

USANA HAND SANITIZER- alcohol liquid
USANA Health Sciences, 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.

USANA Hand Sanitizer

Drug Facts

Active Ingredient(s)

Alcohol 70% v/v

Purpose

Antiseptic

Uses

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

• On open skin wounds. • In children less than 2 months of age.

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.

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 15-30°C. (59- 86°F) • Avoid freezing and excessive heat above 40°C. (104°F)

Inactive ingredients:

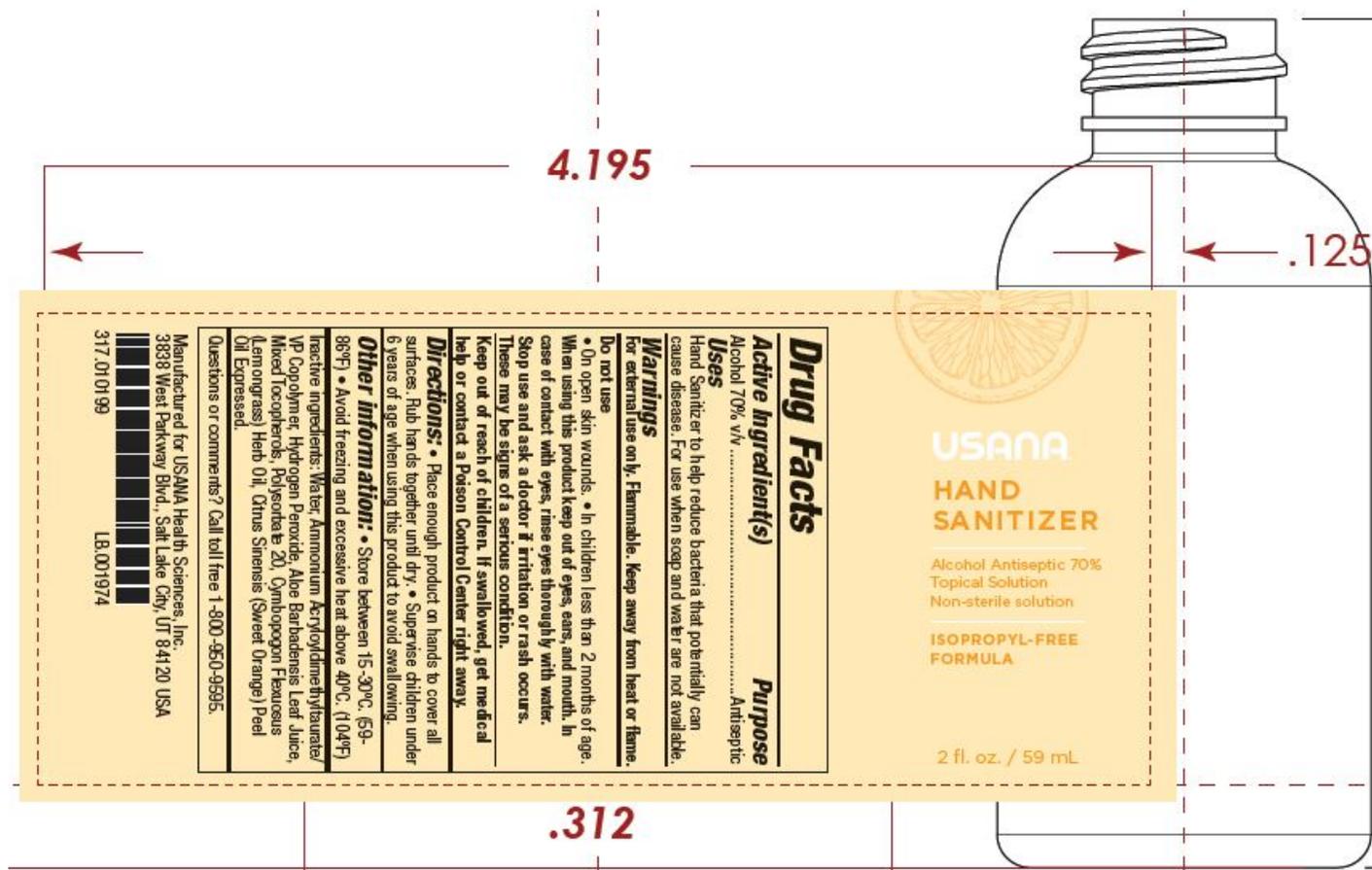
Water, Ammonium Acryloyldimethyltaurate/ VP Copolymer, Hydrogen Peroxide, Aloe Barbadensis

Leaf Juice, Mixed Tocopherols, Polysorbate 20, Cymbopogon Flexuosus (Lemongrass) Herb Oil, Citrus Sinensis (Sweet Orange) Peel Oil Expressed.

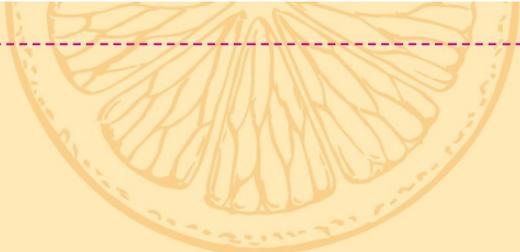
Questions or comments?

Call toll free 1-800-950-9595.

Package Labeling: 59ml



Package Labeling: 240ml



Drug Facts

Active Ingredient(s) **Purpose**

Alcohol 80% v/vAntiseptic

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Manufactured exclusively for USANA Health Sciences, Inc. by Dynamic Blending Specialists Orem, Utah



317.010100

LB.001970

USANA

HAND SANITIZER

Alcohol Antiseptic 80%
Topical Solution
Non-sterile solution

ISOPROPYL-FREE FORMULA

8 FL. oz. / 240 mL

USANA HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:5186 1-070
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: 059QF0K00R)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE (UNII: KBC00G95HI)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

TOCOPHEROL (UNII: R0ZB2556P8)

POLYSORBATE 20 (UNII: 7T1F30V5YH)

CYMOPOGON CITRATUS LEAF (UNII: 06JMS448M5)

ORANGE OIL (UNII: AKN3KSD11B)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51861-070-01	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/15/2020	
2	NDC:51861-070-02	240 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/15/2020	

Labeler - USANA Health Sciences, Inc. (804413250)

Establishment

Name	Address	ID/FEI	Business Operations
Dynamic Blending Specialists, Inc.		085704438	manufacture(51861-070)

Revised: 5/2020

USANA Health Sciences, Inc.