SCOTT FOAM HAND SANITIZER- benzalkonium chloride solution Kimberly-Clark

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.

SCOTT® FOAM HAND SANITIZER

Drug Facts

Active ingredient

Benzalkonium chloride 0.13 %

Purpose

Antiseptic skin cleanser

Uses

For personal hand hygiene to help prevent the spread of bacteria

Warnings

For external use only

When using this product avoid contact with eyes. If contact occurs, rinse with water

Stop use and ask a doctor if irritation develops

Keep out of reach of children. If swallowed, call a poison control centre or get medical help right away.

Directions

For occasional personal domestic use as needed. Supervise children when they use this product. Rub thoroughly into hands for at least 30 seconds. Allow to dry.

Inactive ingredients

Aloe Barbadensis Leaf Juice Powder, Citric Acid, Cocamidopropyl PG-Dimonium Chloride Phosphate, Methylpropanediol, Panthenol, Polyethylene Glycol, Silica, Sodium Hydroxide, Water

Questions?

Call 1-888-346-4652

Distributed in the U.S. by Kimberly-Clark Global Sales, LLC, Roswell, GA 30076-2199

PRINCIPAL DISPLAY PANEL - 40.5 fl oz Bottle Label

12979

Empty &

Discard Pump PLASTIC BOTTLE how2recycle.info

DIN 02481456

For Personal / Domestic Use Only

Scott® Brand

Foam Hand Sanitizer

Benzalkonium Chloride Solution, USP

Alcohol free

SAME GREAT FORMULATION. Now Scott® branded!

LEAVE ON

20-14-557-0-05

1.2 L (40.5 fl oz)

12979



DIN 02481456

For Personal / Domestic Use Only Produit exclusivement destiné à un usage personnel/domestique

Foam Hand Sanitizer

Désinfectant pour les mains en mousse

Benzalkonium Chloride Solution, USP Solution de chlorure de benzalkonium, USP

Alcohol free Sans alcohol



20-14-557-0-05

1.2 L (40.5 fl oz)

Drug Facts / Information sur le médicament

Active ingredient / Ingrédient actif

Purpose / Utilité

Benzalkonium chloride 0.13 % ..

Antiseptic skin cleanser

Uses / Usages

For personal hand hygiene to help prevent the spread of bacteria / Pour l'hygiène personnelle des mains pour aider à prévenir la propagation des bactéries

Warnings / Mises en garde

For external use only / Pour usage externe seulement

When using this product / Lorsque yous utilisez ce produit avoid contact with eyes. If contact occurs, rinse with water / évitez tout contact avec les yeux. En cas de contact, rincer avec de l'eau Stop use and ask a doctor if / Cessez d'utiliser ce produit et consultez un médecin si irritation develops / l'irritation se développe.

Keep out of reach of children. / Garder hors de la portée des enfants. If swallowed, call a poison control centre or get medical help right away. / En cas d'ingestion, appeler un centre antipoison ou obtenir une assistance médicale immédiate.

Directions / Mode d'emploi

For occasional personal domestic use as needed. Supervise children when they use this product. Rub thoroughly into hands for at least 30 seconds. Allow to dry. / Pour usage domestique personnel occasionnel au besoin. Superviser les enfants durant d'utiliser ce produit. Bien frotter le produit dans les mains pendant au moins 30 secondes. Laisser sécher.

Inactive ingredients / Ingrédients inactifs

Aloe Barbadensis Leaf Juice Powder, Citric Acid, Cocamidopropyl PG-Dimonium Chloride Phosphate, Methylpropanediol, Panthenol, Polyethylene Glycol, Silica, Sodium Hydroxide, Water / Jus en poudre de feuille d'Aloe Barbadensis, Acide citrique, Chlorophosphate de cocamidopropyl PG-dimonium, Méthylpropanediol, Panthénol, Poly(éthylène glycol), Silice, Hydroxyde de sodium, Eau

Questions? Call/Composez le 1-888-346-4652

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Mississauga, Ontario L5B 3Y5

www.kcprofessional.com

Re-order #: / Nº de commande : 12979

20-14-558-0-08





SCOTT FOAM HAND SANITIZER

benzalkonium chloride solution

Product Information	Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55118-664		
Route of Administration	TOPICAL				

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Benzalkonium Chloride (UNII: F5UM2KM3W7) (Benzalkonium - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	1.3 mg in 1000 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
CALCIUM (UNII: SY7Q814VUP)		
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)		
COCAMIDO PRO PYL PRO PYLENE GLYCOL-DIMO NIUM CHLO RIDE PHO SPHATE (UNII: H2KVQ74JM4)		
METHYLPRO PANEDIO L (UNII: N8 F53B3R4R)		
PANTHENOL (UNII: WV9CM0O67Z)		
POLYETHYLENE OXIDE 600000 (UNII: 2126FD486L)		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
Sodium Hydroxide (UNII: 55X04QC32I)		
WATER (UNII: 059QF0KO0R)		

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:55118-664-63	6 in 1 CARTON	11/16/2018			
1	NDC:55118-664-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product				
2	NDC:55118-664-65	2 in 1 CARTON	11/16/2018			
2	NDC:55118-664-12	1200 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	11/16/2018		

Labeler - Kimberly-Clark (830997032)

Revised: 10/2019 Kimberly-Clark