

UNIVERSITY MEDICAL PHARMACEUTICALS HAND SANITIZER- alcohol liquid
University Medical Pharmaceuticals Corp.

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.

University Medical PHARMACEUTICALS® HAND SANITIZER

Drug Facts

Active ingredient

Alcohol 70% v/v

Purpose

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

■ on 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 away.

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 to avoid swallowing.

Other information

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

Inactive Ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Glycerin, Tocopheryl Acetate, Purified Water.

NEW

REDUCE BACTERIA THAT POTENTIALLY CAUSE DISEASE

KEEPS HANDS FEELING FRESH AND SOFT

Questions? Call toll free 855-299-8800

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Packaging

NEW

University Medical
PHARMACEUTICALS

HAND SANITIZER

REDUCE BACTERIA THAT POTENTIALLY CAUSE DISEASE

KEEPS HANDS FEELING FRESH AND SOFT

8 FL OZ (236 mL)

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UNIVERSITY MEDICAL PHARMACEUTICALS HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50544-888
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
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	

AMINOMETHYLPROPANOL (UNII: LU49E6626Q)

GLYCERIN (UNII: PDC6A3C0OX)

.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)

WATER (UNII: 059QF0K00R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50544-888-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

Labeler - University Medical Pharmaceuticals Corp. (809706252)

Revised: 5/2020

University Medical Pharmaceuticals Corp.