KLEENEX HAND SANITIZER, GREEN CERTIFIED- 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 Foam Hand Sanitizer, Green Certified

Drug Facts

Active Ingredient

Ethyl Alcohol 70%

Purpose

Antimicrobial

Use

Hand sanitizer to decrease bacteria on the skin.

Warnings

Flammable - Keep product away from fire or flame. 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 and 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.

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 Oleifera Leaf Extract, Citric Acid, Cucumis Sativus (Cucumber) Fruit Extract, Isopropanol, Glycerin, Meadowfoamamidopropyl Betaine, Panthenol, PEG-10 Dimethicone, Water

Questions?

1-888-346-4652

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

PRINCIPAL DISPLAY PANEL - 236 mL Bottle Label

Kleenex®
BRAND

Foam
Hand Sanitizer

green
certified

CERTIFIED
EcoLogo®
Certified Instant
Hand Antiseptic
CCD-170

8 fl oz (236 mL)

20-14-611-0-00

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Kimberly-Clark PROFESSIONAL®

- Fragrance and Dye Free
- No Fluorinated Chemicals
- Contains bio-based Ethanol

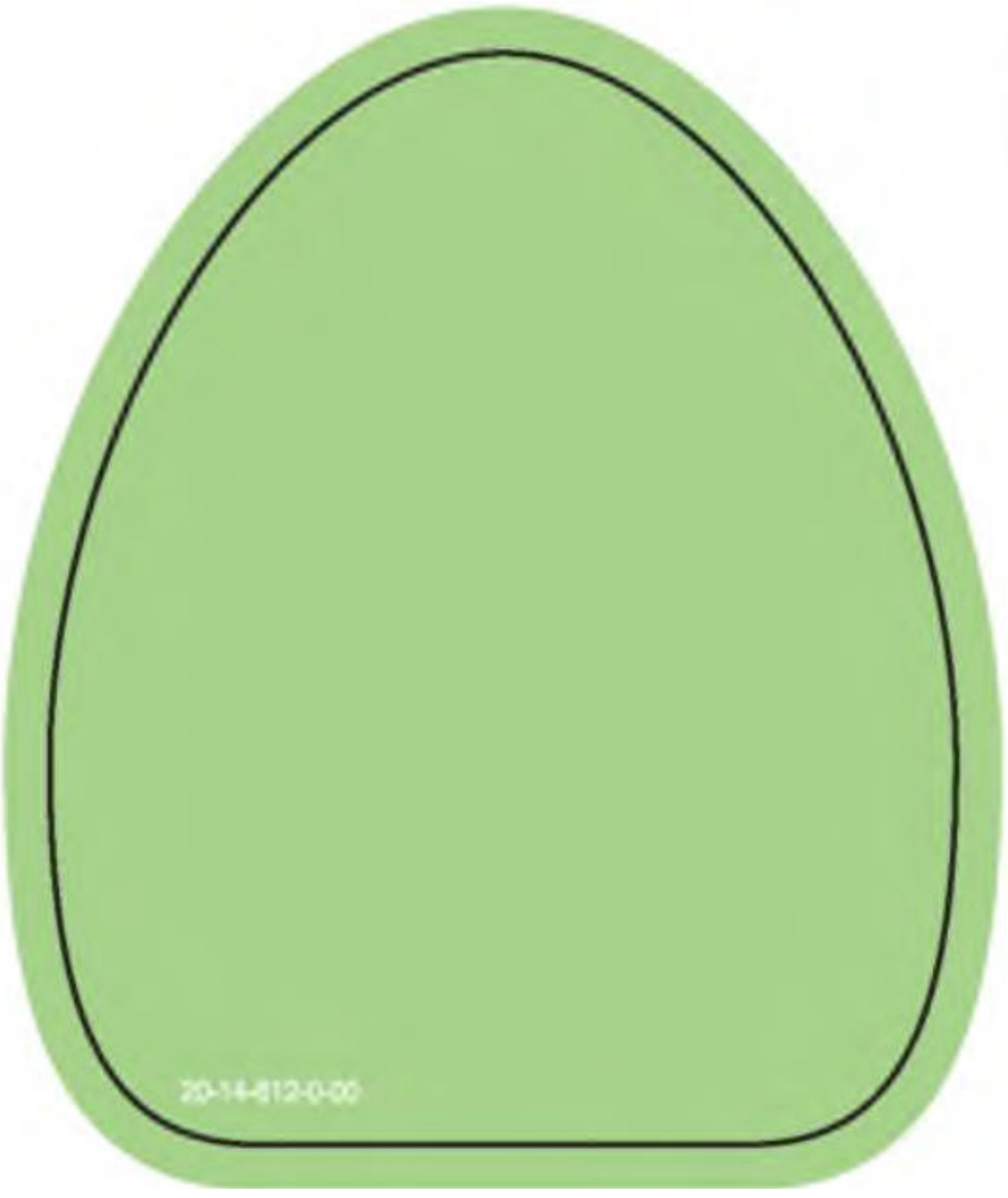
Re-order #: 33947

Made in USA of U.S. and / or non-U.S. materials

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

www.kcprofessional.com

0 36000 33947 5



KLEENEX HAND SANITIZER, GREEN CERTIFIED

alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55118-535
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
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)	
Panthenol (UNII: WV9CM0O67Z)	
Glycerin (UNII: PDC6A3C0OX)	
Aloe Vera Leaf (UNII: ZY81Z83H0X)	
Camellia Oleifera Leaf (UNII: 5077EL0C60)	
Cucumber (UNII: YY7C30VXJT)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55118-535-88	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:55118-535-99	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:55118-535-11	45 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:55118-535-10	1000 mL in 1 BAG; Type 0: Not a Combination Product		
5	NDC:55118-535-12	1200 mL in 1 BAG; 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	03/01/2012	

Labeler - Kimberly-Clark Corporation (006072136)

Revised: 11/2014 Kimberly-Clark Corporation