ANTI-BACTERIAL HAND SANITIZER WITH GOAT MILK VANILLA BEANalcohol liquid

Creative Consumer Products, 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.

Anti-Bacterial Hand Sanitizer with Goat Milk Vanilla Bean

Drug Facts

Active ingredient

Ethyl alcohol 70%

Purpose

Antiseptic

Use

for hand washing to decrease bacteria on the skin.

Warnings

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

Do not use

in the eyes.

Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

Keep out of reach of children.

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

Directions

Wet hands thoroughly with product and allow to dry without wiping.

Inactive ingredients

Water, Isopropyl Alcohol, Glycerin, Fragrance, Acrylates/C10-30 Alkyl Acrylate

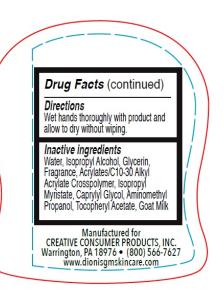
Crosspolymer, Isopropyl Myristate, Caprylyl Glycol, Aminomethyl Propanol, Tocopheryl Acetate, Goat Milk

Package Labeling:









ANTI-BACTERIAL HAND SANITIZER WITH GOAT MILK VANILLA BEAN

alcohol liquid

Product Information

- 1			
1	Product Type	HUMAN OTC DRUG	Item Code (Source)
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roduct Type HUMAN OTC DRUG Item Code (Source) NDC:74028-004

Route of Administration TOPICAL

Active	Ingred	ient/Act	ive Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ISOPROPYL ALCOHOL (UNII: ND2M416302)		
GLYCERIN (UNII: PDC6A3C0OX)		
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)		
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)		
CAPRYLYL GLYCOL (UNII: 00YIU5438U)		
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
GOAT MILK (UNII: XE5K5I4RP7)		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:74028-004- 59	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/28/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/28/2021	

Labeler - Creative Consumer Products, Inc. (079358868)

Revised: 7/2021 Creative Consumer Products, Inc.