

KLEENEX 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.

KLEENEX® INSTANT HAND SANITIZER

Drug Facts

Active Ingredient

Ethyl Alcohol 68% v/v

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, Carbomer, Fragrance, Aminomethyl Propanol, Glycerin

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
MARQUE

Instant Hand
Sanitizer

kills
germs†

8 fl oz (236 mL)

20-14-141-0-09

Kleenex[®]
BRAND
MARQUE

Instant Hand Sanitizer

kills
germs*

8 fl oz (236 mL)

20-14-141-0-08

Drug Facts

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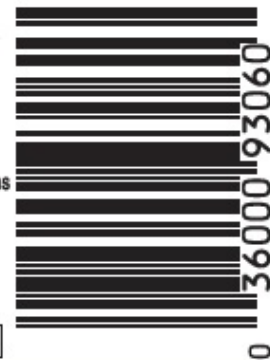
 **Kimberly-Clark**
PROFESSIONAL

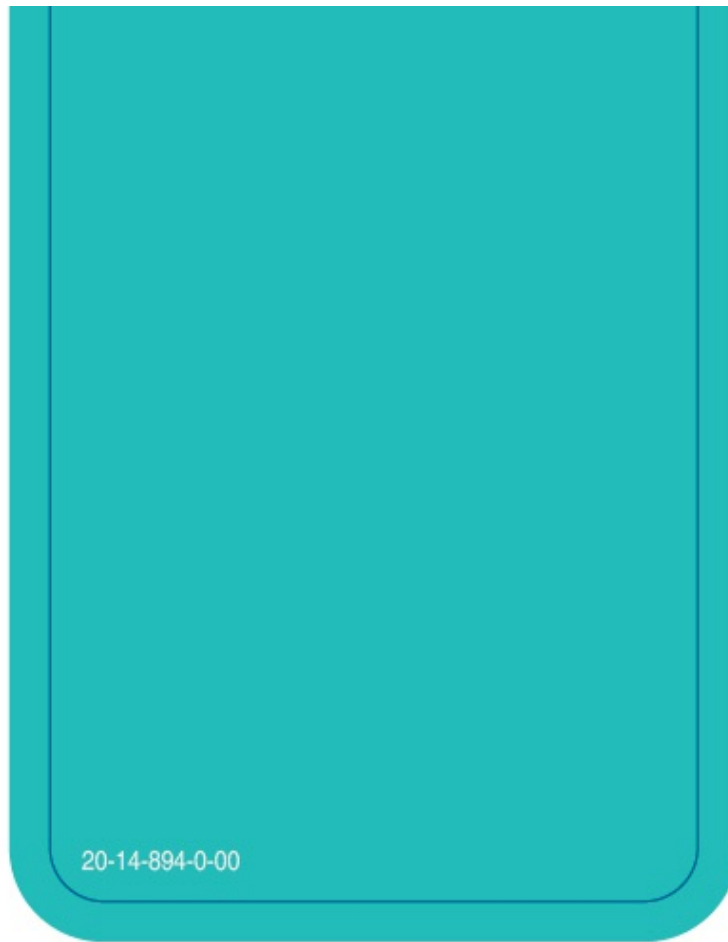
*Kills 99.99% of most common germs that make you sick.

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Distributed in the U.S. by Kimberly-Clark Global Sales, LLC, Roswell, GA 30076-2199
www.kcprofessional.com
Re-order #: 93060



NSF
Hardwood Compounds
Program Listed E3
337322





KLEENEX INSTANT HAND SANITIZER

alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55118-704
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	68 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55118-704-70	6 in 1 CARTON	08/30/2021	
1	NDC:55118-704-88	236 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	08/30/2021	

Labeler - Kimberly-Clark Corporation (830997032)

Revised: 7/2023

Kimberly-Clark Corporation