SCENTED HAND SANITIZER- scented hand sanitizer gel
Beacon Promotions, Inc.

Scented Hand Sanitizer

Drug Facts

Ethyl alcohol 63.5

Put enough product in your palm to cover your hands and rub together briskly until dry

Purpose

Antimicrobial

Use

Hand sanitizer to help decrease bacteria on the skin

Warnings

Flammable. Keep away from fire or flame.

■ For external use only.

■ When using this product do notuse in or near the eyes. In case of contact, rinse eyes thoroughly with water.

■ Stop use andask a doctor if irritation or rash appears and lasts.

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

Keep out of reach of children

Keep out of reach of children

Directions

■ Put enough product in your palm to cover your hands and rub together briskly until dry

■ Children under 6 years of age should be supervised when using this product

Drug Facts

Ethyl alcohol 63.5 w/w

Inactive ingredients

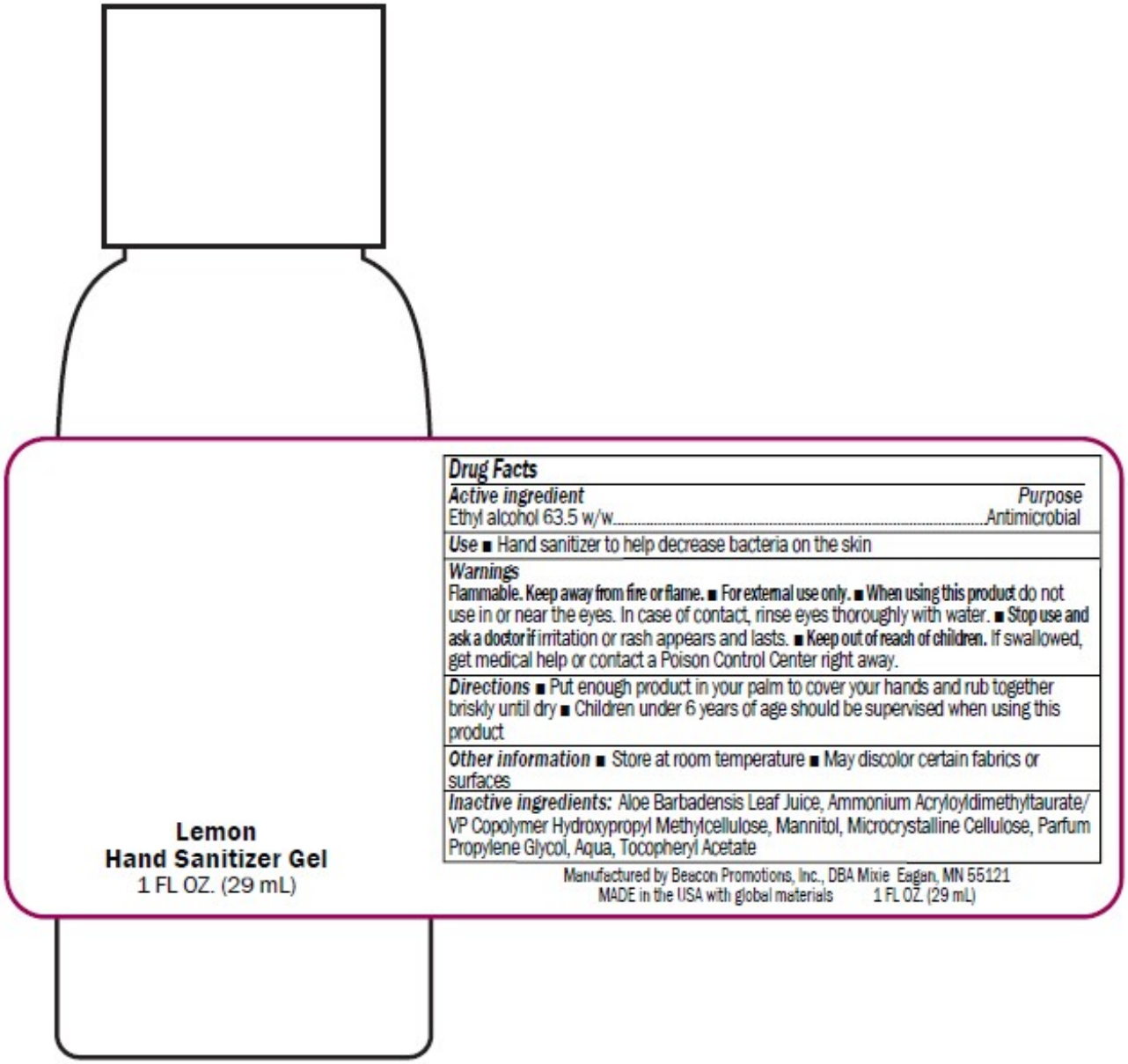
Aloe Barbadensis Leaf Juice, Ammonium Acryloyldimethyltaurate/ VP Copolymer Hydroxypropyl Methylcellulose, Mannitol, Microcrystalline Cellulose, Parfum, Propylene Glycol, Aqua, Tocopheryl Acetate

Other

■ Store at room temperature

■ May discolor certain fabrics or surfaces

Principal Display Panel



SCENTED HAND SANITIZER

scented hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70445-608
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	63.5 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70445-608-01	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
2	NDC:70445-608-02	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
3	NDC:70445-608-03	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
4	NDC:70445-608-04	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
5	NDC:70445-608-05	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
6	NDC:70445-608-06	240 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/20/2024	
7	NDC:70445-608-07	300 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/20/2024	
8	NDC:70445-608-08	480 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
9	NDC:70445-608-09	480 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/20/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	08/20/2024	

Labeler - Beacon Promotions, Inc. (119056382)

Establishment

Name	Address	ID/FEI	Business Operations
Beacon Promotions, Inc.		119056382	manufacture(70445-608)

Revised: 8/2024 Beacon Promotions, Inc.