## OLIKA ULTRA HYDRATING HAND SANITIZER, UNSTOPPABLE- 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.

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## OLIKA Ultra Hydrating Hand Sanitizer, Unstoppable

Ethyl Alcohol 65% v/v. Purpose: Antiseptic

Antiseptic, Hand Sanitizer

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available

For external use only. Flammable. Keep away from heat or flame

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.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

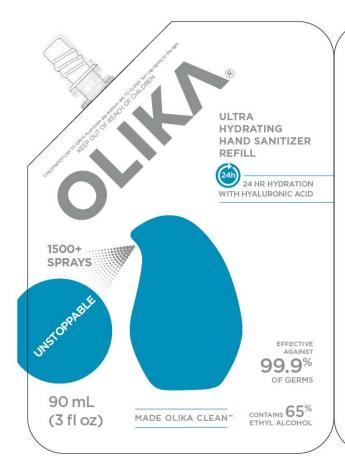
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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.

Store between 15-30C (59-86F)

Avoid freezing and excessive heat above 40C (104F)

water, glycerin, fragrance, aloe vera (aloe barbadensis) leaf juice, hyaluronic acid, denatonium benzoate,



### **ULTRA HYDRATION**

WITH HYALURONIC ACID

Many hand santitzers dry out your hands, causing uncomfortable cracking or peeling that could expose you to more harmful bactera. 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's.



Cruelty-free





OLIKALIFE.COM

Conscious ingredients

More uses than similarly sized gels

## **Drug Facts**

Active Ingredient Alcohol 65% v/v

Purpose Antiseptio

**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.

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

When using keep out of eyes, eas, and mouth in case of contact, mise eyes thoroughly will weller.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serfus condition.

Keep out of track of children. Keep out of reach of children.
If swallowed, get medical help or contact
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Inactive ingredients Water, glycerin, fragrance\*, aloe vera (aloe barbadensis) leaf juice, hyaluronic acid, denatonium benzoate.

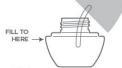
Comments/Questions? (866) 446-5452 (866-44-0LIKA)

formulation

Vegan

This easy to use refill pouch will replenish your OLIKA three or four times for over 1500 sprays.

Simply unscrew the bottom of your OLIKA and fill with sanitizer up to the edge of the reservoir area.

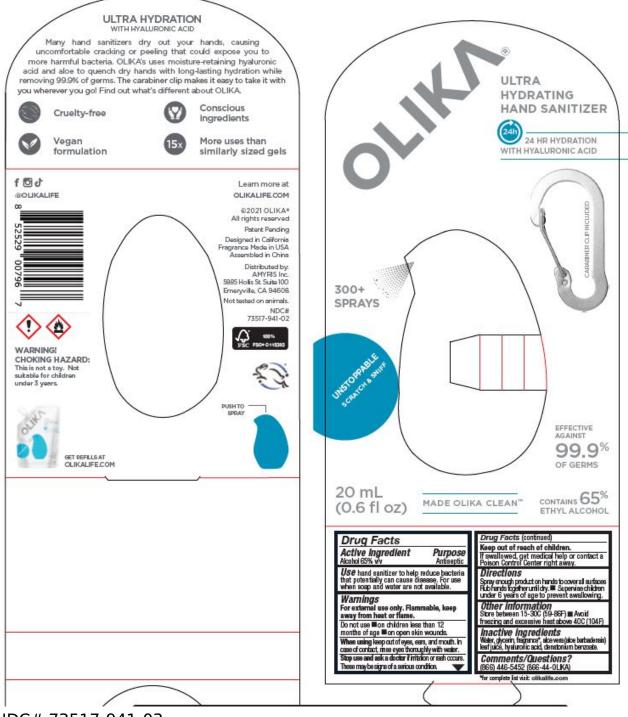


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Designed in California Emeryylle, CA 94608
Fragrance Made in USA
Assembled in China NDC# 73517-941-03





90 ML NDC# 73517-941-03

20 ML NDC# 73517-941-02

# OLIKA ULTRA HYDRATING HAND SANITIZER, UNSTOPPABLE alcohol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73517-941	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL	

Inactive Ingredients				
Ingredient Name	Strength			
GLYCERIN (UNII: PDC6A3C0OX)				
HYALURONATE SODIUM (UNII: YSE9PPT4TH)				
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)				
WATER (UNII: 059QF0KO0R)				
FRAGRANCE 13576 (UNII: 5EM498GW35)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:73517- 941-02	20 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/06/2022	
	2	NDC:73517- 941-03	90 mL in 1 POUCH; Type 0: Not a Combination Product	05/06/2022	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	05/06/2022		

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

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Taizhou Jingshang Cosmetics Technology Co., Ltd.		550819554	manufacture(73517-941)	

Revised: 5/2022 Amyris, Inc