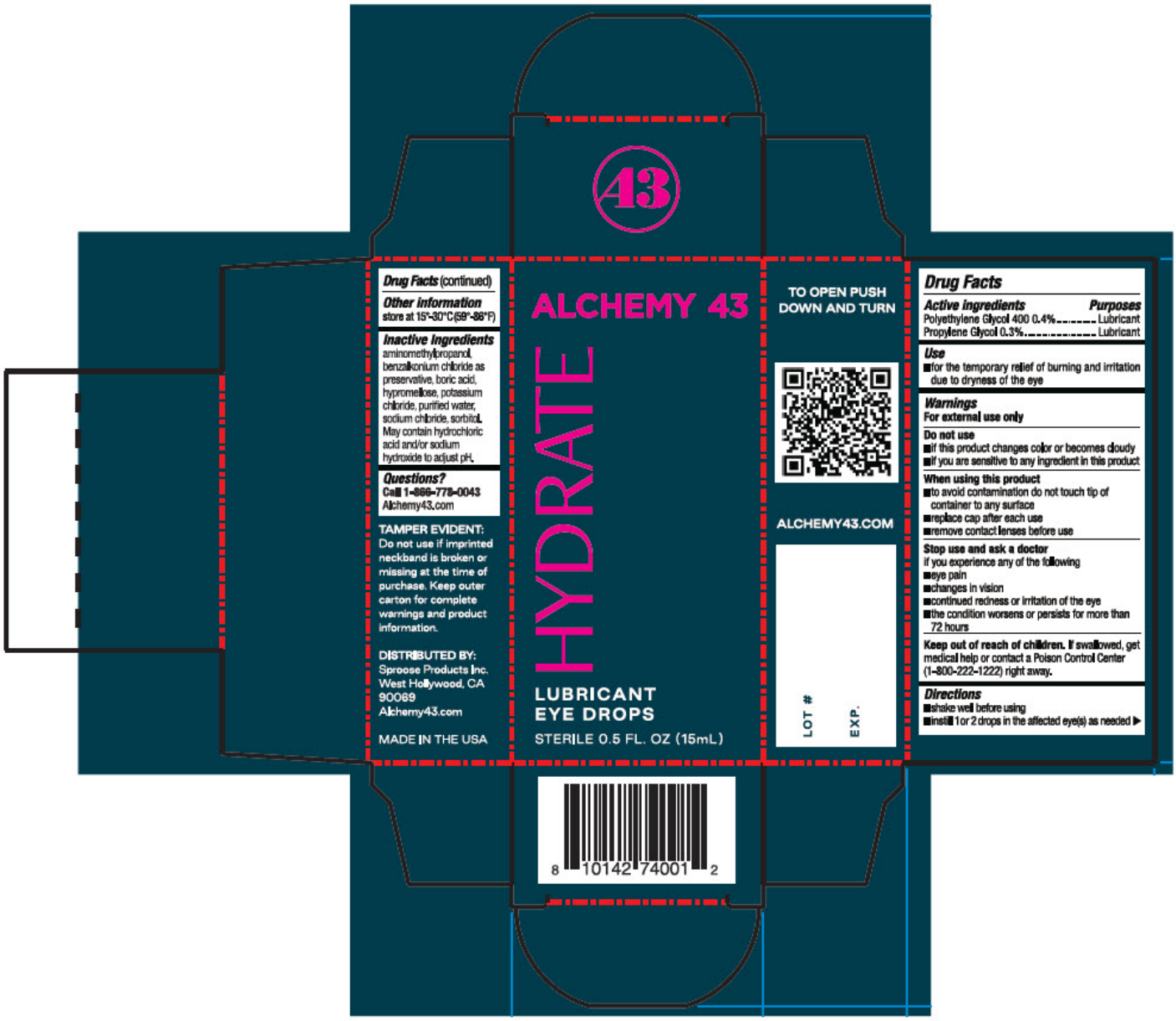
PRINCIPAL DISPLAY PANEL - 15 mL Bottle Carton

ALCHEMY 43

HYDRATE

LUBRICANT
EYE DROPS

STERILE 0.5 FL. OZ (15mL)



HYDRATE LUBRICANT

polyethylene glycol 400 and propylene glycol solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83716-6000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	0.004 g in 1 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	0.003 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SORBITOL (UNII: 506T60A25R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83716-6000-1	1 in 1 CARTON	09/01/2023	
1		15 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M018	09/01/2023	

Labeler - Sproose Products Inc. (119019937)

Establishment

Name	Address	ID/FEI	Business Operations
Samson Pharmaceuticals Inc.		088169581	MANUFACTURE(83716-6000)

Revised: 3/2024 Sproose Products Inc.