

SIGNATURE SELECT ADVANCED HAND SANITIZER WITH ALOE VERA-alcohol solution
Living Better Brands LLC

21130-066 Alcohol sanitizer with Aloe

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Uses

- To decrease bacteria on the skin that could cause disease
- Recommended for repeated use

Warnings

For external use only.

Flammable, keep away from fire or flame

When using this product

Keep out of eyes. In case of contact with eyes, flush thoroughly with water

Avoid contact with broken skin

Do not inhale or ingest

Stop use and ask a doctor if skin irritation develops

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

Directions:

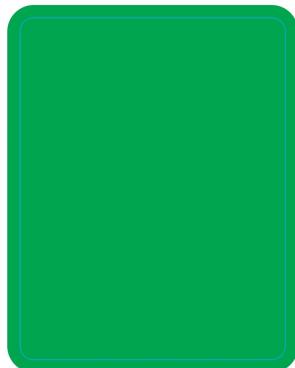
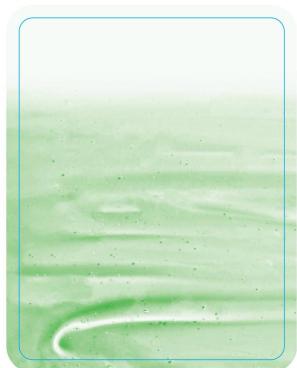
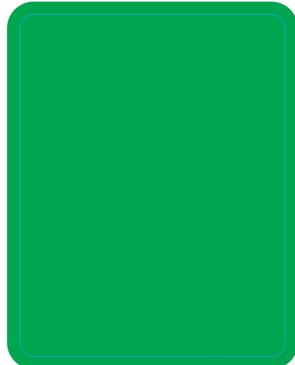
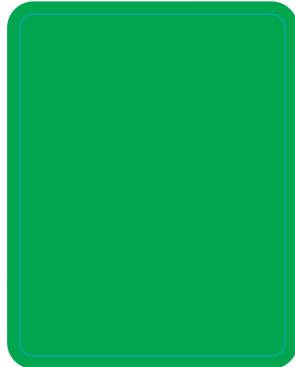
- Wet hands thoroughly with product and allow to dry without wiping
- For children under 6, use only under adult supervision
- Not recommended for infants

Other Information

- Do not store above 105F
- May discolor some fabrics
- Harmful for wood finishes and plastics

Inactive ingredients

Water, Glycerin, Carbomer, Aminomethyl Propanol, Aloe Barbadensis Leaf Extract, Propylene Glycol, Fragrance, Tocopheryl Acetate, Blue 1, Yellow 5



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alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-066
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)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
GLYCERIN (UNII: PDC6A3C00X)	
CARBOMER (UNII: 0A5MM307FC)	
AMINOMETHYL PROPANOL (UNII: LU49E6626Q)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-066-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/26/2026	
2	NDC:21130-066-02	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/26/2026	
3	NDC:21130-066-03	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/26/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	01/26/2026	

Labeler - Living Better Brands LLC (009137209)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Bright Care Co., Ltd.		699496801	manufacture(21130-066)

Revised: 1/2026

Living Better Brands LLC