

ICONIC MEDICAL GROUP HAND SANITIZER - 12OZ 12OZ- hand sanitizer gel
Forsythe Cosmetic Group, Ltd.

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.

DOSAGE & ADMINISTRATION SECTION

354.882mL NDC: 76909-0001-1

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

HAND SANITIZER
12oz Bottle Screen





DIST: ICONIC MEDICAL GROUP, LLC
880 ELKTON DR, STE A
CO SPRINGS, CO 80907
WWW.ICONICMEDICALGROUP.COM



MFG: COLOR CLUB
FREEPORT, NY 11520



8 50018 22604 2



ADVANCED
HAND SANITIZER

ALCOHOL ANTISEPTIC 75%
TOPICAL SOLUTION

Fragrance Free
Antiseptic Hand Rub
Non-sterile Solution

MADE IN THE USA

12 FL OZ (354 mL)

Drug Facts

Active ingredient[s]	Purpose
Ethanol 75% v/v	Antiseptic

Use[s]
Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

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

Do not use

- 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 thoroughly 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 way.

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.

Other Information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients glycerin, hydrogen peroxide, xanthan gum, distilled water.

— DIELINE EDGE
- - BLEED AREA

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INACTIVE INGREDIENT SECTION

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INDICATIONS & USAGE SECTION

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OTC - KEEP OUT OF REACH OF CHILDREN SECTION

354.882mL NDC: 76909-0001-1

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

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OTC - ACTIVE INGREDIENT SECTION

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76909-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	266.1615 mL in 354.882 mL

Inactive Ingredients

Ingredient Name	Strength
XANTHAN GUM (UNII: TTV12P4NEE)	17.009712 mL in 354.882 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	10.64646 mL in 354.882 mL
GLYCERIN (UNII: PDC6A3C0OX)	10.64646 mL in 354.882 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76909-0001-1	354.882 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Forsythe Cosmetic Group, Ltd. (038650495)

Registrant - Forsythe Cosmetic Group, Ltd. (038650495)

Establishment

Name	Address	ID/FEI	Business Operations
Forsythe Cosmetic Group, Ltd.		038650495	manufacture(76909-0001)

Revised: 4/2020

Forsythe Cosmetic Group, Ltd.