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

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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

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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.

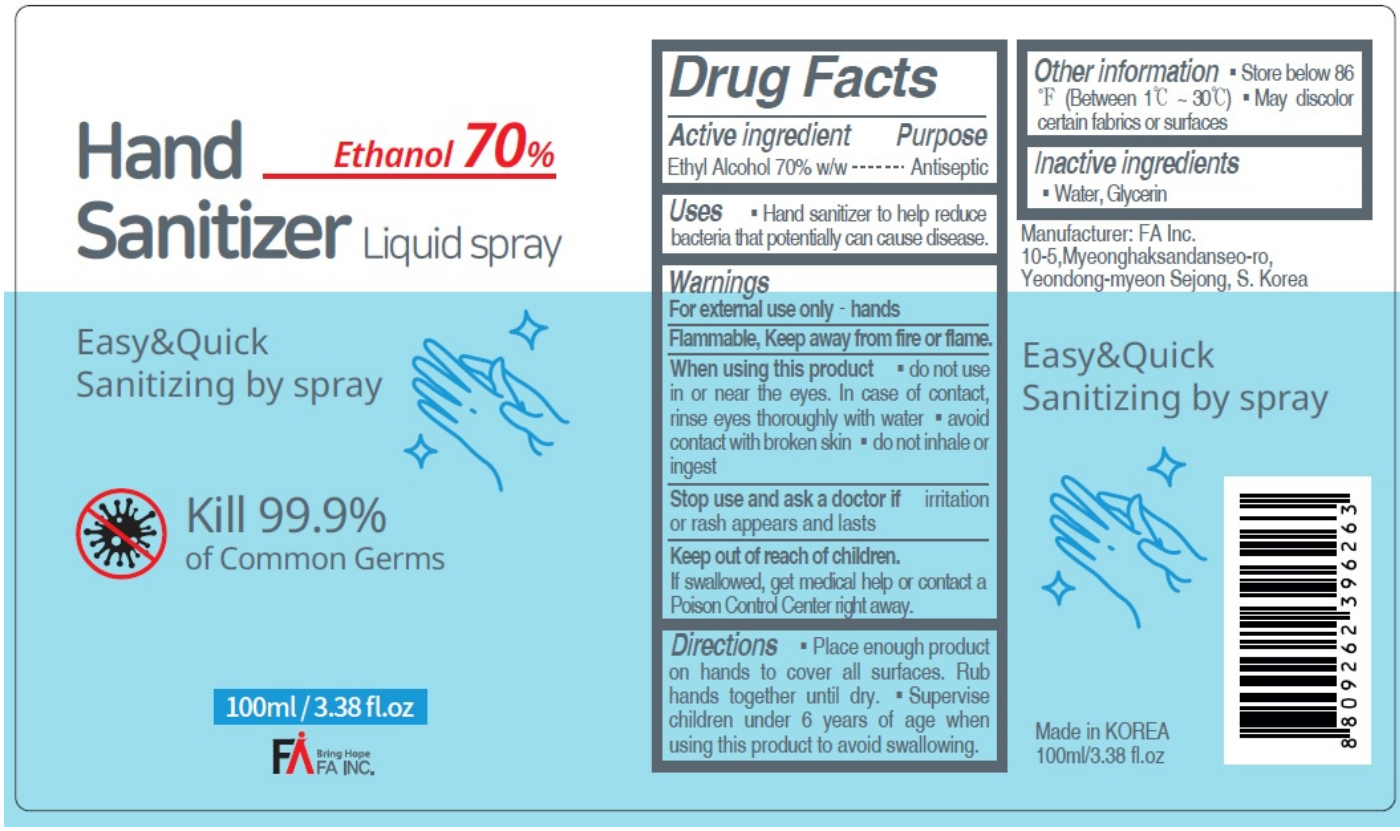
Other Information

Other Information

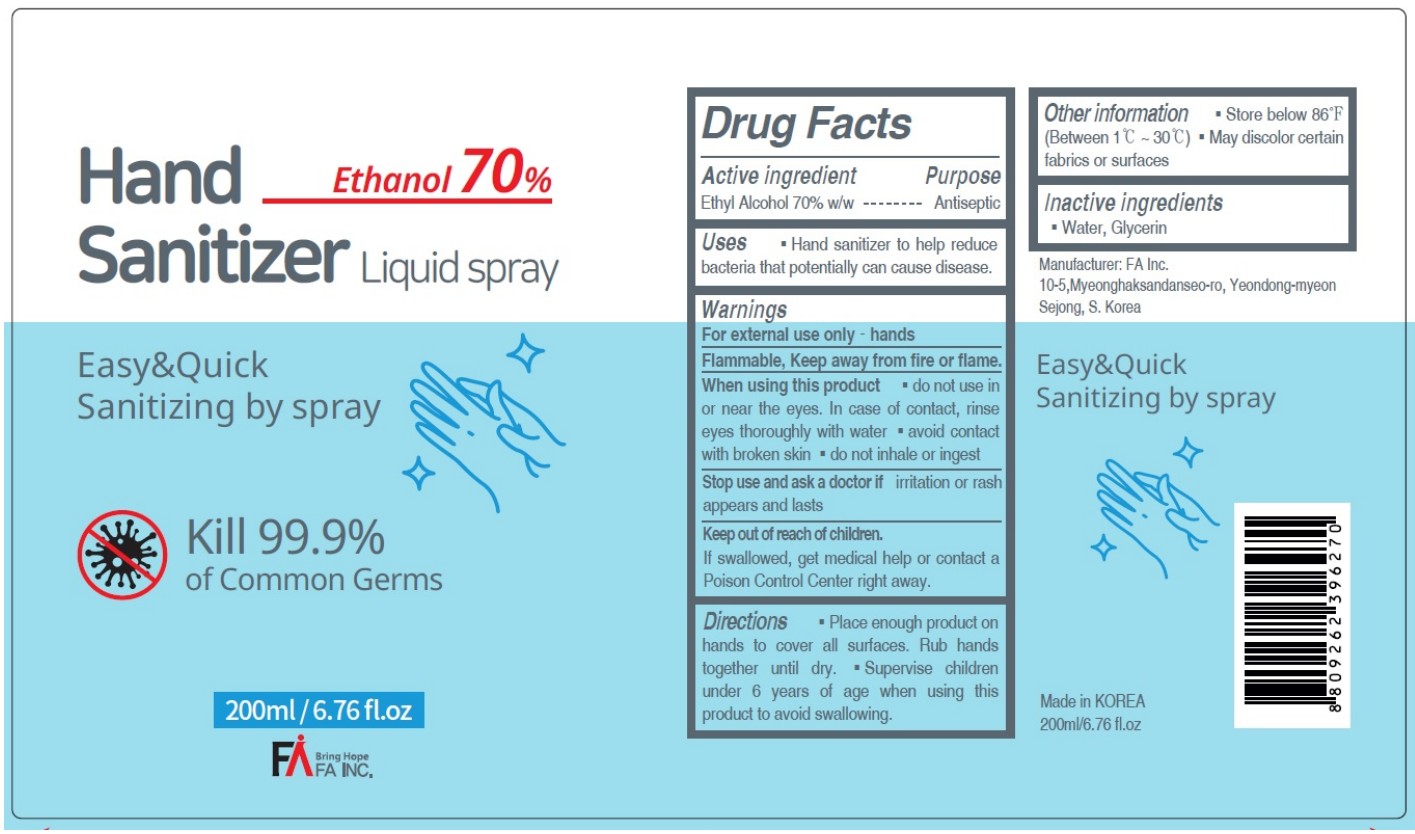
■ Store below 86□(Between 1~30□)

■ May discolor certain fabrics or surfaces

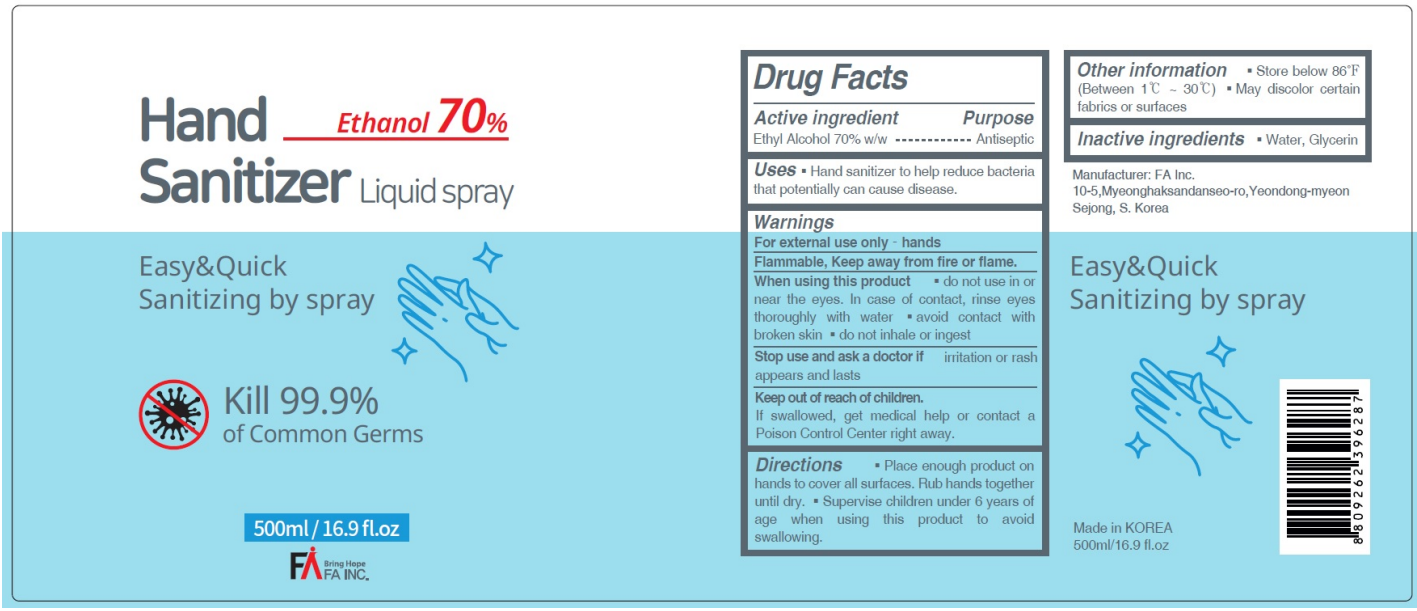
Package Label: FA Hand Sanitizer Liquid 70 100mL



Package Label: FA Hand Sanitizer Liquid 70 200mL



Package Label: FA Hand Sanitizer Liquid 70 500mL



FA HAND SANITIZER LIQUID 70

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74932-310
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:74932-310-01	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	
2	NDC:74932-310-02	200 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	
3	NDC:74932-310-03	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	

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.