OLIKA ULTRA HYDRATING HAND SANITIZER, BRAVE- 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 Ultra Hydrating Hand Sanitizer, Brave

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.

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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)

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



ULTRA HYDRATION

WITH HYALIVONIC 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

More uses than

similarly sized gels

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

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



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OLIKALIFE.COM

NDC# 73517-951-03





Drug Facts

Active Ingredient Alcohol 65% v/v

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.

away from heat or flame.

Do not use on children less than 12 months of age on children less than 12 months of age on children less than 12 months of age on contain less of contact, rinse eyes throughly will weller. Stop use and ask a doctor if irritation or rash cours. 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.

Directions

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Refill empty bottle.

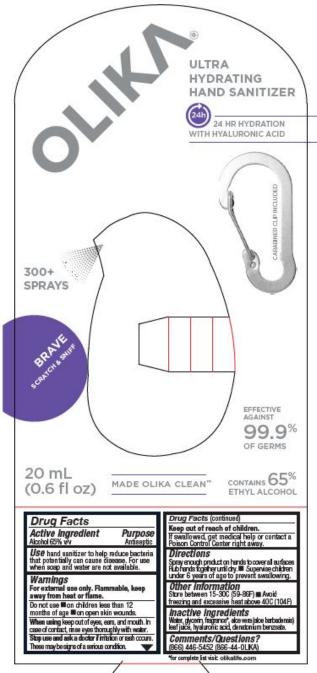
Spray enough product on hands to cover all surfaces. Rub hands together until dry.

Supervise children under 6 years of age to prevent swallowing.

Other information
Stre between 15-30C (59-86F) Makoid
reczing and excessive heat above 40C (104F)
Inactive ingredients
Water, glycerin, fragrance*, aloe vera
(aloe barbadenss) leaf juice, hyaluronic
acid, denatonium benzoate.

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





90 ML NDC# 73517-951-03

20 ML NDC# 73517-951-02

OLIKA ULTRA HYDRATING HAND SANITIZER, BRAVE

alcohol gel

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73517-951		
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			
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)				
HYALURONATE SODIUM (UNII: YSE9PPT4TH)				
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
FRAGRANCE 13576 (UNII: 5EM498GW35)				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73517- 951-02	20 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/09/2022	
2	NDC:73517- 951-03	90 mL in 1 POUCH; Type 0: Not a Combination Product	05/09/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/09/2022	

Labeler - Amyris, Inc (185930182)

Establishment				
Name	Address	ID/FEI	Business Operations	
Taizhou Jingshang Cosmetics Technology Co., Ltd.		550819554	manufacture(73517-951)	

Revised: 5/2022 Amyris, Inc