

HAND SANITIZER- alcohol gel
GFA Production (Xiamen) Co., Ltd.

Hand Sanitizer

Drug Facts

Active ingredient

Ethyl alcohol 70% v/v

Purpose

Antiseptic

Uses

- For handwashing to decrease bacteria on the skin
- Recommended for repeated use.

Warnings

Flammable, keep away from fire or flame

For external use only.

Do not use

- in the eyes.

Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

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.

Other information

Store at 15° to 25°C (59° to 77°F)

Inactive ingredients

Carbomer, propylene glycol, purified water, titanium dioxide

Package Labeling:

Directions for safe use:

For hand washing to decrease bacteria on the skin that may cause disease.

Warning:

Intended for use exclusively by adults, and use only as directed. Excessive use may cause irritation to skin and urinary tract. Discontinue use if rash, redness, or itching occurs. Consult your physician if irritation persists. Keep out of reach of children.



Genuine First Aid



Hand Sanitizer

- Rinse-Free Hand Sanitizer Gel
- Safety Instant Hand Refreshing Hand Cleanser Gel
- Effective 99%

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TO REPORT A SERIOUS ADVERSE EVENT, CONTACT +1 (727) 441-3222





Manufacturer:

GFA Production Xiamen Co., Ltd
 No. 20 Huli Industrial Park Meixi Road,
 Tong'an, Xiamen Fujian, China 361100
 Made in China



6 970529 421584

Net Wt. 6.75 fl oz
(200ml)

HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50814-051
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50814-051-02	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/01/2020	

Labeler - GFA Production (Xiamen) Co., Ltd. (421256261)

Revised: 10/2023

GFA Production (Xiamen) Co., Ltd.