## 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.

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#### Medi Fect Label

Indications for use: For hospital and professional use only. Medi-Fect is intended to be used as a handwash 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



# Medi-Fect™ Antiseptic Hand-Wash (with Aloe Vera)



## Manufactured by Medical Chemical Corp. Torrance, CA 90501

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NDC# 012745-177E

#### 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	<b>Basis of Strength</b>	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: ZY8 1Z8 3H0 X)	0.5 g in 100 mL			
DIISO PRO PYLAMINE (UNII: BR9 JL I40 NO)	0.505 g in 100 mL			
WATER (UNII: 059QF0KO0R)				

Packaging		
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#	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	

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: 9/2019 Medical Chemical Corporation