FOASU DAILY HAND SANITIZER GEL- alcohol gel HANSCOS Co.,Ltd.

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.

FOASU DAILY HAND SANITIZER GEL

Active ingredient

Ethyl alcohol 70%

Purpose

Antimicrobial

Use

• Hand sanitizer to help reduce bacteria on the skin

Warnings

Flammable. Keep away from fire or flame.

For external use only

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 a doctor if irritation or rash appears and lasts

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Put nough produt in your palm to cover hands and rub hands together briskly until dry.
- Children under 6 years of age should be supervised when using this product

Other information

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces

Inactive ingredients

WATER, GLYCERIN, PEG-60 HYDROGENATED CASTOR OIL, CARBOMER, TRIETHANOLAMINE, CAMELLIA SIMENSIS LEAF EXTRACT, ALOE BARBADENSIS LEAF EXTRACT, MENTHA PIPERITA (PEPPERMINT) OIL, LIMONENE



DAILY HAND SANTIETS GEL



16.9 FL.OZ (500 ml)

	Drug Facts	
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HANSCOS Co.,Ltd.

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www.Hanscos.com

MADE IN KOREA



FOASU DAILY HAND SANITIZER GEL

alcohol gel

Product Information

HUMAN OTC DRUG Product Type NDC:77964-001 Item Code (Source)

Route of Administration TOPICAL

Active Ingredient/Active Moiety

PEPPERMINT OIL (UNII: AV092KU4JH) LIMO NENE, (+)- (UNII: GFD7C86Q1W)

Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 0.7 g in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
PEG-60 HYDROGENATED CASTOR OIL (UNII: 02NG325BQG)				
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)				
TROLAMINE (UNII: 903K93S3TK)				
GREEN TEA LEAF (UNII: W2ZU1RY8B0)				
ALOE VERA LEAF (UNII: ZY81Z83H0 X)				

Packaging					
#	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:77964-001-	20 mL in 1 TUDE: Type 0: Not a Combination Broduct	05/15/2020		

1	01	30 IIIL III 1 100E, 19Pe 0. NOLA COMOMANON FIOUNCE	03/13/2020	
2	NDC:77964-001- 02	50 mL in 1 TUBE; Type 0: Not a Combination Product	05/15/2020	
3	NDC:77964-001- 03	100 mL in 1 TUBE; Type 0: Not a Combination Product	05/15/2020	
4	NDC:77964-001- 04	250 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/15/2020	
5	NDC:77964-001- 05	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/15/2020	

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	05/15/2020			

Labeler - HANSCOS Co.,Ltd. (688494423)

Registrant - HANSCOS Co.,Ltd. (688494423)

Establishment				
Name	Address	ID/FEI	Business Operations	
HANSCOS Co.,Ltd.		688494423	manufacture (77964-001)	

Revised: 5/2020 HANSCOS Co.,Ltd.