INSTANT HAND SANITIZER- alcohol gel Zidac Laboratories 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.

Instant Hand Sanitizer

DRUG FACTS

Active Ingredient

Ethyl Alcohol 77%v/v

Purpose

Antimicrobial

Uses

- Hand sanitizer helps reduce bacteria on the skin
- Recommended for repeated use

Warnings

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

When using this product

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor

if irritation or rash occurs and lasts.

Keep out of reach of children.

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

Directions

- Place product in palm of hand
- Rub hands together until dry
- Supervise children under 6 years of age when using product

Storage

Store below 110°F (43°C)

Inactive ingredients

Water (Aqua), Glycerin, Carbomer, Aminomethyl Propanol, Fragrance (Parfum), Aloe Barbadensis Leaf Juice, Propylene Glycol

Package Labeling: 100ml



INSTANT HAND SANITIZER

KILLS 99.9% OF GERMS

- Fast drying
- No water
- No sticky residue
- With Aloe Vera

3.38fl.oz. @ 100ml

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Produced in the UK / Zidac Laboratories Ltd, Unit 5 Merlin Park, Airport Service Road, Portsmouth, PO3 5FU United Kingdom





Package Labeling: 500ml





DRUG FACTS

Active Ingredient Ethyl Alcohol 77%v/v

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KILLS 99.9% OF GERMS

- Fast drying
- No water
- No sticky residue
- With Aloe Vera

16.9fl.oz. ⊕ 500ml





Package Labeling:5000ml







INSTANT HAND SANITIZER

77% Alcohol Gel

KILLS 99.999% OF GERMS

- Fast drying
- No water
- No sticky residue
- With Aloe Vera







roduced in the UK
idac Laboratories Ltd, Unit 5
Aerlin Park, Airport Service Road
ortsmouth, PO3 5FU, United Kir

DRUG FACTS

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Ethyl Alcohol 77%v/v

Purpose

Antimicrobial

Uses

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INSTANT HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:79699-000

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)
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ALCOHOL

0.77 mL in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)				
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:79699-000-	$3\ \ 100\ mL$ in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2020		
2 NDC:79699-000-	6 500 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2020		
3 NDC:79699-000-	9 5000 mL in 1 CAN; Type 0: Not a Combination Product	09/01/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	08/01/2020		

Labeler - Zidac Laboratories Ltd (222761253)

Revised: 10/2020 Zidac Laboratories Ltd