

SUAVE- 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.

Suave Hand Sanitizer

SUAVE HAND SANITIZER - Ethyl Alcohol gel

Suave Hand Sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol 75.6%

Purpose

Antiseptic

Uses

help decrease bacteria on the hands

Warnings

- **For external use only**
- **Flammable. Keep away from fire or flame**
- **Avoid contact with eyes**, in case of contact, rinse eyes thoroughly with water immediately.
- **If irritation develops**, discontinue use and consult a doctor.
- **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

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

Inactive ingredients

Cetyl Lactate, Glycerin, Hydroxypropylcellulose, Lauryl Lactate, Myristyl Lactate, Water (Aqua)

Questions?

Call us at 1-800-782-8301.

Packaging



The image shows the front of a Suave Hand Sanitizer bottle. At the top is the Suave logo in a dark blue wave. Below it, the words "HAND SANITIZER" are printed in large, bold, blue capital letters. At the bottom, an orange banner contains the text "Kills 99.9% of germs". The volume "8 FL. OZ. (236mL)" is printed in white at the very bottom.

EFFECTIVE AT ELIMINATING OVER 99.9% OF MANY COMMON HARMFUL GERMS AND BACTERIA.

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 Empty Before Recycling
how2recycle.info
PLASTIC BOTTLE

THE SUAVE PROMISE:
WE GUARANTEE YOUR FULL SATISFACTION OR WE WILL REPLACE OR REFUND YOUR PURCHASE.

 Unilever

DIST. BY ©UNILEVER
TRUMBULL, CT 06611

QUESTIONS OR COMMENTS?
VISIT WWW.SUAVE.COM
OR CALL US AT 1-800-782-8301.
68428122

smartlabel app enabled 



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SUAVE

hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1742
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	756 mg in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
CETYL LACTATE (UNII: A7EVH2RK4O)				
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
LAURYL LACTATE (UNII: G5SU0BFK7O)				
MYRISTYL LACTATE (UNII: 1D822OC34X)				
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-1742-1	236 mL in 1 CONTAINER; Type 0: Not a Combination Product	06/15/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	06/15/2020		

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

Revised: 12/2022

Conopco Inc. d/b/a/ Unilever