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

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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

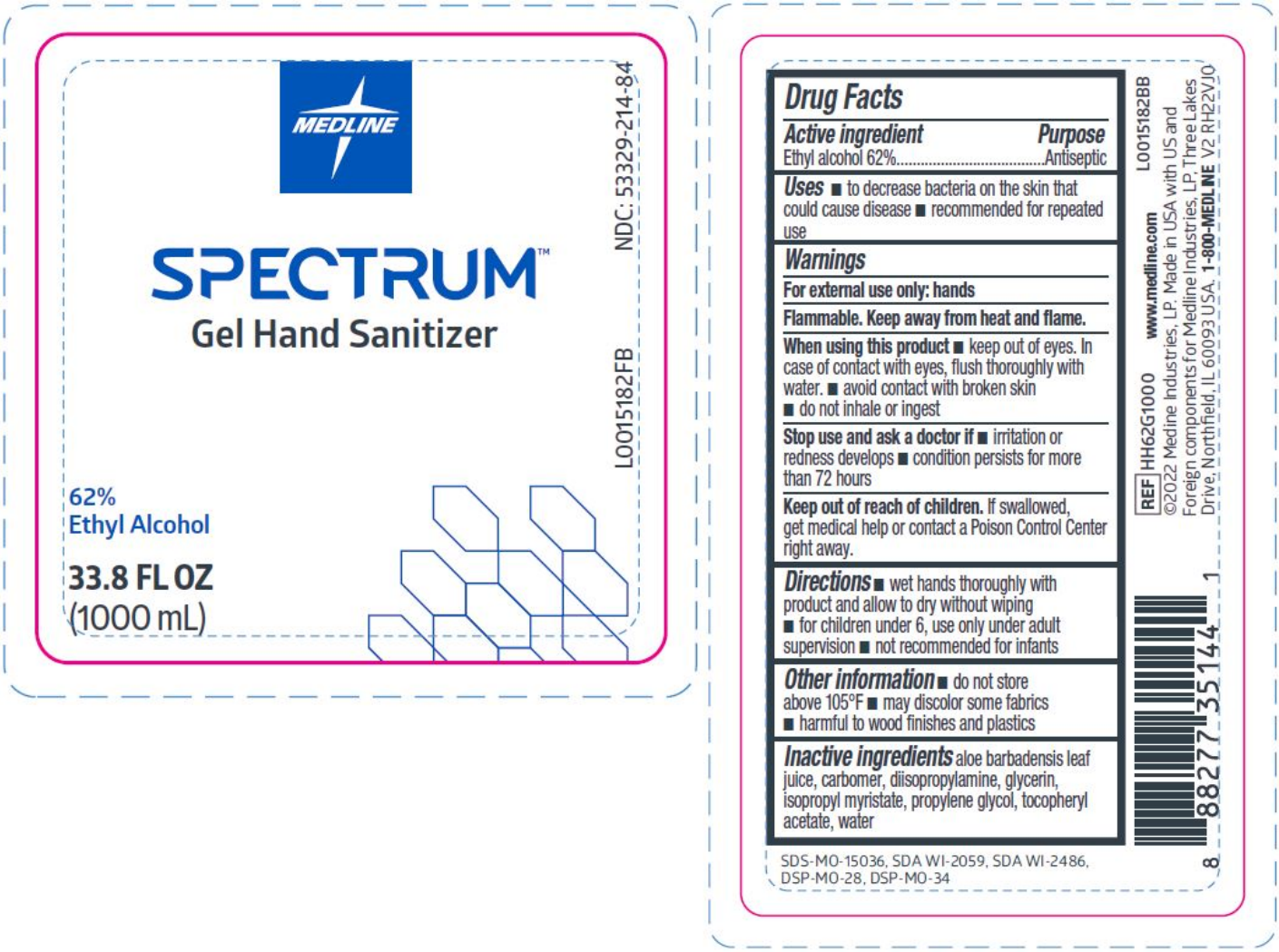
www.medline.com

1-800-MEDLINE (633-5463)

REF: HH62G1000

V2 RH22VJO

Package Label



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**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
DIISOPROPYLAMINE (UNII: BR9JLI40NO)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53329-214-13	59.147 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2018	12/06/2022
2	NDC:53329-214-70	443.6 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2018	01/31/2022
3	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 Drug	505G(a)(3)	04/09/2018	

Labeler - Medline Industries, LP (025460908)

Registrant - Medline Industries, LP (025460908)

Revised: 5/2024 Medline Industries, LP