HAND SANITIZER- ethyl alcohol gel
Steris 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.

Soft n Sure Hand Sanitizer

Active ingredient

Ethyl Alcohol 65%

Purpose

Antiseptic

Uses

Healthcare Personnel Handwash to decrease bacteria on the skin before contact with patients under medical care or treatment

Warnings

For external use only-hands

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

skin irritation develops. If condition persists more than 72 hours consult a doctor.

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

Other information

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

Inactive ingredients

benzophenone-4, carbomer, fragrance, glycerin, isopropyl myristate, propylene glycol, tocopheryl acetate, water

Questions or Comments

800-548-4873

adverse reaction section

Effective at eliminating 99.99% of many common harmful germs and bacteria in as little as 15 seconds*

Soft' N Sure is a registered trademark of STERIS Corporation. Dignity Health and its logo are tradmarks of Dignity Health

Manufactured for:

STERIS Corporation

7501 Page Avenue

St. Louis, MO 63133 USA

800-548-4873 www.steris.com

951.001/951AB

principal display panel

NDC 0519-8802-13

Dignity Health Hand Sanitizer Soft N SureAntiseptic Hand Gel

Moisturizing Fast Acting

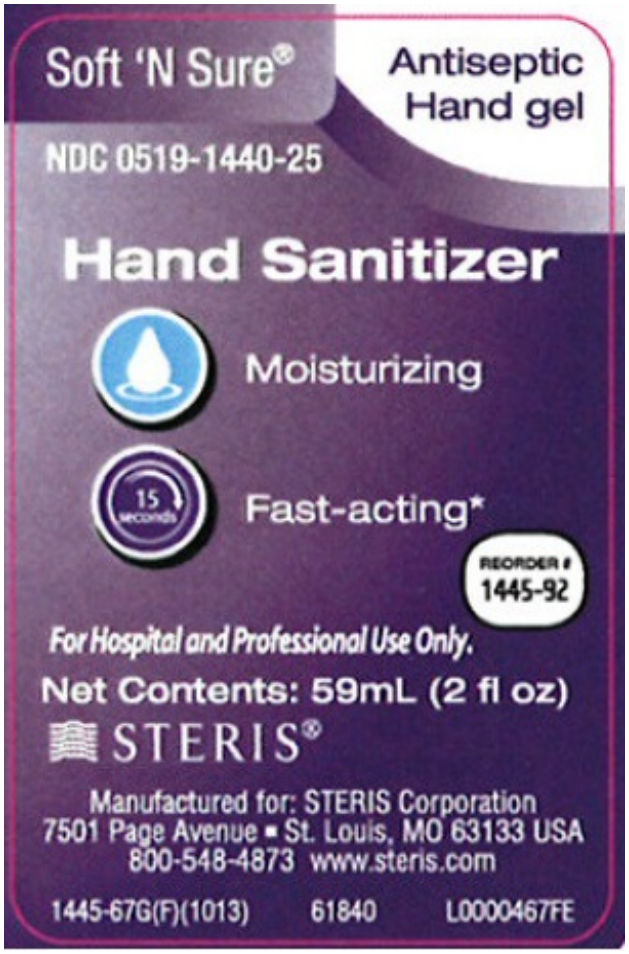
8D02-26

For Hospital and Professional use Only

6542

8D02-XEA (F) (814)

NET CONTENTS: 444 mL (15 FL OZ)



HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0519-8802
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	578 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SULISOBENZONE (UNII: 1W6L629B4K)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
glycerin (UNII: PDC6A3C0OX)	
isopropyl myristate (UNII: 0RE8K4LNJS)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0519-8802-13	444 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/08/2014	
2	NDC:0519-8802-41	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/08/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/08/2014	

Labeler - Steris Corporation (139424188)

Registrant - Vi-Jon (088520668)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(0519-8802)

Revised: 11/2019 Steris Corporation