SCOTT ANTIMICROBIAL FOAM- 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® Antimicrobial Foam Soap

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.1% w/w

Purpose

Antiseptic cleanser

Use

For personal hand hygiene to help prevent the spread of certain bacteria.

Warnings

- **For External Use Only. Do not ingest.** If swallowed, get medical help or contact a Poison Control Center immediately.
- **Use with caution in children.**
- **Avoid contact with eyes. In contact occurs, flush eyes with water.**
- **Discontinue use & consult a health care practitioner** if irritation develops.
- **Keep out of reach of children.**
- **Do not use** if you are allergic to any ingredients.

Directions

Apply small amount and lather in hands for at least 30 seconds.

Adults: Supervise children when using this product. Do not dilute product. For occasional use as needed.

Other Information

- Report serious side effects from this product to 1-877-561-6587

Inactive Ingredients

Aloe Barbadensis Leaf Juice Powder, Citric Acid, Coco-Betaine, Glycerin, Panthenol, Sodium Benzoate, Sodium Chloride, Sodium Hydroxide, Tocopheryl Acetate, Water/Eau.

Questions?

1-888-346-4652

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

PRINCIPAL DISPLAY PANEL - 33.8 fl oz Bottle Label

Scott®
Brand

Antimicrobial Foam Soap

Benzalkonium
Chloride Solution, NF

For Domestic / Personal Care Use

Kills 99.9% of most common germs†

1 Liter
(33.8 fl oz)

DIN: 02458675

†Enterococcus faecalis VRE; MDR,
Enterococcus faecium, Pseudomonas aeruginosa,
Escherichia coli, Staphylococcus epidermidis,
Salmonella typhi, Streptococcus pyogenus,
Listeria monocytogenes, Proteus vulgaris,
Salmonella enterica

20-29-382-0-01

Scott® Brand

Antimicrobial Foam Soap
Savon En Mousse antimicrobien

Benzalkonium
Chloride Solution, NF

For Domestic / Personal Care Use
Destiné à un usage domestique/personnel

20-29-382-0-01

Kills 99.9% of most common germs†

†Enterococcus faecalis VRE; MDR, Enterococcus faecium, Pseudomonas aeruginosa, Escherichia coli, Staphylococcus epidermidis, Salmonella typhi, Streptococcus pyogenus, Listeria monocytogenes, Proteus vulgaris, Salmonella enterica

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- **Discontinue use & consult a health care practitioner** if irritation develops.
- **Keep out of reach of children.**
- **Do not use** if you are allergic to any ingredients.

Directions: Apply a small amount and lather hands with water for at least 30 seconds. Rinse Well. Adults: Supervise children when using this product. Do not dilute product. For occasional use as needed.

Other Information

- Report serious side effects from this product to 1-877-561-6587

Drug Facts (cont.)

Inactive Ingredients: Aloe Barbadensis Leaf Juice Powder, Citric Acid, Coco-Betaine, Glycerin, Panthenol, Sodium Benzoate, Sodium Chloride, Sodium Hydroxide, Tocopheryl Acetate, Water/Eau.

Questions? 1-888-346-4652

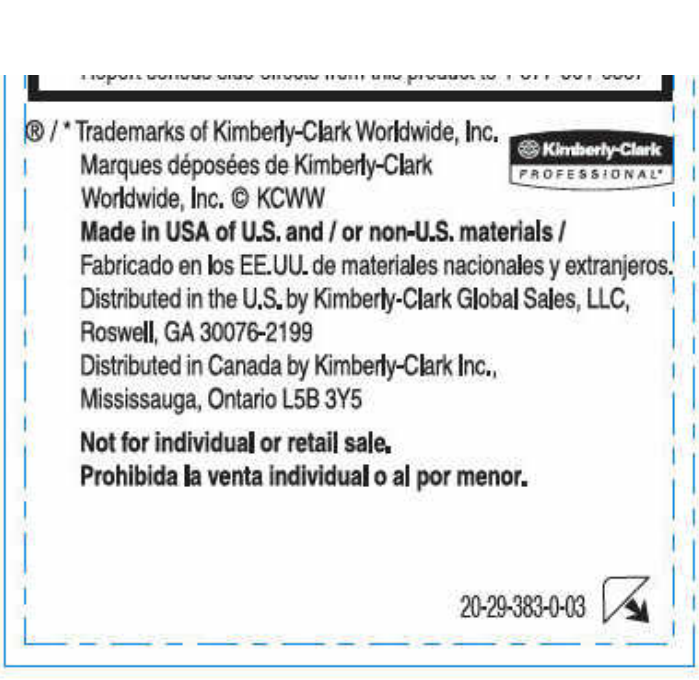
Au sujet de ce médicament

Ingrédient medicinal: ***Usage:***
Chlorure de benzalkonium 0,1% p/p.....Nettoyant antiseptique

Utilisation: Pour l'hygiène des mains afin de prévenir la propagation de certaines bactéries.

Mises en garde:

- **Pour usage externe seulement. Ne pas ingérer.** Si une personne ingère de ce produit, consulter un médecin ou communiquer avec un centre antipoison immédiatement.
- **Utiliser le produit avec précaution chez les enfants.**



SCOTT ANTIMICROBIAL FOAM

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55118-235
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Benzalkonium Chloride (UNII: F5UM2KM3W7) (Benzalkonium - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	1 mg in 1000 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Coco-Betaine (UNII: 03DH2IZ3FY)	
Glycerin (UNII: PDC6A3C0OX)	
Sodium Chloride (UNII: 451W47IQ8X)	
Sodium Benzoate (UNII: OJ245FE5EU)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
Aloe (UNII: V5VD430YW9)	
Panthenol (UNII: WV9CM0O67Z)	
.Alpha.-Tocopherol Acetate (UNII: 9E8X80D2L0)	
Sodium Hydroxide (UNII: 55X04QC32I)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
	NDC:55118-235			

1	NDC:55118-235-63	3 in 1 CARTON	12/01/2016	
1	NDC:55118-235-10	1000 mL in 1 CARTRIDGE; 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	12/01/2016	

Labeler - Kimberly-Clark (830997032)

Revised: 10/2019 Kimberly-Clark