

**FIND YOUR HAPPY PLACE- sunkissed ocean waves hand sanitizer gel
Conopco Inc. d/b/a/ Unilever**

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.

Find Your Happy Place Sunkissed Ocean Waves Hand Sanitizer

FIND YOUR HAPPY PLACE SUNKISSED OCEAN WAVES HAND SANITIZER - Ethyl Alcohol gel

Find Your Happy Place Sunkissed Ocean Waves Hand Sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol 70 %

Purpose

Antiseptic

Uses

Hand sanitizer to help reduce bacteria on the skin

Warnings

For external use only

Flammable. Keep away from fire or flame. Do not store in car .

When using this product do not use in or near the eyes.

Stop use and ask doctor if irritation or redness develop.

- **Keep out of reach of children except under adult supervision.**

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

Directions

Wet hands thoroughly with product and rub lightly until dry. Do not wipe off or rinse.

Other information

Store below 105°F (40°C). May discolor fabrics or surfaces.

Inactive ingredients

Water (Aqua), Fragrance (Parfum), Glycerin, Isopropyl Alcohol, Butylene Glycol, Carbomer, Aminomethyl Propanol, Isopropyl Myristate, Tocopheryl Acetate, Theobroma Cacao (Cocoa) Seed Butter, Butyrospermum Parkii (Shea) Butter, Tocopherol, Blue 1 (CI 42090), Red 33 (CI 17200).

Questions or comments?

1-800-404-0580

Packaging



Drug Facts	
Active ingredient	Purpose
Ethyl Alcohol (70% v/v)	Antiseptic
Uses Hand sanitizer to help reduce bacteria on the skin	
Warnings	
For external use only	
Flammable. Keep away from fire or flame. Do not store in car.	
When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.	
Stop use and ask doctor if irritation or redness develop. ▶	

FIND YOUR HAPPY PLACE

sunkissed ocean waves hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1830
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	700 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
COCOA BUTTER (UNII: 512OYT1CRR)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
SHEA BUTTER (UNII: K49155WL9Y)	
TOCOPHEROL (UNII: R0ZB2556P8)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-1830-1	59 mL in 1 CONTAINER; Type 0: Not a Combination Product	02/15/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/15/2021	

Labeler - Conopco Inc. d/b/a/ Unilever (001375088)

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