ANTISEPTIC HAND SANITIZER GEL 128 OZ- alcohol gel
Infinity Global, Inc.

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.

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (80%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
b. Glycerol (1.45% v/v).
c. Hydrogen peroxide (0.125% v/v).
d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Alcohol 72% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

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 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 Center right away.

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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.

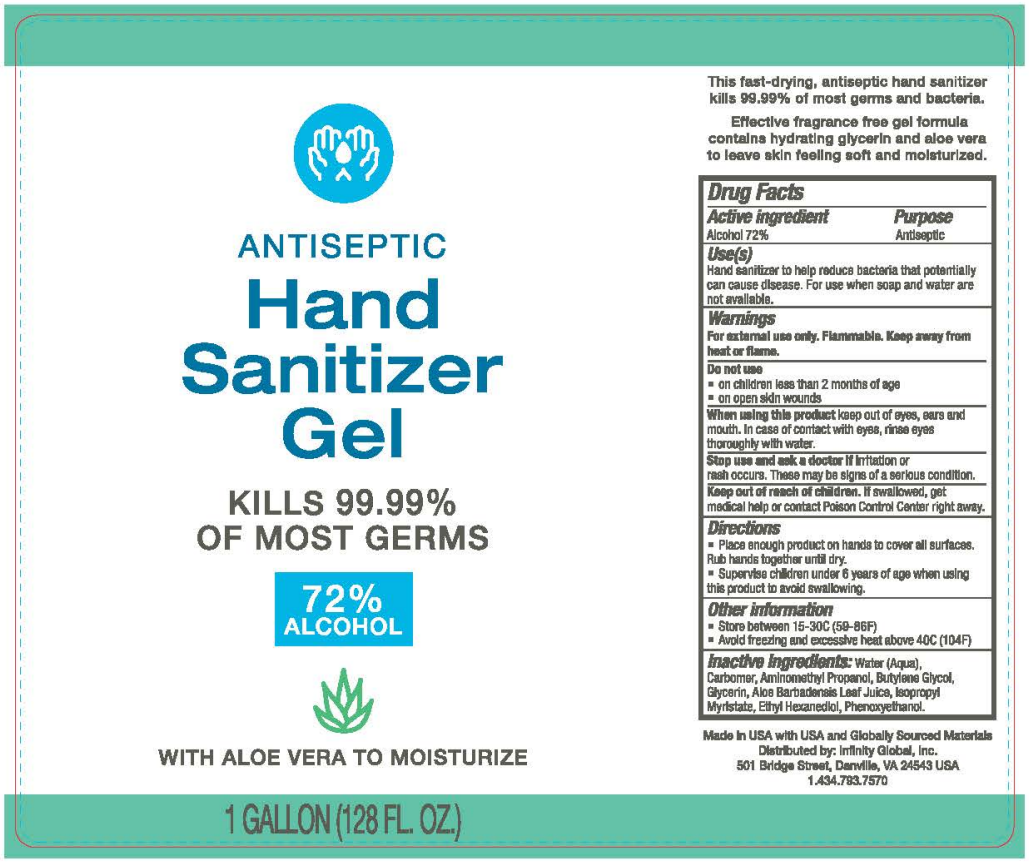
Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Water, Carbomer, Aminomethyl Propanol,Butylene Glycol, Glycerin, Aloe Barbadensis Leaf Juice, Isopropyl Myristate, Ethyl Hexanediol, Phenoxyethanol

Package Label - Principal Display Panel



Customer: **WORMSER**
Folder: **WORMSER•13-67•artwork**
File: **Infinity_Gallon_HandSanitizerLabel_NW_V.ai**
SYSPRO#: **TBD** PRECISION#: **TBD**
SIZE: **5" x 6" w/.125" cr**
JOB#: **TBD**
UNWIND: **TBD**
SUBSTRATE: **TBD** PREPARED BY: **JR**
DATE: **5/28/2020** PROOF #: **1**

Color Guide:
- dieline does not print
- deco area does not print
- PMS 7479
- PMS 33385
- PMS 306
- PMS 307
- PMS 418

PRECISION LABEL CORPORATION
50 Marcus Blvd.
Hauppauge, NY 11788
tel 631.293.4200
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IMPORTANT: Please review proof for typographical errors, format, style and color breaks. This is a representation of the desired color. Please reference PANTONE® color guide for a more accurate color.

PLEASE NOTE: Before proceeding, all proofs and revised proofs MUST be signed and returned.

☐ Approved ☐ Not Approved

NDC: 78679-001-01

ANTISEPTIC HAND SANITIZER GEL 128 OZ

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78679-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	72 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	0.01 mL in 100 mL
WATER (UNII: 059QF0KO0R)	27.65597 mL in 100 mL
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.06303 mL in 100 mL
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	0.01 mL in 100 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.0098 mL in 100 mL
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	0.001 mL in 100 mL
ETHOHEXADIOL (UNII: M9JGK7U88V)	0.00015 mL in 100 mL
PHENOXYETHANOL (UNII: HIE492ZZ3T)	0.00005 mL in 100 mL
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	0.25 mL in 100 mL

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:78679-001-01	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/08/2020	

Labeler - Infinity Global, Inc. (609927566)

Registrant - Nu-World Corporation (628045858)

Establishment

Name	Address	ID/FEI	Business Operations
Nu-World Corporation		628045858	manufacture(78679-001)

Revised: 6/2020 Infinity Global, Inc.