MED MATE INSTANT HAND SANITIZER- alcohol gel United Specialty Solutions

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.

MED Mate Instant Hand Sanitizer

Active Ingredient(s)

Ethyl Alcohol 70%. Purpose: Antimicrobial

Purpose

Antimicrobial, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria on the skin that could cause disease Recommended for repeated use,

Warnings

Flammable. Keep away from heat or flame.

For external use only.

When using this product do not use in or near the eyes. In case of contact, rinse eyes throughly with water.

Stop and ask a doctor if irritation or rash appears and last.

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

Directions

- Place a thumbnail size amount in your palm.
- Rub hands together briskly until dry.
- Children under 6 years of age must be supervised when using Med-Made

Inactive ingredients

Water, Glycerin, Propylene Glycol, Aloe Vera, Carbomer, Vitamin E

Questions or comments?

Call 1-888-760-7018

Package Label - Principal Display Panel

MED MAte

Instant Hand Sanitizer

Fight against 99% of germs, bacteria, and viruses

Made with 70% Ethanol

Made in the U.S.A.

16.9 FL OZ (473mL)

MED MATE INSTANT HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:80263-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

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70 mL in 100 mL

Inactive Ingredients

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Ingredient Name	Strength
.ALPHATOCOPHEROL (UNII: H4N855PNZ1)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KOOR)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

Packaging

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#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:80263-001-01	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/25/2020		
2	NDC:80263-001-02	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/25/2020		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/25/2020	

Labeler - United Specialty Solutions (115068941)

Establishment							
Name	Address	ID/FEI	Business Operations				
BILLE CROSS LABORATORIES INC		008298879	manufacture(80263-001)				

Revised: 8/2020 United Specialty Solutions