

SPECTRUM HAND SANITIZER WIPES- alcohol wipes cloth
Medline Industries, Inc.

SPECTRUM HAND SANITIZER WIPES

Active ingredient

Ethyl alcohol 70%

Purpose

Antiseptic

Uses

to decrease bacteria on the skin that could cause disease

Warnings

For external use only: hands

Flammable. Keep away from fire 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

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, glycerin, isopropanol, propanediol, tocopheryl acetate, water

Uses

to decrease bacteria on the skin that could cause disease

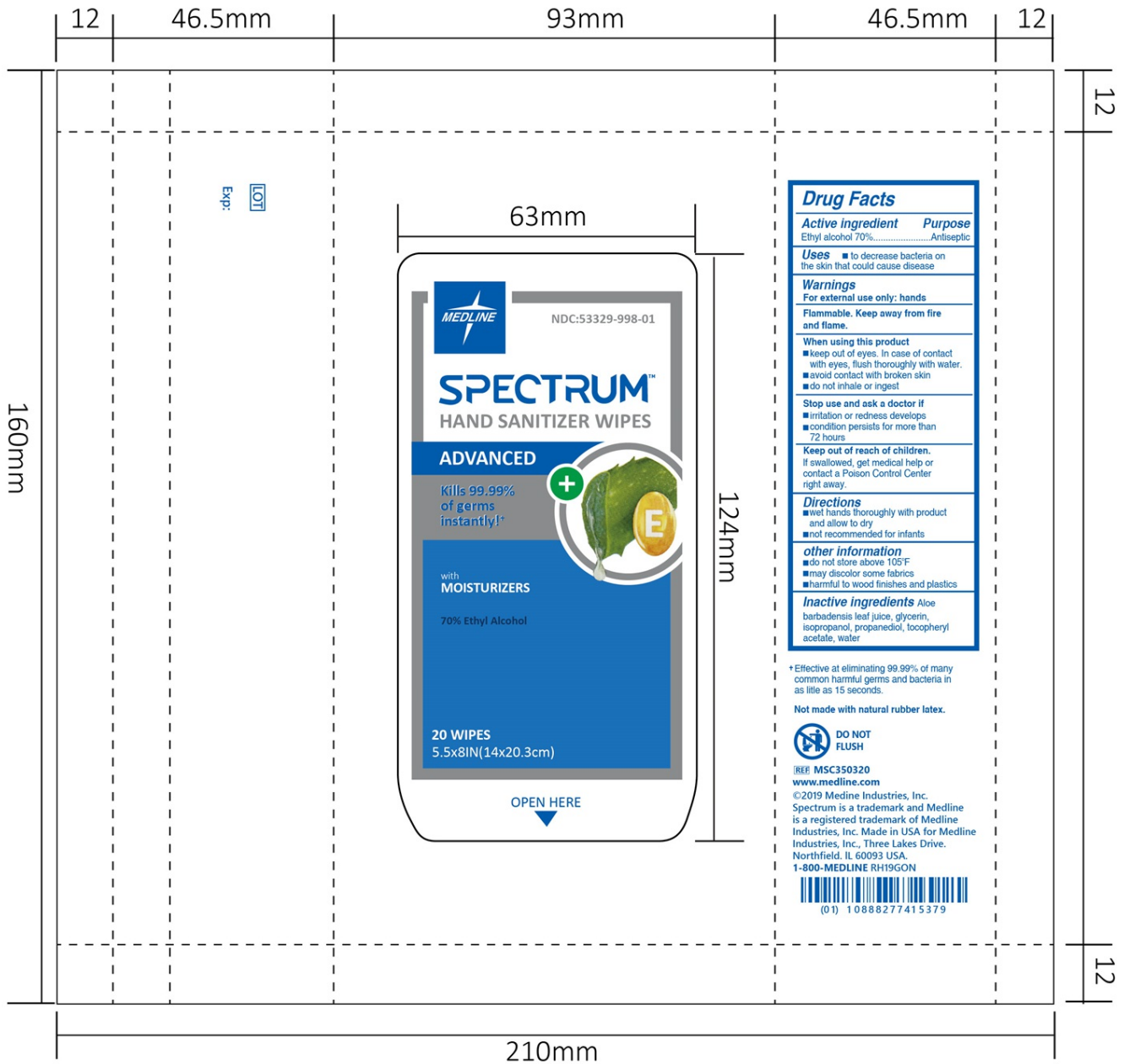
recommended for repeated use

Directions

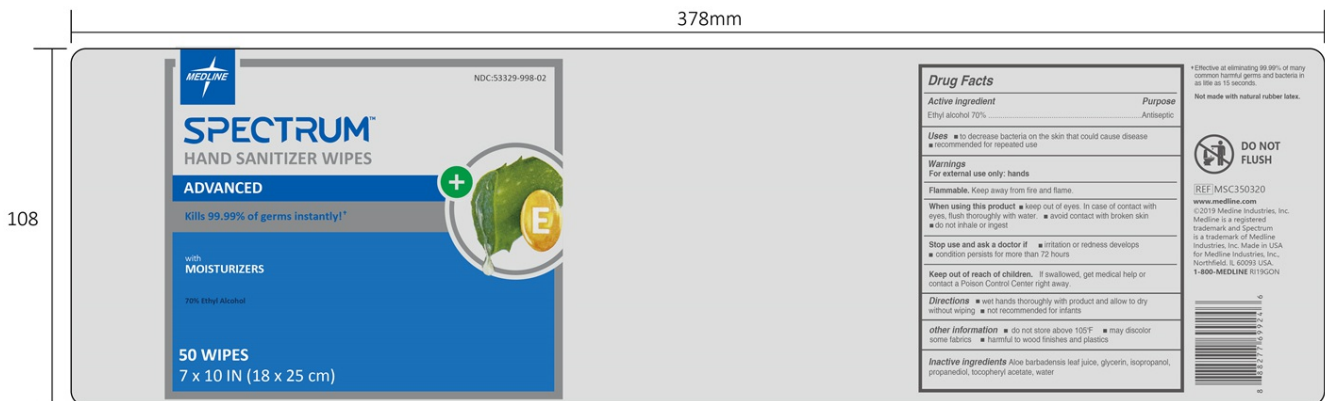
wet hands thoroughly with product and allow to dry without wiping

not recommended for infants

20 WIPES, NDC: 53329-998-01



50 WIPES, NDC: 53329-998-02



160 WIPES, NDC: 53329-998-03



SPECTRUM HAND SANITIZER WIPES

alcohol wipes cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53329-998
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PROPANEDIOL (UNII: 5965N8W85T)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53329-998-01	20 in 1 PACKAGE	04/30/2021	
1		1.32 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
2	NDC:53329-998-02	50 in 1 CANISTER	04/30/2021	

2		2.55 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
3	NDC:53329-998-03	160 in 1 CANISTER	04/30/2021	
3		1.2 mL in 1 APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/30/2021	

Labeler - Medline Industries, Inc. (025460908)

Revised: 10/2023

Medline Industries, Inc.