UNSCENTED SANITIZER- unscented sanitizer gel
Beacon Promotions, Inc.

Unscented 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

Antimicrobial

Warnings

Warnings
Flammable. Keep away from fire or flame.

■ For external use only.

■ When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

■ Stop use and ask 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

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

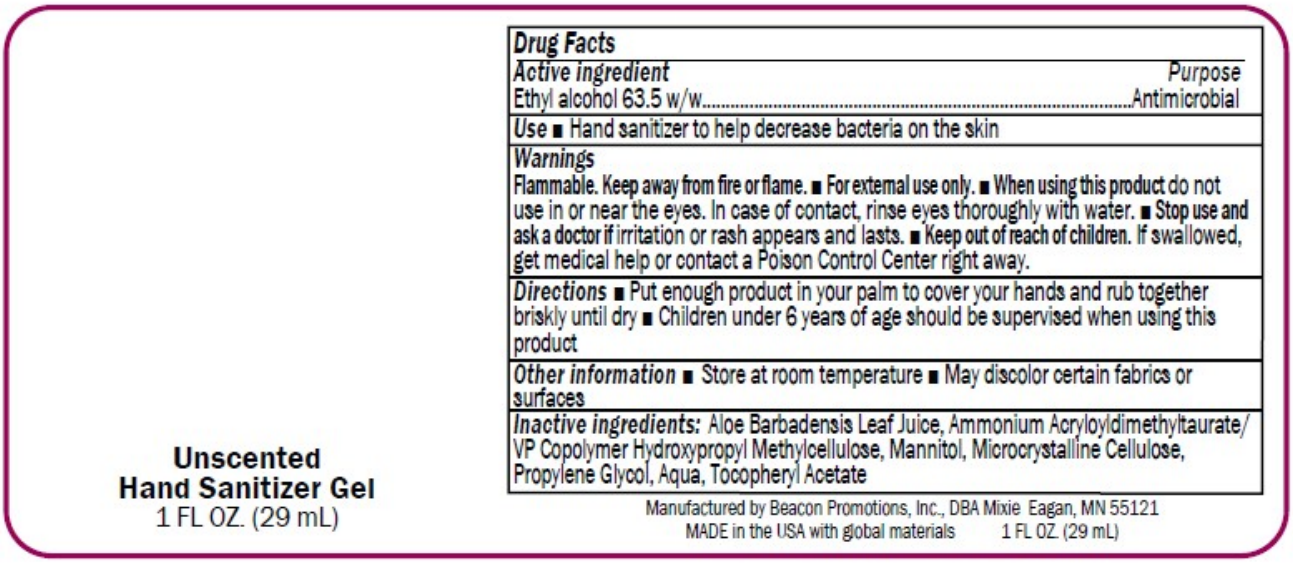
Inactive Ingredients

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

Other Information

- Store at room temperature
- May discolor certain fabrics or surfaces

Principal Display Panel



UNSCENTED SANITIZER

unscented sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70445-609
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**

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70445-609-01	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
2	NDC:70445-609-02	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
3	NDC:70445-609-03	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
4	NDC:70445-609-04	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
5	NDC:70445-609-05	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
6	NDC:70445-609-07	300 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/20/2024	
7	NDC:70445-609-06	240 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/20/2024	
8	NDC:70445-609-08	480 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/20/2024	
9	NDC:70445-609-09	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(70445-609)

Revised: 8/2024 Beacon Promotions, Inc.