CV MEDICATED- triclosan soap
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.

CV® Medicated Lotion Soap

Drug Facts

Active ingredient

0.5% Triclosan

Purpose

Antiseptic

Use

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

Warnings

For external use only

Discontinue use if irritation and redness develop. If irritation persists for more than 3 days, consult a physician.

When using this product do not get it in the eyes; this product causes eye irritation upon direct contact. In case of eye exposure, rinse thoroughly with water. If eye irritation persists, contact a physician.

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

Wet skin and spread a small amount on the hands. Wash hands, rinse thoroughly with water, and repeat as necessary between patient contacts.

Inactive ingredients

Water, alkali salts of fatty acids, sorbitol, disodium cocoamphodiacetate, cocoamide DEA, glycerin, acrylates/PEG-10 maleate/styrene copolymer, hydrolyzed collagen, tetrasodium EDTA, sodium chloride, iodopropynyl butylcarbamate, fragrance

Questions or comments?

800-548-4873 www.steris.com

PRINCIPAL DISPLAY PANEL - 1 Liter Bottle Label

CV® Medicated

Lotion
Soap

NDC 0519-6262-41

Hand Soap

Moisturizing

Antimicrobial

15
seconds
Fast-acting

REORDER #
6262-87

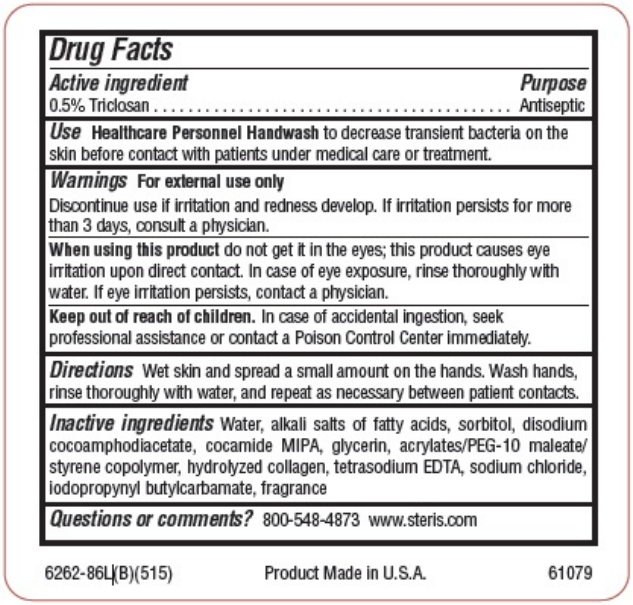
1 Liter SDS (33.8 fl oz) (1.05 qt)

STERIS®

STERIS Corporation ■7501 Page Avenue ■ St. Louis, MO 63133 ■ USA

6262-86L(F)(515)

61080



CV MEDICATED

triclosan soap

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0519-6262
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
COCO MONOISOPROPANOLAMIDE (UNII: 21X4Y0VTB1)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHACRYLATE/METHOXY PEG-10 MALEATE/STYRENE COPOLYMER (UNII: 39DK5WQ2PR)	
GELATIN HYDROLYSATE (PORCINE SKIN, MW 3000) (UNII: 0K9R94573C)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Product Characteristics

Color	WHITE (Off-white)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0519-6262-94	72 in 1 CASE		
1	NDC:0519-6262-03	118 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0519-6262-92	18 in 1 CASE		
2	NDC:0519-6262-13	444 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:0519-6262-87	12 in 1 CASE		
3	NDC:0519-6262-41	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/27/2003	

Labeler - STERIS Corporation (139424188)

Establishment

Name	Address	ID/FEI	Business Operations
STERIS Corporation		139424188	MANUFACTURE(0519-6262)

Revised: 5/2015

STERIS Corporation