HAND SANITIZER- alcohol gel ningbo fareast industry co.,ltd

Reference Label Set Id: a68674ba-455a-4d63-e053-2a95a90aed00

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.

alcohol 75% v/v purpose :antiseptic alcohol 70% v/v purpose :antiseptic alcohol 60% v/v purpose :antiseptic

to help reduce bacteria that potentially can cause disease.for use when soap and water are not available for external use only,flammable,keep away from heat or flame

. in children less than 2 months of age .on open skin wounds

when using this product keep out of eyes, ears, and mouth in case of contact with eyes, rinse eyes thorough with water

stop use and ask a doctor if irritation or rash occurs, these may be signs of a serious condition. keep out of reach of children, if swallowed, get medical help or contact a poison control center right away.

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place enough product on hands to cover all surface, rub hands together until dry supervise children under 6 years when using this product to avoid swallowing

package label principal display panel



Active ingredients

Purpose

Ethyl Alcohol 70%

.....Antiseptic

Uses ■ Hand sanitizer to help decrease bacteria on the skin ■ When water, soap & towel are not available. ■ Recommended for repeated use.

Warnings

- For external use only
- Keep away from fire or flame
- Do not apply around eyes. Do not use in ears & mouth.
- When using this product. avoid contact with eyes.
- In case of contact flush eyes with water
- Stop use and ask a doctor it redness or irritation develop and persist for more than 72 hours
- Keep out of reach of children. Children must be supervised in use of this product

Directions

Pump as needed into your palms and thoroughly spread on both hands. Rub into skin until dry

Other Information Store at 20C(88" to 77"F) may discolor fabrices

Inactive ingredients

Water, Glyerin, Carbomer, Hydrogen Peroxide, Trolamine.





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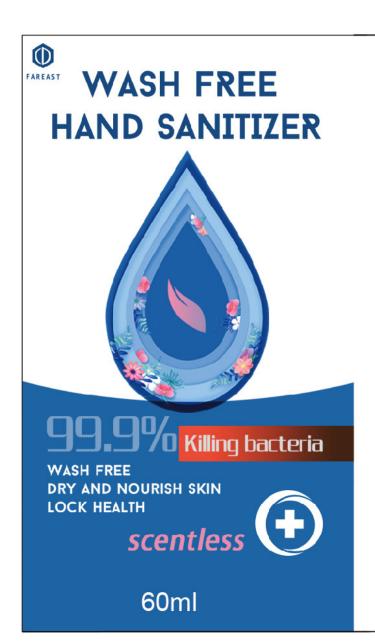
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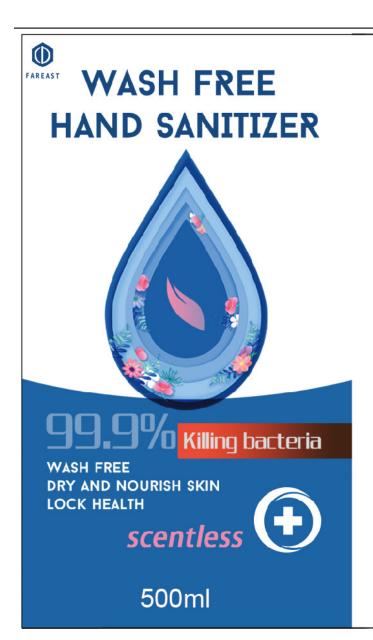
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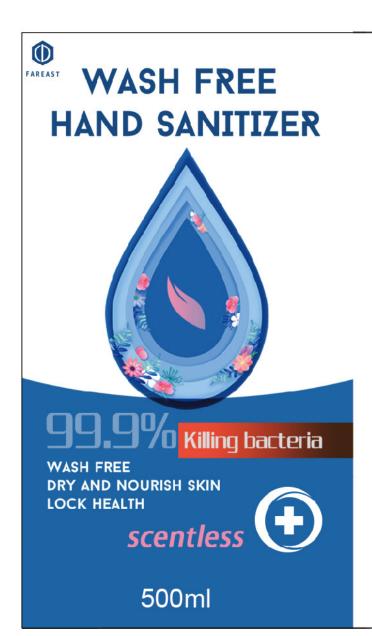
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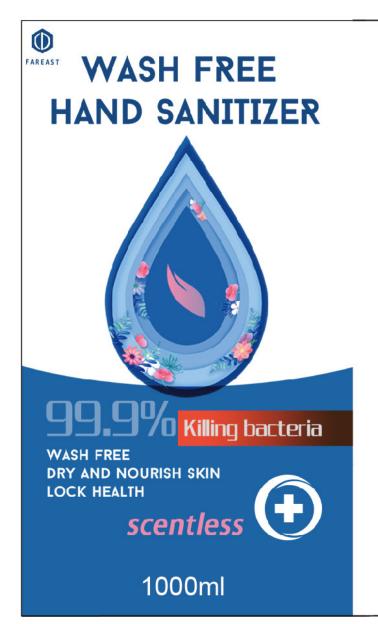
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made in china

