DONE HAND SANITIZER ISOPROPYL ALCOHOL- is opropyl alcohol gel Olein Recovery Corporation

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.

Done Hand Sanitizer Isopropyl Alcohol spray

Drug Facts

Active Ingredient

Isopropyl Alcohol 75%

Purpose

Antiseptic

Uses

Health care personnel hand rub to help reduce bacteria that potentially can cause disease.

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 wounds

When using this product

Keep out of eyes, ears and mouth. In case of contact with eyes, rinse eyes thoroughly 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

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 59°-86°F)
- Avoid freezing and excessive heat above 104°F

Inactive Ingredients

Carbomer, Water, fragrance

Package Labeling:



REFRESHING SPRAY

Hand Sanitizer

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*Effective at eliminating many common harmful germs and bacteria in as little as 15 seconds.



10 OZ (296 mL) 283 g

PART NO. DON-301-10



MANUFACTURED BY: OLEIN RECOVERY, CORP., YABUCOA PR 00767

DONE HAND SANITIZER ISOPROPYL ALCOHOL

isopropyl alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:77142-005

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL UNII:ND2M416302)

ISOPROPYL
ALCOHOL

ISOPROPYL
ALCOHOL

in 1 mL

Inactive Ingredients

Ingredient Name

CARBOMER HO MO PO LYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)

WATER (UNII: 059 QF0 KO0R)

Packaging

l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC ·77142-005-00	296 mL in 1 BOTTLE: Type 0: Not a Combination Product	0.4/21/20.20	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/21/2020	

Labeler - Olein Recovery Corporation (188543446)

Revised: 5/2020 Olein Recovery Corporation