KLEENEX MOISTURIZING INSTANT 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.

MARQUE
kleenex®
BRAND
MOISTURIZING INSTANT HAND SANITIZER

Drug Facts

Active Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

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 right away.

Directions

Wet hands thoroughly with product and allow to 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

Water, Glycerin, Dimethicone, Carbomer, Petrolatum, Aminomethyl Propanol, Fragrance, Tocopheryl Acetate, Panthenol, Hydroxypropylcellulose, Ceteth-10, Steareth-21, Poloxamer 335, Aloe Barbadensis Leaf

Questions?

1-888-346-4652

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

www.kcprofessional.com

PRINCIPAL DISPLAY PANEL - 1.5 mL Pouch Label

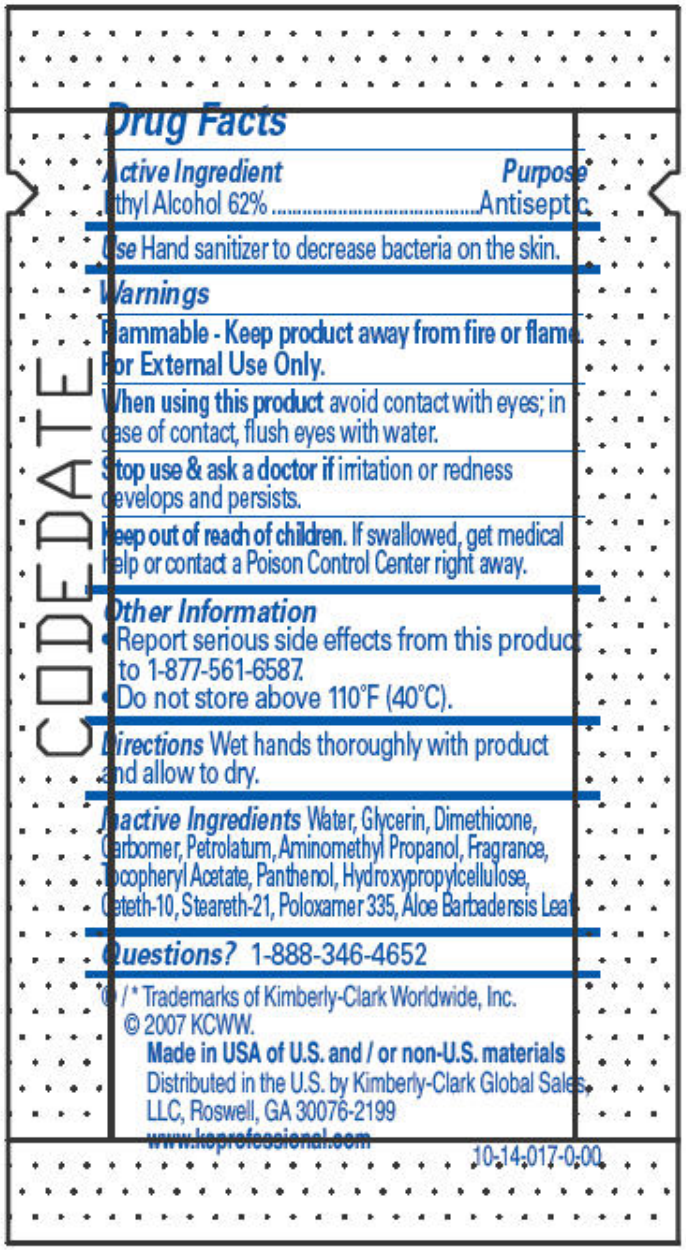
MARQUE
Kleenex*
BRAND

Moisturizing Instant
Hand Sanitizer

Contains Aloe & Vitamin E

Single Use

.05
Fl Oz
(1.5 mL)



KLEENEX MOISTURIZING INSTANT HAND SANITIZER

alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55118-480
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	
Dimethicone (UNII: 92RU3N3Y1O)	
Petrolatum (UNII: 4T6H12BN9U)	
Aminomethylpropanol (UNII: LU49E6626Q)	
Panthenol (UNII: WV9CM0O67Z)	
Hydroxypropyl Cellulose (Type H) (UNII: RFW2ET671P)	
Ceteth-10 (UNII: LF9X1PN3XJ)	
Steareth-21 (UNII: 53J3F32P58)	
Poloxamer 335 (UNII: G6DQL26D50)	
Aloe Vera Leaf (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55118-480-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product		
2	NDC:55118-480-88	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:55118-480-22	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:55118-480-12	1200 mL in 1 CARTRIDGE; Type 0: Not a Combination Product		
5	NDC:55118-480-55	15 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
6	NDC:55118-480-11	30 mL 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	03/01/2011	

Labeler - Kimberly-Clark Corporation (006072136)

Establishment

Name	Address	ID/FEI	Business Operations
Marietta Corporation		010765394	MANUFACTURE(55118-480) , ANALYSIS(55118-480) , RELABEL(55118-480)

Revised: 9/2014 Kimberly-Clark Corporation