KINDEST KARE ADVANCED ANTIMICROBIAL HANDWASH- benzalkonium chloride liquid
SC Johnson Professional USA, Inc.

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.

Kindest Kare® Advanced Antimicrobial Handwash

Drug Facts

Active ingredient

Benzalkonium Chloride 0.7%

Purpose

Antiseptic

Uses

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

Warnings

For external use only.

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

Stop use and ask a doctor if irritation and redness develop and persist for more than 3 days.

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

Directions

Wet hands and apply a small amount of product. Wash hands, rinse thoroughly with water, and repeat as necessary between patient contacts.

Inactive ingredients

Aqua (Water), Hexylene Glycol, Cocamidopropylamine Oxide, PEG- 80 Sorbitan Laurate, Hydroxypropyl Methylcellulose, PEG-150 Distearate, PEG-8 Dimethicone, Sorbitol,

Glycerin, Cocamidopropyl PG-Dimonium Chloride Phosphate, Cocamide MIPA, Soyamidopropylamine Oxide, Lauramine Oxide, Phenoxyethanol, Potassium Hydroxide, Citric Acid

Questions or comments?

1-866-783-0422

PRINCIPAL DISPLAY PANEL - 1 Liter Bottle Label

SCJ PROFESSIONAL
HEALTHCARE

Kindest Kare®

NDC 11084-806-41

Hand Soap
Antimicrobial Handwash
Advanced

Health Personnel Handwash

Moisturising

Fragrance Free

Manufactured for:
SC Johnson Professional USA, Inc.
Charlotte, NC 28217
1-866-783-0422 www.scjp.com
Pat. www.scjp.com/patents
Made in Canada

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

REORDER #
6267-87

deb
SKIN CARE

SAP # 4000000170


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KINDEST KARE ADVANCED ANTIMICROBIAL HANDWASH

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-806
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.7 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
COCAMIDOPROPYLAMINE OXIDE (UNII: M4SL82J7HK)	
PEG-80 SORBITAN LAURATE (UNII: 239B50Y732)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
PEG-8 DIMETHICONE (UNII: GIA7T764OD)	
SORBITOL (UNII: 506T60A25R)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
COCO MONOISOPROPANOLAMIDE (UNII: 21X4Y0VTB1)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-806-41	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2018	
2	NDC:11084-806-13	444 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2018	06/01/2022

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT			

OTC MONOGRAPH NOT FINAL	part333E	02/01/2018	

Labeler - SC Johnson Professional USA, Inc. (607378015)

Establishment

Name	Address	ID/FEI	Business Operations
STERIS Corporation		139424188	MANUFACTURE(11084-806)

Establishment

Name	Address	ID/FEI	Business Operations
APEX International, Inc.		015226132	MANUFACTURE(11084-806)

Revised: 7/2021 SC Johnson Professional USA, Inc.