

HAND SANITIZER- alcohol spray

MedZone Products, LLC

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.

Hand Sanitizer

DRUG FACTS

Active Ingredient

Ethyl Alcohol 62.5% v/v

Purposes

Antiseptic handwash

Uses

for handwashing to decrease bacteria on the skin

Warnings

For external use only | Flammable, keep away from fire or flame, heat, sparks, and sources of static discharge

Do not use

In eyes | In children less than 2 months of age | On open skin wounds

When using this product

If in eyes, rinse promptly and thoroughly with water | Discontinue use if irritation and redness develops

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

Apply product onto hands, spread thoroughly and rub dry | Supervise children under 6 years of age when using this product to avoid swallowing.

Other

For additional information, see Safety Data Sheets (SDS) | For emergency medical information in USA and Canada, call 1-888-255-3924 | For emergency medical information worldwide, call +1-813-248-0573 | Store between 15-30°C (59-86°F) | Avoid freezing and excessive heat above 40°C (104°F)

Inactive Ingredients

Purified Water, Glycerin, Coconut Oil, Jojoba Oil, Almond Oil

Questions?

Toll Free 866-MEDZONE (866-633-9663)

DISTRIBUTED BY

MedZone Products LLC

9300 Marshall Dr, Ste. 200,
Lenexa, Kansas 66215

PRINCIPAL DISPLAY PANEL - 52 mL Bottle Label

MedZone®
SINCE 2001

Hand
Sanitizer

SPRAY

MADE IN USA

KILLS
99.99%
OF COMMON ILLNESS CAUSING GERMS
IN 15 SECONDS

1.75 FL OZ (52mL)



ANTISEPTIC HANDWASH | DAILY USE, TRAVEL, & SPORTS | RINSE FREE

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NDC# 70338-605-75

SDS-IN-20015

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Lenexa, Kansas 66215 USA
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HAND SANITIZER

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70338-605
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	0.625 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0K00R)	
Glycerin (UNII: PDC6A3C0OX)	
Coconut Oil (UNII: Q9L0O73W7L)	
Jojoba Oil (UNII: 724GKU717M)	
Almond Oil (UNII: 66YXD4DKO9)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70338-605-75	52 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/01/2020	

Labeler - MedZone Products, LLC (080083739)**Establishment**

Name	Address	ID/FEI	Business Operations
Hubot Healthcare LLC		081084880	MANUFACTURE(70338-605)

Revised: 3/2020

MedZone Products, LLC