KLEENEX FOAM HAND SANITIZER- benzalkonium chloride solution
Kimberly-Clark

KLEENEX® Foam Hand Sanitizer

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%w/w

Purpose

Antiseptic cleanser

Use

For personal hand hygiene to help prevent the spread of bacteria

Warnings

For external use only. Do not ingest. Use with caution in children.

Do not use if you are allergic to any ingredients.

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

Apply small amount and rub thoroughly into hands for at least 30 seconds. Allow to dry. Not to be rinsed off with water (i.e. not to be used as a handwash). For occasional and personal domestic use. Supervise children when they use this product.

Other information

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

Inactive ingredients

Aloe Barbadensis Leaf Juice Powder, Citric Acid, Cocamidopropyl PG-Dimonium Chloride Phosphate, Methylpropanediol, Panthenol, PEG-14M, Silica, Sodium Hydroxide, Water

Questions?

1-888-346-4652

Distributed in the U.S. by Kimberly-Clark Global Sales, LLC, Roswell, GA 30076-2199
Distributed in Canada by Kimberly-Clark Inc., Mississauga, Ontario L5B 3Y5

PRINCIPAL DISPLAY PANEL - 532 mL Bottle Label

Kleenex®

Foam Hand Sanitizer

alcohol
free

DIN: 02443252

For Personal / Domestic Use Only

18 fl oz (532 mL)

Kleenex® BRAND MARQUE

Foam Hand Sanitizer
Désinfectant pour les mains en mousse

alcohol free
sans alcool

DIN 02443252

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

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Info-médicament

Ingrédient actif — **Utilité**
Chlorure de benzalkonium, 0,13 % p/p.....Nettoyant antiseptique

Usage
Pour l'hygiène des mains afin de prévenir la propagation des bactéries

Mode d'emploi Appliquer une petite quantité de ce produit et frotter vigoureusement sur les mains pendant au moins 30 secondes. Laisser sécher. Ne pas rincer le produit à l'eau (ce produit ne doit pas être utilisé comme un savon pour les mains). Pour usage domestique personnel et occasionnel. Superviser les enfants qui utilisent ce produit.

Autres renseignements
Signaler tout effet secondaire grave relatif à ce produit au 1-877-561-6587

Mises en garde
Pour usage externe seulement. Ne pas ingérer. Utiliser le produit avec précaution chez les enfants.

Cesser d'utiliser et consulter un médecin si une irritation ou une rougeur se produit et persiste. En cas de contact, rincer les yeux avec de l'eau.

Garder hors de la portée des enfants. En cas d'ingestion, appeler un centre antipoison ou consulter un médecin immédiatement.

Ne pas utiliser ce produit en cas d'allergie à l'un des ingrédients.

Signaler tout effet secondaire grave relatif à ce produit au 1-877-561-6587.

Ingrédients inactifs Poudre de jus d'aloès officinal, acide citrique, cocamidopropyl-PG-dimonium chloride phosphate, méthylpropanediol, panthénol, PEG-14M, silice, hydroxide de sodium, eau

Questions? 1-888-346-4652

Kimberly-Clark PROFESSIONAL

• Fragrance and Dye Free / Sans parfum ni colorant

Made in Mexico of U.S. and/or Non-US materials /
Fabriqué au Mexique à partir de matériaux américains et non américains /
Fabricado en México con materiales estadounidenses y/o no estadounidenses
Distributed in the U.S. by Kimberly-Clark Global Sales, LLC, Roswell, GA 30076-2199
Distributed in Canada by Kimberly-Clark Inc., Mississauga, Ontario L5B 3Y5
www.kcprofessional.com

0 36000 45827 5

Re-order # / N° de commande : 45827

20-14-861-0-02

KLEENEX FOAM HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55118-701
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
Water (UNII: 059QF0KO0R)	
Methylpropanediol (UNII: N8F53B3R4R)	
Cocamidopropyl Propylene Glycol-Dimonium Chloride Phosphate (UNII: H2KVQ74JM4)	
Polyethylene Oxide 600000 (UNII: 2126FD486L)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
Silicon Dioxide (UNII: ETJ7Z6XBU4)	
Aloe (UNII: V5VD430YW9)	
Panthenol (UNII: WV9CM0O67Z)	
Sodium Hydroxide (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End

#	Item Code	Package Description	Date	Date
1	NDC:55118-701-64	24 in 1 CARTON	09/01/2015	
1	NDC:55118-701-11	45 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		
2	NDC:55118-701-99	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/01/2015	
3	NDC:55118-701-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	09/01/2015	
4	NDC:55118-701-12	1200 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	09/01/2015	
5	NDC:55118-701-67	4 in 1 CARTON	09/01/2015	
5	NDC:55118-701-18	532 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	09/01/2015	

Labeler - Kimberly-Clark (830997032)

Establishment

Name	Address	ID/FEI	Business Operations
Tri-Pac Inc.		020844956	MANUFACTURE(55118-701) , LABEL(55118-701) , PACK(55118-701)

Establishment

Name	Address	ID/FEI	Business Operations
Cyan Labs		812754130	MANUFACTURE(55118-701) , LABEL(55118-701) , ANALYSIS(55118-701) , PACK(55118-701)

Revised: 12/2024 Kimberly-Clark