OLIKA HYDRATING HAND SANITIZER-CLARITY- alcohol gel
Amyris

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 Hydrating Hand Sanitizer-Clarity

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.

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 barbadensis leaf extract, hyaluronic acid, ganoderma lucidum (mushroom) extract, denatonium benzoate

90 ML NDC# 73517-949-01

20 ML NDC# 73517-949-02

OLIKA HYDRATING HAND SANITIZER-CLARITY

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73517-949
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
HYALURONIC ACID (UNII: S270N0TRQY)	
GANODERMA LUCIDUM STEM (UNII: U8PA41532G)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73517-949-01	90 mL in 1 POUCH; Type 0: Not a Combination Product	07/29/2022	
2	NDC:73517-949-02	20 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	07/29/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/29/2022	

Labeler - Amyris (185930182)

Establishment

Name	Address	ID/FEI	Business Operations
Taizhou Jingshang Cosmetics Technology Co., Ltd.		550819554	manufacture(73517-949)

Revised: 7/2022 Amyris