INSTANT HAND SANITIZER- instant hand sanitizer gel NANXIONG SINGLONG CHEMICALS 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.

80533-001 INSTANT HAND SANITIZER

Active Ingredient(s)

Ethyl Alcohol 75%v/v

Purpose

Antimicrobial

Use

Hand sanitizer to help reduce bacteria on 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.

Do not use

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Directions

Put enough product in your palm to cover hands and rub hands together briskly until dry Children under 6 years of age should be supervised

Other information

Store below 86°F/30°C May discolor certain fabrics or surfaces

Inactive ingredients

water, Glycerin, Aloe Barbadensis Leaf Extract, Acry1ates/C10-30 Alkyl Acrylate Crosspolymer, Triethanolamine

Package Label - Principal Display Panel



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Manufactured by

MADE IN CHINA

Nanxiong Singlong Chemicals Co.,Ltd. No.3, Development Avenue 2nd Road, Zhuji Industrial Park, Nanxiong City, Shaoguan City, Guangdong Province, China





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60ML

INSTANT HAND SANITIZER

instant hand sanitizer gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:80533-001

TOPICAL **Route of Administration**

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive	Ingredients
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mactive ingredients	
Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
WATER (UNII: 059QF0KO0R)	

TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80533-001-01	8 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:80533-001-03	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
3	NDC:80533-001-04	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
4	NDC:80533-001-02	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
5	NDC:80533-001-06	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
6	NDC:80533-001-05	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
7	NDC:80533-001-09	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
8	NDC:80533-001-08	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
9	NDC:80533-001-07	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - NANXIONG SINGLONG CHEMICALS CO.,LTD. (552128700)

Establishment			
Name	Address	ID/FEI	Business Operations
NANXIONG SINGLONG CHEMICALS CO.,LTD.		552128700	manufacture (80533-001)

Revised: 9/2020 NANXIONG SINGLONG CHEMICALS CO.,LTD.