KLEENEX ANTIMICROBIAL SKIN CLEANSER- benzalkonium chloride solution
Kimberly-Clark 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.

KLEENEX® Antimicrobial Foam Skin Cleanser

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Use

For handwashing to decrease bacteria on the skin.

Warnings

For External Use Only.

When using this product avoid contact with eyes; in case of contact, flush eyes with water.

Stop use & ask a doctor if irritation or redness develops or persists.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Wet hands and apply a palmful to hands. Scrub thoroughly for 15-20 seconds. Rinse and dry thoroughly.

Other Information

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

Inactive Ingredients

Water/Eau/Aqua, Cocamidopropylamine Oxide, Polysilicone-20, Cetrimonium Chloride, Di-PPG-2 Myreth-10 Adipate, Benzyl Alcohol, Polymethacrylamidopropyltrimonium Chloride, Tetrasodium Iminodisuccinic Acid, Citric Acid, Aminomethyl Propanol

Questions?

1-888-346-4652

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

PRINCIPAL DISPLAY PANEL - 1 Liter Bottle Label

***Kleenex*®**
BRAND

Antimicrobial Foam
Skin Cleanser

Triclosan Free
1 Liter (33.8 fl oz)

20-14-736-0-00

KLEENEX ANTIMICROBIAL SKIN CLEANSER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55118-440
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

Benzalkonium Chloride (UNII: F5UM2KM3W7) (Benzalkonium - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	0.013 mg in 1 L

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
Cocamidopropylamine Oxide (UNII: M4SL82J7HK)	
Cetrimonium Chloride (UNII: UC9PE95IBP)	
Di-PPG-2 Myreth-10 Adipate (UNII: 4IN301M0KJ)	
Benzyl Alcohol (UNII: LKG8494WBH)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
Aminomethylpropanol (UNII: LU49E6626Q)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55118-440-10	1 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:55118-440-12	1.2 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:55118-440-15	1.5 L in 1 BOTTLE, PLASTIC; 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	06/20/2013	

Labeler - Kimberly-Clark Corporation (006072136)

Revised: 11/2014 Kimberly-Clark Corporation