#### 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

238910\_041320

# Alcohol Antiseptic 80% Topical Solution

# Antiseptic Hand Sanitizer Non-Sterile Solution

Made in Dayton, Ohio by Your Friends at



buckeyevodka.com

3.5 fl oz | 103.5 ml DO NOT STORE ABOVE 86F SAMPLE - NOT FOR SALE NOT FOR CONSUMPTION





	-	
Drug Facts Active Ingre Alcohol 80%	edient[s] 5 v/v	<i>Purpose</i> Antiseptic
	zer to help reduce bacteria use when soap and water	
Warnings For externa	l use only. Flammable. Kee	p away from heat or flame
<ul> <li>Do not use</li> <li>in children</li> <li>on open sl</li> </ul>	less than 2 months of age kin wound	
	this product keep out of e	
	d ask a doctor if irritation of serious condition.	or rash occurs. These may
	reach of children. If swallo bison Control Center right a	
together u	until dry.	over all surfaces. Rub hands ge when using this product
<ul> <li>Supervise to avoid st</li> <li>Other Inform</li> <li>Store betw</li> <li>Inactive Ing</li> </ul>	mation veen 15-30C (59-86F)	
Inactive Ing	redients Hydrogen Peroxid	e, Glycerol, Distilled Water

# 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)					
<b>Glycerin</b> (UNII: PDC <b>Water</b> (UNII: 059QF					
Water (UNII: 059QF					
		Package Description		Marketing Star Date	t Marketing End Date
Water (UNII: 059QF Packaging # Item Code NDC:61429,005	0KOOR)	<b>Package Description</b> 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	<b>Date</b>	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	<b>Date</b>	U
Water (UNII: 059QF         Packaging         #       Item Code         1       NDC:61429-005-20         2       NDC:61429-005-30	0KO0R) 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	ation (	Date 04/20/2020 04/20/2020	Date
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 <b>hformat</b>	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	<b>Business Operations</b>
MiracleCorp Products		604479543	MANUFACTURE(61429-005)

Revised: 4/2020

MiracleCorp Products