# ANTIBACTERIAL HAND SANITIZER- ethyl alcohol gel ASC Marketing Ltd.

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#### 73145-009, ASC ANTIBACTERIAL HAND SANITIZER

#### Active ingredient

Ethyl alcohol 70%

#### **Purpose**

Antiseptic

#### Uses

For handwashing to decrease bacteria on the skin.

### Warnings

**Flammable. Keep away from fire and flame.** For external use only. Do not use in eyes. In case of contact rinse eyes thoroughly with water. **Discontinue use** if irritation or redness develop. If condition persists for more than 72 hours consult a doctor.

## Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Centre right away.

#### **Directions**

Wet hands thoroughly for with product and allow to dry without wiping.

## Inactive Ingredients

Water, Glycerin, Propylene Glycol, Fragrance, Aloe Barbadensis Leaf Juice, Maltodextrin.

#### INSTANT HAND SANITIZER SPRAY

Drug Facts	Drug Facts Continued	
Active Ingredients Purpose Ethyl Alcohol 70%	Directions: Wet hands thoroughly with product and allow to dry without wiping.	
Use - For hand washing to decrease bacteria on the skin.	Inactive Ingredients: Water, Glycerin, Propylene Glycol, Fragrance, Aloe Barbadensis Leaf Juice, Maltod extrin.	
water. Discontinue use if irritation and redness develop. If condition persists for more than 72 hours consult a doctor. Keep out of reach of children. If	Made in China for: (L) 8295 ASC Marketing 7615 Othello Ave, Unit H-I San Diego, CA 92111	
	8ml/0.27 fl oz. Labeled in USA	

## ANTIBACTERIAL HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73145-009
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	<b>Basis of Strength</b>	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			

GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:73145- 009-01	8 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/16/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	11/16/2020	

## Labeler - ASC Marketing Ltd. (117025198)

Revised: 4/2025 ASC Marketing Ltd.