INSTANT HAND SANITIZER- alcohol spray ASC Marketing LTD

Instant Hand Sanitizer

Drug Facts

Active Ingredients

Ethyl Alcohol 70%

Purpose

Antiseptic

Use

For hand washing to decrease bacteria on the skin.

Warnings

For external use only. Flammable. Keep away from fire and flame.

Do not use

in the eyes. In case of contact, rinse eyes thoroughly with 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 swallowed get medical help or contact Poison Control Center right away.

Directions:

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

Inactive Ingredients:

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

Package Labeling:

HSSP 8ML

INSTANT HAND SANITIZER SPRAY

Prug Facts Drug Facts Continued		
	Directions: Wet hands thoroughly with product and allow to dry without wiping.	
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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 swallowed, get medical help or contact a Poison Control Center right away.	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	

Package Labeling:10ml

HSBTLCC

INSTANT HAND SANITIZER SPRAY

Drug Facts	Drug Facts Continued	
Active IngredientsPurposeEthyl Alcohol 70%Antiseptic	Directions: Wet hands thoroughly with product and allow to dry without wiping.	
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	Net Wt. 0.33 FL OZ (10 ml) Labeled in USA	

INSTANT HAND SANITIZER

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73145-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
MALTODEXTRIN (UNII: 7CVR7L4A2D)		

ı	Packaging			
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date
:	NDC:73145- 010-01	8 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/21/2020	
:	NDC:73145- 010-02	10 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/23/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	11/21/2020	

Labeler - ASC Marketing LTD (117025198)

Revised: 12/2023 ASC Marketing LTD