

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.

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

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73871-378
Route of Administration	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.