MEDI FECT ANTISEPTIC HAND WASH- ethyl alcohol liquid
Medical Chemical Corporation

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.

Medi Fect Label

Indications for use: For hospital and professional use only. Medi-Fect is intended to be used as a hand-wash to reduce bacteria that can potentially cause disease. Recommended for repeated use.

Ingredients: 70% v/v ethyl alcohol, propylene glycol, emolients (polysorbate 80, cetyl alcohol, acetylated lanolin alcohol), carbomer, diazolidinyl urea, methyl paraben, aloe vera and propyl paraben. Contains emollients and skin conditioners. Contains no added fragrance or dyes.

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Warnings: Flammable, keep away from fire or flame. For external use only. Do no use in the eyes. Discontinue use if irritation or redness develops. Keep out of reach of children. In case of ingestion contact poison control center immediately.

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Directions: Place a 'palmful' (about 5 g) of product in one hand. Spread on both hands and rub into the skin until dry (approximately 1 to 2 minutes). Place a smaller amount (2.5 grams) into one hand, spread over both hands to wrist, and rub into skin until dry (approximately 30 seconds).

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medifectlabel.jpg

MED CHEM®

Medi-Fect™ Hand Sanitizer

With Aloe vera

Drug Facts

Active Ingredients — **Purpose**

Ethyl alcohol, 70% v/v — Germicide

Uses

Intended for use as a hand sanitizer to reduce pathogenic bacteria. Recommended for repeated use. For external use on the skin only. Not for use on the eyes.

Warnings

Danger: Highly flammable liquid and vapor. Keep away from heat, sparks, open flames and hot surfaces. No smoking. Keep container tightly closed. Use only non-sparking tools. Take precautions against static discharge. In case of fire, use fire extinguishers approved for alcohol fires. In case of ingestion contact a poison control center. Discontinue use if irritation or redness develops. Keep out of reach of children.

Directions

Spray about 5 g (1 tsp.) on to one hand and spread over both hands to the wrist. Rub into the skin until dry. Repeat.

Other Information.

Keep tightly closed and protected from light. Store at room temperature,

Inactive ingredients ● Purified water ● Emollients (Laurel, Myristal and Cetyl Lactate) ● Aloe Vera ● Carbomer ● Parabens ●Diisopropylamine

Contains no added fragrances or dyes.

NDC 12745-177

Manufactured in the USA by Medical Chemical Corp., 19430 Van Ness Ave. Torrance Ca. 90501

MEDI FECT ANTISEPTIC HAND WASH

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:12745-177
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	59.86 g in 100 mL
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4) (DIAZOLIDINYL UREA - UNII:H5RIZ3MPW4)	DIAZOLIDINYL UREA	1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)	0.5 g in 100 mL
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	0.5 g in 100 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.5 g in 100 mL
DIISOPROPYLAMINE (UNII: BR9JLI40NO)	0.505 g in 100 mL
WATER (UNII: 059QF0KO0R)	

Packaging

		Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12745-177-01	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/14/2001	
2	NDC:12745-177-02	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/14/2001	
3	NDC:12745-177-03	3785 mL in 1 BOTTLE, PLASTIC; Type 1: Convenience Kit of Co-Package	05/14/2001	
4	NDC:12745-177-04	18927 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/14/2001	
5	NDC:12745-177-05	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/14/2001	
6	NDC:12745-177-06	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/14/2001	
7	NDC:12745-177-07	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/14/2001	
8	NDC:12745-177-08	208198 mL in 1 DRUM; Type 0: Not a Combination Product	05/14/2001	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/14/2001	

Labeler - Medical Chemical Corporation (008496861)

Establishment

Name	Address	ID/FEI	Business Operations
Medical Chemical Corporation		008496861	manufacture(12745-177)

Revised: 10/2020 Medical Chemical Corporation