

SANITIZE ASAP HAND SANITIZER- isopropyl alcohol spray
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 SPRAY

Drug Facts

Active ingredient:

Isopropyl Alcohol 70%

Purpose

Antiseptic

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

Flammable, keep away from heat and flame.

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.

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

Inactive Ingredients:

Alcohol, Deionized water, Hydroxypropyl guar, Eucalyptus globulus

ANTI-BACTERIAL

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

KILLS BACTERIA

MADE IN USA

LABORATORY CRAFTED - ANTI-BACTERIAL BLEND FROM THE RESEARCH
LABORATORIES OF NWH.

Packaging

ANTI-BACTERIAL • ANTI-BACTERIAL • ANTI-BACTERIAL



**BE SAFE
BE CAUTIOUS**

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**HAND SANITIZER
SPRAY**

**KILLS
BACTERIA!**

**MADE IN
USA**

8FL OZ (236 ML)

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NDC 73871-192-24

SANITIZE ASAP HAND SANITIZER

isopropyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73871-192
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
GUARAPROLOSE (3500 MPAS AT 1%) (UNII: 3A1I7376TC)	
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73871-192-24	236 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/31/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/31/2020	

Labeler - New World Holdings, Inc. (081183610)

Establishment

Name	Address	ID/FEI	Business Operations
New World Holdings, Inc.		081183610	manufacture(73871-192)

Revised: 3/2020

New World Holdings, Inc.