# SPECTRUM 62% HAND SANITIZER- ethyl alcohol gel Medline Industries, LP

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.

-----

#### 214 Spectrum 62% Hand Sanitizer

#### Active ingredient

Ethyl alcohol 62% v/v

#### **Purpose**

Antiseptic

#### Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

#### Warnings

For external use only: hands

Flammable. Keep away from heat and flame.

#### When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water
- avoid contact with broken skin
- do not inhale or ingest

#### Stop use and ask a doctor if

- irritation or redness develops
- condition persists for more than 72 hours

#### Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use onlyunder adult supervision
- not recommended for infants

#### Other information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

#### Inactive ingredients

aloe barbadensis leaf juice, carbomer, diisopropylamine, glycerin, isopropyl myristate, propylene glycol, tocopheryl acetate, water

#### **Manufacturing Information**

Manufactured for:

Medline Industries, LP

Three Lakes Drive, Northfield, IL 60093 USA

Made in USA with US and foreign components

www.medline.com

1-800-MEDLINE (633-5463)

REF: HH62G1000

V2 RH22VJO

#### Package Label



#### ©2022 Medine Industries, LP. Made in USA with US and Foreign components for Medline Industries, LP. Three Lakes Drive, Northfield, IL 60093 USA, **1-800-MEDLNE** VZ RH22VJO L0015182BB **Drug Facts** Active ingredient Purpose Ethyl alcohol 62%. .Antiseptic Uses ■ to decrease bacteria on the skin that could cause disease ■ recommended for repeated Warnings For external use only: hands Flammable. Keep away from heat and flame. When using this product ■ keep out of eyes. In case of contact with eves, flush thoroughly with water. ■ avoid contact with broken skin do not inhale or ingest Stop use and ask a doctor if irritation or redness develops ■ condition persists for more than 72 hours Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Directions ■ wet hands thoroughly with product and allow to dry without wiping ■ for children under 6, use only under adult supervision In not recommended for infants 4 Other information ■ do not store above 105°F ■ may discolor some fabrics harmful to wood finishes and plastics Inactive ingredients aloe barbadensis leaf juice, carbomer, diisopropylamine, glycerin, 88 isopropyl myristate, propylene glycol, tocopheryl acetate, water

SDS-M0-15036, SDA WI-2059, SDA WI-2486,

DSP-MO-28, DSP-MO-34

00

#### **SPECTRUM 62% HAND SANITIZER**

ethyl alcohol gel

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:53329-214

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name

Basis of Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

62 mL in 100 mL

# Inactive Ingredients Ingredient Name Strength GLYCERIN (UNII: PDC6A3C00X) .ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)

CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)

DIISOPROPYLAMINE (UNII: BR9JLI40NO)
ISOPROPYL MYRISTATE (UNII: ORE8K4LNJS)

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)

WATER (UNII: 059QF0KO0R)

ALOE VERA LEAF (UNII: ZY81Z83H0X)

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
<b>1</b> NDC:53329-214-13	59.147 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2018	12/06/2022	
NDC:53329- 214-70	443.6 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2018	01/31/2022	
<b>3</b> NDC:53329-214-84	1000 mL in 1 BAG; Type 0: Not a Combination Product	04/09/2018		
4 NDC:53329- 214-85	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2018		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	04/09/2018			
inidi					

## Labeler - Medline Industries, LP (025460908)

## Registrant - Medline Industries, LP (025460908)

Revised: 8/2023 Medline Industries, LP