OLIKA HYDRATING HAND SANITIZER, SANDALWOOD- 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 Hydrating Hand Sanitizer- Sandalwood

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

20 mL 73517-737-02

90 mL 73517-737-01







OLIKA HYDRATING HAND SANITIZER, SANDALWOOD

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73517-737

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M) ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M)

Inactive Ingredients				
Ingredient Name	Strength			
GANODERMA LUCIDUM STEM (UNII: U8PA41532G)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
HYALURONIC ACID (UNII: S270N0TRQY)				
GLYCERIN (UNII: PDC6A3C0OX)				
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)				
WATER (UNII: 059QF0KO0R)				
FRAGRANCE 13576 (UNII: 5EM498GW35)				

Packaging							
# Item Code	Package Description	Marketing Start Date	Marketing End Date				
NDC:73517- 737-01	90 mL in 1 POUCH; Type 0: Not a Combination Product	07/29/2022					
NDC:73517- 737-02	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	07/01/2022		
illiai				

Labeler - Amyris, Inc (185930182)

Establishment						
Name	Address	ID/FEI	Business Operations			
Taizhou Jingshang Cosmetics Technology Co., Ltd.		550819554	manufacture(73517-737)			

Revised: 7/2022 Amyris, Inc