FA HAND SANITIZER LIQUID 70- alcohol spray FA Inc.

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.

ACTIVE INGREDIENT

Active ingredients: Ethyl Alcohol 70% w/w

INACTIVE INGREDIENT

Inactive ingredients:

Water, Glycerin

PURPOSE

PURPOSE: Antiseptic

WARNINGS

Warnings:

For external use only. Flammable. Keep away from heat or flame

When using this product ■ do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water ■ avoid contact with broken skin ■ do not inhale or ingest

Stop use and ask a doctor if irritation or rash appears and lasts

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children.

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

Uses

Uses

■ Hand sanitizer to help reduce bacteria that potentially can cause disease.

Directions

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

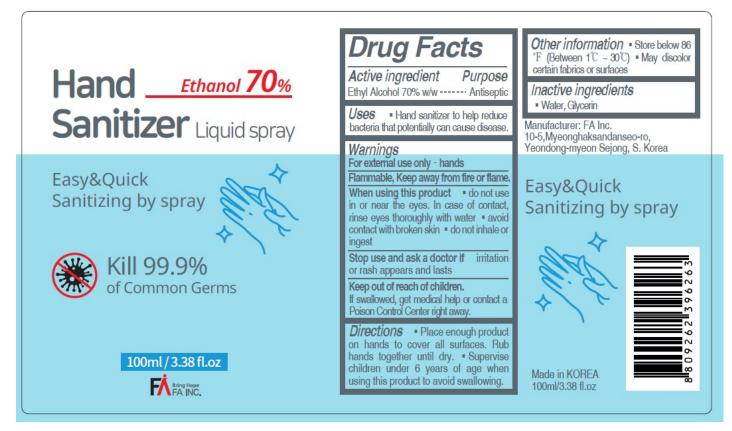
Other Information

Other Information

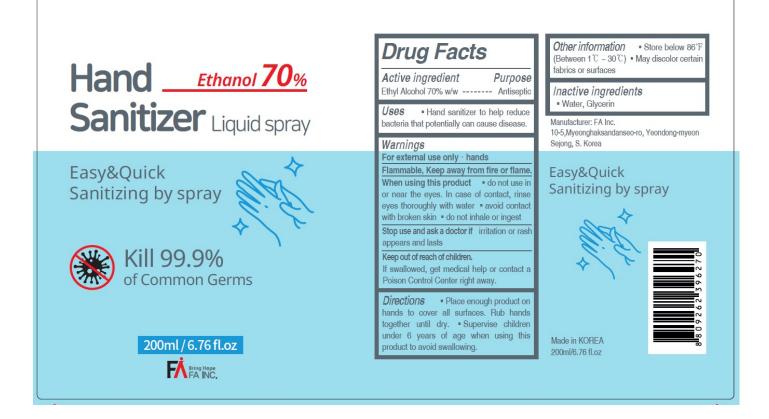
■ Store below 860(Between 1~300)

■ May discolor certain fabrics or surfaces

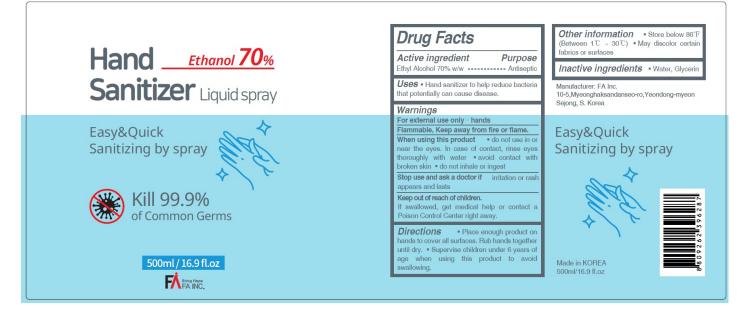
Package Label: FA Hand Sanitizer Liquid 70 100mL



Package Label: FA Hand Sanitizer Liquid 70 200mL



Package Label: FA Hand Sanitizer Liquid 70 500mL



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			ltem Code (Source)			NDC:/4932-310	
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Ingredient Name				Basis of Strength		Strength	
Alcohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)				Alcohol		70 mL in 100 mL	
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Item Code		Package Description			•	Marketing End Date	
NDC:74932-310- 01	100 mL in 1 BOTT Product	'LE, SPRAY; Type 0: Not a Combination		04/01/2020			
NDC:74932-310- 02	200 mL in 1 BOT Product	TLE, SPRAY; Type 0: Not a Combination		04/01/2020			
NDC:74932-310- 03	500 mL in 1 BOT Product	OTTLE, SPRAY; Type 0: Not a Combination		0 4/0 1/20 20			
	eoduct Type oute of Administ ctive Ingredie cohol (UNII: 3K99 active Ingred active Ingred der (UNII: 059QF ycerin (UNII: PDC bckaging Item Code NDC:74932-310- 02 NDC:74932-310-	roduct Information roduct Type pute of Administration Ctive Ingredient/Active Moid Ingredients Ctive Ingredients Ctive In	roduct Information roduct Type HUMAN OTC DRUG TOPICAL TOPICAL TOPICAL Ctive Ingredient/Active Moiety Ingredient Name cohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) active Ingredients Ingredient Name active Ingredients Ingredient Name ter (UNII: 059QF0K00R) ycerin (UNII: PDC6A3C00X) Ackaging Item Code Package Description NDC:74932-310- 01 00 mL in 1 BOTTLE, SPRAY; Type 0: Not a Comb Product NDC:74932-310- 20 0 mL in 1 BOTTLE, SPRAY; Type 0: Not a Comb Product NDC:74932-310- 20 0 mL in 1 BOTTLE, SPRAY; Type 0: Not a Comb Product NDC:74932-310- 20 0 mL in 1 BOTTLE, SPRAY; Type 0: Not a Comb Product NDC:74932-310- 20 0 mL in 1 BOTTLE, SPRAY; Type 0: Not a Comb Product NDC:74932-310- 20 0 mL in 1 BOTTLE, SPRAY; Type 0: Not a Comb	roduct Information boduct Type HUMAN OTC DRUG Item Cod pute of Administration TOPICAL TOPICAL trive Ingredient/Active Moiety Tingredient Name cohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) active Ingredients active Ingredients active Ingredients ter (UNII: 059QF0K00R) yeerin (UNII: PDC6A3C00X) Ackaging Item Code Package Description NDC:74932-310- 100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 200 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	roduct Information oduct Type HUMAN OTC DRUG Item Code (Source) pute of Administration TOPICAL Strive Ingredient/Active Moiety Strive Ingredient/Active Moiety Sohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) Alcohol Sohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) Alcohol active Ingredients active Ingredients active Ingredients stree (UNII: 059QF0K00R) geerin (UNII: PDC6A3C0OX) street (UNII: PDC6A3C0OX) street (UNII: PDC6A3C0OX) street (UNII: 00 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 20 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product NDC:74932-310- 500 mL in 1 BOTTLE	roduct Information roduct Type HUMAN OTC DRUG HUMAN OTC DRUG Item Code (Source) TOPICAL TOPICAL TOPICAL TOPICAL TIgredient/Active Moiety Time Code Ingredient Name Topic Ingred	

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333A	04/01/2020					

Labeler - FA Inc. (557816103)

Registrant - FA Inc. (557816103)

Establishment

Name	Address	ID/FEI	Business Operations	
FA Inc.		557816103	manufacture(74932-310)	

Revised: 5/2020

FA Inc.