

**SANITIZE ASAP HAND SANITIZER- ethyl alcohol gel**  
**New World Holdings, 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.*

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**SANITIZE ASAP HAND SANITIZER**

***Drug Facts***

***Active ingredient:***

Ethyl Alcohol Denatured.....80%

***Purpose:***

Antiseptic

***Inactive Ingredients:***

Deionized Water, Hydroxypropyl Cellulose, Glycerin, Hydrogen Peroxide.

**Uses:** to decrease bacteria on the skin that could interfere with good health, recommended for repeated use.

**Directions:** wet the hands thoroughly with the product and rub hands until dry.

**Keep out of reach of children, keep out of the eyes, avoid contact with broken skin, if swallowed get medical help or contact a poison control center.**

**Flammable, keep away from heat and flame.**

**ANTI-BACTERIAL**

**KILLS BACTERIA!**

**MADE IN USA**

**LABORATORY CRAFTED - ANTI-BACTERIAL BLEND**

**FROM THE RESEARCH LABORATORIES OF NWH-USA**

**BE SAFE**

**BE CAUTIOUS**

These statements have not been evaluated by the Food and Drug Administration.

This product is not intended to diagnose, treat cure or prevent any disease.

**SANITIZEASAP.COM**

New World Cosmetics and Medicinals.

1080 Holland Drive. Suite 1,

Boca Raton, Florida 33487

**Packaging**

ANTI-BACTERIAL • ANTI-BACTERIAL • ANTI-BACTERIAL



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**HAND SANITIZER  
GEL**

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16FL OZ (473 ML)

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**NDC 73871-378-16**

**SANITIZE ASAP HAND SANITIZER**

ethyl alcohol gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:73871-378
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.8 mL in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73871-378-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

**Labeler** - New World Holdings, Inc. (081183610)

Revised: 5/2020

New World Holdings, Inc.