SANSPRAY- antibacterial sanitizer spray spray
Beacon Promotions Inc

Alcohol Sanitizer Spray

Drug facts

Ethyl Alcohol (62%)

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Spray enough product on hands to cover
all surfaces. Rub hands together until dry.

Inactive ingredients

Deionized Water, Glycerin, Propylene Glycol, Tocopherol, Fragrance, Aloe Vera.

Purpose

Antiseptic

Purpose

Anticeptic

Directions

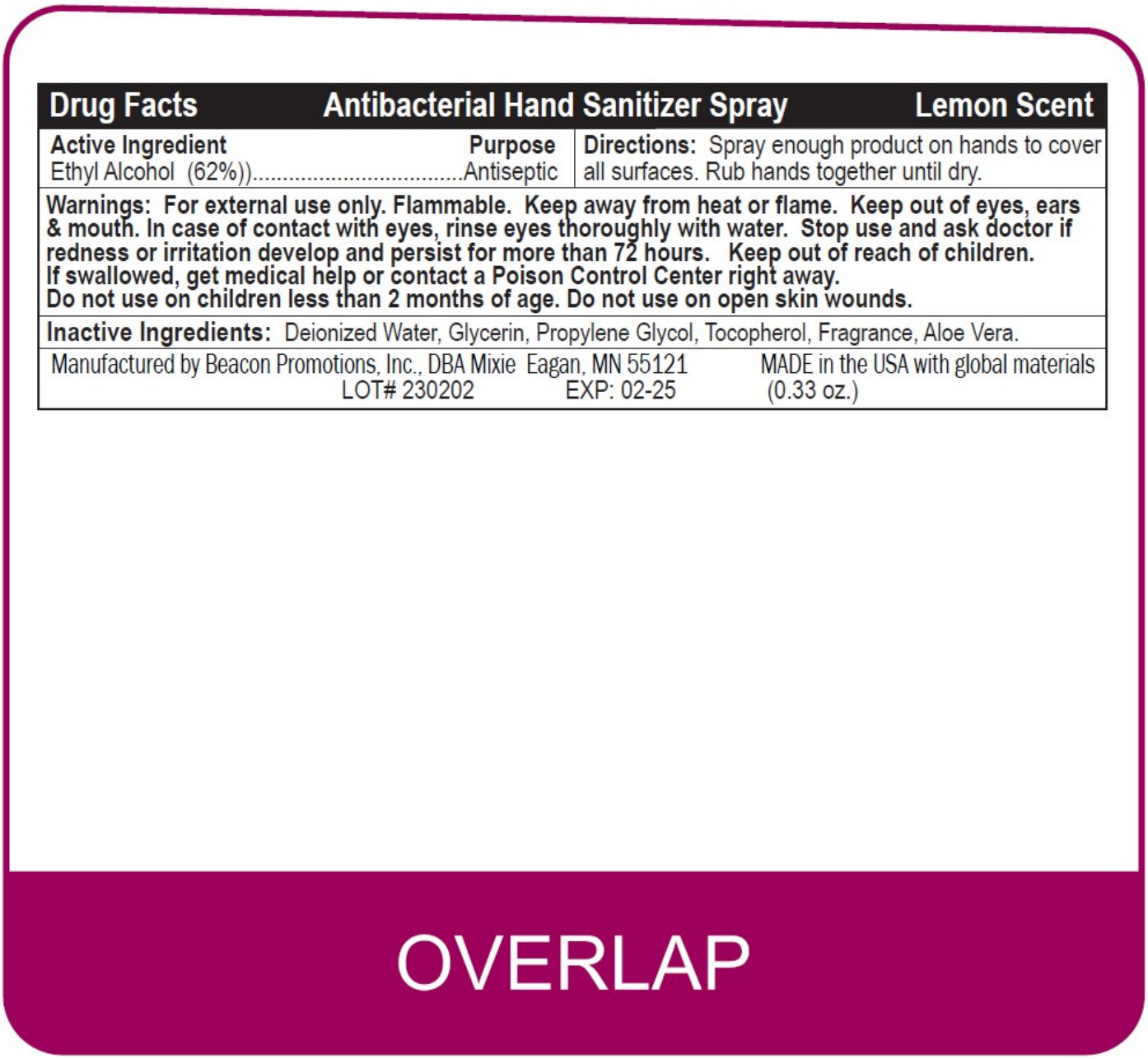
Spray enough product on hands to cover
all surfaces. Rub hands together until dry.

Warnings

For external use only. Flammable. Keep away from heat or flame. Keep out of eyes, ears & mouth. In case of contact with eyes, rinse eyes thoroughly with water. Stop use and ask doctor if
redness or irritation develop and persist for more than 72 hours. Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center right away.
Do not use on children less than 2 months of age. Do not use on open skin wounds.

Keep out of reach of children

Keep out of reach of children



SANSPRAY

antibacterial sanitizer spray spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70445-605
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE (UNII: V5VD430YW9)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70445-605-01	9.76 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/04/2024	
2	NDC:70445-605-02	19.8 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/04/2024	
3	NDC:70445-605-03	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/04/2024	
4	NDC:70445-605-04	120 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/04/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/04/2024	

Labeler - Beacon Promotions Inc (119056382)

Registrant - Beacon Promotions Inc. (119056382)

Establishment

Name	Address	ID/FEI	Business Operations
Beacon Promotions Inc.		119056382	manufacture(70445-605)

Revised: 1/2024 Beacon Promotions Inc