MOISTURIZER HAND SANITIZER- benzalkonium chloride solution
Rinati Skin, LLC

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.

RINATI - MOISTURIZING HAND SANITIZER (76849-101)

ACTIVE INGREDIENT

BENZALKONIUM CHLORIDE 0.13%

PURPOSE

ANTIMICROBIAL

USE

FOR HAND SANITIZING TO DECREASE BACTERIA ON THE SKIN.

RECOMMENDED FOR REPEATED USE.

WARNINGS

FOR EXTERNAL USE ONLY.

WHEN USING THIS PRODUCT AVOID CONTACT WITH EYES.

IN CASE OF EYE CONTACT, FLUSH EYES WITH WATER. STOP USE AND ASK A DOCTOR IF IRRITATION, OR REDNESS DEVELOPS OR IF CONDITION PERSISTS FOR MORE THAN 72 HOURS.

KEEP OUT OF REACH OF CHILDREN.

IF SWALLOWED, GET MEDICAL HELP.

DIRECTIONS

SHAKE WELL BEFORE USE.

SPRAY A SMALL AMOUNT OF PRODUCT INTO PALM OF HAND.

RUB THOROUGHLY OVER ALL SURFACES OF BOTH HANDS.

RUB HANDS TOGETHER BRISKLY UNTIL DRY.

INACTIVE INGREDIENTS

WATER, PROPYLENE GLYCOL, LINUM USITATISSIMUM (FLAX) SEED EXTRACT, OCIMUM

TENUIFLORUM LEAF EXTRACT, MELIA AZADIRACHTA LEAF EXTRACT, GLUCONOLACTONE, CITRIC ACID, SODIUM BENZOATE, EUCALYPTUS GLOBULUS OIL

MOISTURIZER HAND SANITIZER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76849-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
FLAX SEED (UNII: 4110YT348C)	
OCIMUM TENUIFLORUM TOP (UNII: 34T63W8ULS)	
AZADIRACHTA INDICA LEAF (UNII: HKY915780T)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76849-101-11	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/04/2020	

Labeler - Rinati Skin, LLC (109530208)

Revised: 10/2022

Rinati Skin, LLC