ANTISEPTIC HAND SANITIZER- alcohol spray MiracleCorp Products

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Antiseptic Hand Sanitizer

Drug Facts

Active Ingredient[s]

Alcohol 80% v/v

Purpose

Antiseptic

Use[s]

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wound

When using this product keep out of eyes, ears and mouth. In case of contact with eyes, rinse throughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

• Store between 15-30C (59-86F)

Inactive Ingredients

Hydrogen Peroxide, Glycerol, Distilled Water

PRINCIPAL DISPLAY PANEL - 103.5 ml Bottle Label

Alcohol Antiseptic 80% Topical Solution

Antiseptic Hand Sanitizer Non-Sterile Solution

Made in Dayton, Ohio by Your Friends at

Buckeye™ VODKA

buckeyevodka.com

MiracleCorp Premium Pet Products miraclecorp.com

3.5 fl oz | 103.5 ml

DO NOT STORE ABOVE 86F SAMPLE - NOT FOR SALE NOT FOR CONSUMPTION

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ANTISEPTIC HAND SANITIZER

alcohol spray

Product Information

Product Type		HUMAN OTC DRUG Item (Code (S	ource)	NDC:61429-005
Route of Administ	ration	TOPICAL			
Active Ingredie	ent/Active	Moiety			
Ingredient Name			Ba	sis of Strength	Strength
Alcohol (UNII: 3K99	lcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M) Alcohol			ol	80 mL in 100 mL
Inactive Ingred	ients				
Ingredient Name					Strength
Hydrogen Peroxide	(UNII: BBX	060 AN9 V)			
Glycerin (UNII: PDC6A3C0OX)					
Glycerin (UNII: PDC Water (UNII: 059QF					
Water (UNII: 059QF					
		Package Description		Marketing Star Date	t Marketing End Date
Water (UNII: 059QF Packaging # Item Code NDC:61429,005	0KOOR)	Package Description 1 BOTTLE, PLASTIC; Type 0: Not a Combinat	ion (U
Water (UNII: 059QF Packaging # Item Code 1 NDC:61429-005-20 NDC:61420-005	0KO0R) 103.5 mL in Product		tion	Date	0
Water (UNII: 059QF Packaging # Item Code 1 NDC:61429-005- 20 NDC:61429-005-	0KO0R) 103.5 mL in Product 473.18 mL i	1 BOTTLE, PLASTIC; Type 0: Not a Combinat	tion	Date	J
Water (UNII: 059QF Packaging Item Code NDC:61429-005-20 NDC:61429-005-30	0KO0R) 103.5 mL in Product 473.18 mL i Product	a 1 BOTTLE, PLASTIC; Type 0: Not a Combinat	tion	Date	U
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Water (UNII: 059QF Packaging Item Code NDC:61429-005-20 NDC:61429-005-30	0KOOR) 103.5 mL in Product 473.18 mL i Product hformat	in 1 BOTTLE, PLASTIC; Type 0: Not a Combinat in 1 BOTTLE, PLASTIC; Type 0: Not a Combina ion Application Number or Monograph Cita	tion Ma	Date 04/20/2020 04/20/2020	Date

Labeler - MiracleCorp Products (604479543)

Establishment			
Name	Address	ID/FEI	Business Operations
MiracleCorp Products		604479543	MANUFACTURE(61429-005)

Revised: 4/2020

MiracleCorp Products