

HAND SANITIZER- alcohol gel

Amyris, 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.

Olika Fragrance Free

Active Ingredient(s)

Alcohol 65% 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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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

Aqua

Glycerin

Aloe Barbadensis Leaf Extract

Sodium Hyaluronate

Package Label - Principal Display Panel

ULTRA HYDRATION
WITH HYALURONIC ACID

Many hand sanitizers dry out your hands, causing uncomfortable cracking or peeling that could expose you to more harmful bacteria. OLIKA's uses moisture-retaining hyaluronic acid and aloe to quench dry hands with long-lasting hydration while removing 99.9% of germs. Find out what's different about OLIKA.



Cruelty-free



Conscious ingredients



Vegan formulation



15x More uses than similarly sized gels

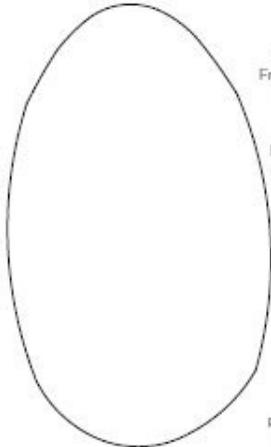
f t i s
@OLIKALIFE



WARNING!
CHOKING HAZARD:
This is not a toy. Not suitable for children under 3 years.



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Patent Pending
Designed in California
Fragrance Made in USA
Assembled in China

Manufactured for:
AMYRIS Inc.
Emeryville, CA 94608
Not tested on animals.
NDC#
XXXXX-XXX-XX



OLIKA®

ULTRA HYDRATING HAND SANITIZER



24 HR HYDRATION
WITH HYALURONIC ACID

500+
SPRAYS

FRAGRANCE
FREE

EFFECTIVE
AGAINST
99.9%
OF GERMS

30 mL
(1.0 fl oz)

MADE OLIKA CLEAN™

CONTAINS **65%**
ETHYL ALCOHOL

Drug Facts

Active Ingredient Ethyl Alcohol 65% v/v
Purpose Antiseptic

Use Helps reduce bacteria on skin.

Warnings

For external use only. Flammable, keep away from fire or flame.

Do not use on children less than 12 months of age on open skin wounds.

When using keep out of eyes, ears, and mouth. In case of contact, 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.

Drug Facts (continued)

If swallowed, get medical help or contact a Poison Control Center right away.

Other information

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

Directions

Spray liquid into hands and rub until dry. ■ Supervise children under 6 years of age to prevent swallowing.

Inactive ingredients

Aloe vera (aloe barbadensis) leaf juice, denatonium benzoate*, glycerin, hyaluronic acid, and water.

Comments/Questions?

(866) 446-5452 (866-44-OLIKA)

*Alcohol denaturant. For more info visit: olikalife.com

30 mL NDC: 73517-767-01

HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73517-767
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73517-767-01	30 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	09/01/2021	
2	NDC:73517-767-02	20 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	09/01/2021	
3	NDC:73517-767-03	90 mL in 1 POUCH; Type 0: Not a Combination Product	09/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Amyris, Inc (185930182)**Establishment**

Name	Address	ID/FEI	Business Operations
Taizhou Jingshang Cosmetics Technology Co., Ltd.		550819554	manufacture(73517-767)

Revised: 1/2023

Amyris, Inc