75% alcohol concent

1000ml NDC: 78623-375-01

500ml NDC: 78623-375-02

237ml NDC: 78623-375-03

100ml NDC: 78623-375-04

60ml NDC: 78623-375-05

30ml NDC: 78623-375-06

70% alcohol concent

1000ml NDC: 78623-370-01

500ml NDC: 78623-370-02

237ml NDC: 78623-370-03

100ml NDC: 78623-370-04

60ml NDC: 78623-370-05

30ml NDC: 78623-370-06

60% alcohol concent

1000ml NDC: 78623-360-01 500ml NDC: 78623-360-02 237ml NDC: 78623-360-03 100ml NDC: 78623-360-04 60ml NDC: 78623-360-05 30ml NDC: 78623-360-06

HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78623-370
Route of Administration	TOPICAL		

	Active Ingredient/Active Moiety			
ı	Ingredient Name	Basis of Strength	Strength	
ı	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
TROLAMINE (UNII: 903K93S3TK)		
HYDROGEN PERO XIDE (UNII: BBX060AN9V)		
GLYCERIN (UNII: PDC6A3C0OX)		
CARBO XYPO LYMETHYLENE (UNII: 0 A5MM30 7FC)		

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78623-370-01	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021	
2	NDC:78623-370-02	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021	
3	NDC:78623-370-03	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021	
4	NDC:78623-370-04	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021	
5	NDC:78623-370-05	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021	
6	NDC:78623-370-06	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/31/2020	

HAND SANITIZER

alcohol gel

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:78623-375

Route of Administration TOPICAL

Active Ingredient/Active Moiety

8		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
CARBO XYPO LYMETHYLENE (UNII: 0 A5MM30 7FC)			
WATER (UNII: 059QF0KO0R)			
TROLAMINE (UNII: 903K93S3TK)			
GLYCERIN (UNII: PDC6A3C0OX)			
HYDRO GEN PERO XIDE (UNII: BBX060 AN9 V)			

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:78623-375-01	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021		
2	NDC:78623-375-02	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021		
3	NDC:78623-375-03	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021		
4	NDC:78623-375-04	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021		
5	NDC:78623-375-05	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021		
6	NDC:78623-375-06	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021		

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OTC monograph not final	part333A	01/04/2021	

HAND SANITIZER

alcohol gel

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Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	75 mL in 100 mL	

Inactive Ingredients			
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TROLAMINE (UNII: 903K93S3TK)			
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)			
CARBO XYPO LYMETHYLENE (UNII: 0 A5MM30 7FC)			
WATER (UNII: 059QF0KO0R)			

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6	NDC:78623-360- 06	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/04/2021			

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Labeler - ningbo fareast industry co.,ltd (554535808)

Revised: 1/2021 ningbo fareast industry co.,ltd