SCOTT MOISTURIZING FOAM HAND SANITIZER (ULTRA)- alcohol 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® Moisturizing Foam Hand Sanitizer (Ultra)

Drug Facts

Active Ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Use

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

Warnings

Flammable - Keep away from open flame and sources of heat.

For External Use Only.

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

Stop use & 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 Center 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 Sinensis Leaf Extract, Citric Acid, Cucumis Sativus (Cucumber) Fruit Extract, Glycerin, Isopropanol, Meadowfoamamidopropyl Betaine, Panthenol, PEG-10 Dimethicone, Water/Eau

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 - 33.8 fl oz Bottle Label

34700

SAME GREAT FORMULATION. Now Scott® branded!

 $Scott^{\mathbb{R}}$

Brand

Moisturizing

Foam Hand Sanitizer

ULTRA Leave On

Empty & Discard Pump PLASTIC BOTTLE

20-14-658-0-03

how2recycle.info

ECOLOGO PRODUCT CERTIFIED FOR REDUCED ENVIRONMENTAL IMPACT. VIEW SPECIFIC ATTRIBUTES EVAUATED: UL.COM/EL UL 2783

NPN: 80030706

For Personal / Domestic Use Only

1 Liter (33.8 fl oz)

34700



SAME GREAT FORMULATION. Now Scott® branded!

Moisturizing Foam Hand Sanitizer Désinfectant hydratant en mousse pour les mains







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Au sujet des médicaments

Ingrédient médicinal Usage Alcohol éthylique, 70 % v/v....Antiseptique

Utilisation Pour l'hygiène des mains afin de prévenir la propagation de certaines bactéries. Tue les bactéries ou les germes néfastes.

Avertissements INFLAMMABLE. Tenir éloigné des flammes nues et des sources de chaleur. Pour usage externe seulement

Lorsque vous utilisez ce produit, évitez tout contact avec les yeux. En cas de contact, rincez les yeux à l'eau.

Arrêtez l'utilisation et consultez un practitien de la santé si une irritation ou Poison Control Center immediately.

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⊕ Kimberly-Clark

PROFESSIONAL™

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Etats-Unis à partir de matériaux provenant des Etats-Unis ou
d'aileurs, Distributed in the US. by Kimberly-Clark Globel Sales, LLC,
Rosvell, GA 30076-2199. Distributed in Carrada by Kimberly-Clark Inc.
Messissausa, Ordano LSB 375. www.kcprofessional.com
Re-order #/ N° de commande: 34700

une rougeur se produit ou persiste.

Tenir loin de la portée des enfants. Si une personne ingère de ce produit, consultez un médecin ou communiquez 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 dans le cadre de votre toilette quotidienne.

Autre Information Signalez tous les effets secondaires graves relatif à ce produit au 1-877-561-6587. Ne pas entreposer à une température supérieure à 40 °C (110 °F).

Ingrédients non médicinaux : voir ci-dessus

- · Fragrance and Dye Free / Sans parfum ni colorant
- Latex, Nitrile, and Vinyl glove compatible / Compatible avec les gants en latex, en nitrile et en vinyle
- CHG compatible / Compatible avec le CHG



SCOTT MOISTURIZING FOAM HAND SANITIZER (ULTRA)

alcohol solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:55118-532

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			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
Betaine (UNII: 3SCV180C9W)				
GREEN TEA LEAF (UNII: W2ZU1RY8B0)				
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)				
CUCUMBER (UNII: YY7C30 VXJT)				
Glycerin (UNII: PDC6A3C0OX)				
ISOPROPYL ALCOHOL (UNII: ND2M416302)				
MEADO WFO AMAMIDO PRO PYL BETAINE (UNII: HNV0 L650 LG)				
Panthenol (UNII: WV9CM0O67Z)				

PEG-10 DIMETHICO NE (600 CST) (UNII: 8 PR7V1SVM0)

Water (UNII: 059QF0KO0R)

]	Packaging					
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
:	NDC:55118-532-63	6 in 1 CARTON	04/25/2018			
	NDC:55118-532-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product				
2	NDC:55118-532-65	2 in 1 CARTON	04/25/2018			
2	NDC:55118-532-12	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	04/25/2018				

Labeler - Kimberly-Clark (830997032)

Revised: 10/2019 Kimberly-Clark