

ALOESAFE ANTISEPTIC HAND SANITIZER- 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.

AloeSafe

Indications for Use

Indications for use: AloeSafe is intended to be used as a waterless hand sanitizer to reduce microorganisms that can cause disease. Recommended fo repeated use.

Ingredients

Ingredients: Ethyl alcohol 62% (active ingredient), purified water, carbomer, aloe vera gel, diisopropylamine, vitamin E, and lavender oil.

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Warnings

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Directions

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AloeSafe Label



AloeSafe™

Antiseptic Hand Sanitizer
with Aloe Vera, Lavender Oil,
and Vitamin E



16 Fl. Oz. (473ml)

Your hands deserve only the best! Hardy's AloeSafe™ contains no dyes, parabens, or propylene glycol. Moisturizers and conditioners leave hands smooth, soft, and silky. Kills 99.99% of bacteria on contact.

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Manufactured for:
Hardy Diagnostics
1430 W. McCoy Lane
Santa Maria, CA 93455
(800) 266-2222
www.HardyDiagnostics.com



Cat. no.
AS16HD

ALOESAFE ANTISEPTIC HAND SANITIZER

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:12745-180
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	60 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
POLYACRYLIC ACID (450000 MW) (UNII: KD3S7H73D3)	0.25 g in 100 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.2 g in 100 mL
LAVENDER OIL (UNII: ZBP1YXW0H8)	0.02 g in 100 mL
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	0.04 g in 100 mL
WATER (UNII: 059QF0KO0R)	28.03 g in 100 mL
DIISOPROPYLAMINE (UNII: BR9JLI40NO)	0.2 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12745-180-01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/14/2019	
2	NDC:12745-180-02	120 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2010	
3	NDC:12745-180-03	480 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/01/2010	

Labeler - Medical Chemical Corporation (008496861)

Registrant - Medical Chemical Corporation (008496861)

Establishment

Name	Address	ID/FEI	Business Operations
Medical Chemical Corporation		008496861	manufacture(12745-180)

Revised: 1/2022

Medical Chemical Corporation