KLEENEX ULTRA MOISTURIZING HAND SANITIZER- alcohol 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® ULTRA Moisturizing Foam Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Uses

For personal hand hygiene to help prevent the spread of bacteria. Kills harmful bacteria or germs.

Warnings

For external use only

Flammability warning

Keep away from open flame and sources of heat.

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

Stop use and ask a health care practitioner if irritation or redness develops or persists.

Keep out of reach of children. If swallowed, get medical help or contact a poison control centre immediately.

Directions

Use enough foam to cover your hands. Rub hands together briskly until dry. Use as part of your daily cleansing routine.

Other information

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

Do not store above 110 °F (40 °C).

Inactive Ingredients

Aloe Barbadensis Leaf Extract, Betaine, Camellia Sensis Leaf Extract, Citric Acid, Cucumis Sativus Fruit Extract (Cucumber), Glycerin, Isopropanol, Meadowfoam Amidopropyl Betaine, Panthenol, PEG-10 Dimethicone, Water/Eau/Aqua

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 - 1L Bottle Label

34700

Kleenex® BRAND MARQUE ULTRA

Moisturizing Foam Hand Sanitizer

SAME GREAT FORMULATION. Now Kleenex® Branded

For Personal / Domestic Use Only

1L (33.8 fl oz)

Empty &
Discard Pump
PLASTIC
BOTTLE

how2recycle.info

NPN 80030706

ECOLOGO PRODUCT CERTIFIED FOR REDUCED ENVIRONMENTAL IMPACT. VIEW SPECIFIC ATTRIBUTES EVAUATED: UL.COM/EL UL 2784

20-14-833-0-00



34700

Moisturizing Foam **Hand Sanitizer**

Hydratant en mousse

Désinfectant pour les mains

SAME GREAT FORMULATION. Now Kleenex® Branded

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



1L (33.8 fl oz)



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Warnings For external use only

Flammability warning - Keep away from open

Au sujet des médicaments

Ingrédient actif Utilité Alcool éthylique, 70 % v/v... Nettoyant antimicrobien

Usages Pour l'hygiène des mains afin de prévenir la propagation des bactéries. Tue les bactéries ou les germes nu sibles.

Mises en garde Pour usage externe seulement flame and sources of heat.

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Questions? 1-888-346-4652



Avertissement - inflammabilité Tenir éloigné des flammes nues et des sources de chaleur.

Lorsque vous utilisez ce produit éviter tout contact avec les yeux. En cas de contact, rincer abondamment les yeux avec de l'eau.

Cesser d'utiliser et consulter un médecin si une irritation ou une rougeur se produit et persiste.

Garder hors de la portée des enfants.

En cas d'ingestion, consulter un médecin ou communiquer avec un centre antipoison immédiatement.

Instructions Couvrez entièrement vos mains de mousse. Frottez-vous vigoureusement les mains jusqu'à ce qu'elles soient sèches. Utilisez ce produit en faisant votre toilette quotidienne.

Autres renseignements

Signaler tout effet secondaire grave relatif à ce produit au 1-877-561-6587. Ne pas entreposer à plus de 40 °C (110 °F).

Ingrédients inactifs Extrait de feuilles d'aloès officinal, bétaine, extrait de feuilles de Camellia Sinensis, acide citrique, extrait des fruits du (Cucumis sativus), glycérine, isopropanol, bétaine amidopropy l à base d'huile d'écume des prés, panthénol, diméthicone PEG-10, eau

Questions ? 1-888-346-4652





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www.kcprofessional.com Re-order#/ No de commande : 34700

20-14-834-0-00

KLEENEX ULTRA MOISTURIZING HAND SANITIZER

alcohol solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:55118-530

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)

Alcohol (UNII: 3K9958V90M) Alcohol 70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength	
Water (UNII: 059QF0KO0R)		
Isopropyl Alcohol (UNII: ND2M416302)		
Betaine (UNII: 3SCV180C9W)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
GREEN TEA LEAF (UNII: W2ZU1RY8B0)		
Panthenol (UNII: W/9CM0O67Z)		
Glycerin (UNII: PDC6A3C0OX)		
MEADOW FOAMAMIDOPROPYL BETAINE (UNII: HNV0L650LG)		
Aloe Vera Leaf (UNII: ZY81Z83H0X)		
Cucumber (UNII: YY7C30VXJT)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:55118- 530-88	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2012		
2	NDC:55118- 530-99	472 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2012		
3	NDC:55118- 530-11	44 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2012		
4	NDC:55118- 530-10	1000 mL in 1 BAG; Type 0: Not a Combination Product	03/01/2012		
5	NDC:55118- 530-12	1200 mL in 1 BAG; Type 0: Not a Combination Product	03/01/2012		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	03/01/2012	

Labeler - Kimberly-Clark Corporation (830997032)

Establishment			
Name	Address	ID/FEI	Business Operations
Tri-Pac Inc.		020844956	MANUFACTURE(55118-530), LABEL(55118-530), PACK(55118-530)

Establishment			
Name	Address	ID/FEI	Business Operations
Cyan Labs		812754130	MANUFACTURE(55118-530) , LABEL(55118-530) , ANALYSIS(55118-530) , PACK(55118-530)