KLEENEX REVEAL MOISTURIZING FOAM HAND SANITIZER- alcohol solution
Kimberly-Clark

Kleenex® Reveal™ 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 doctor 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. Supervise children when they use this product. For occasional and personal domestic use. Rub thoroughly into hands for at least 30 seconds. Allow to 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 - 532 mL Bottle Label

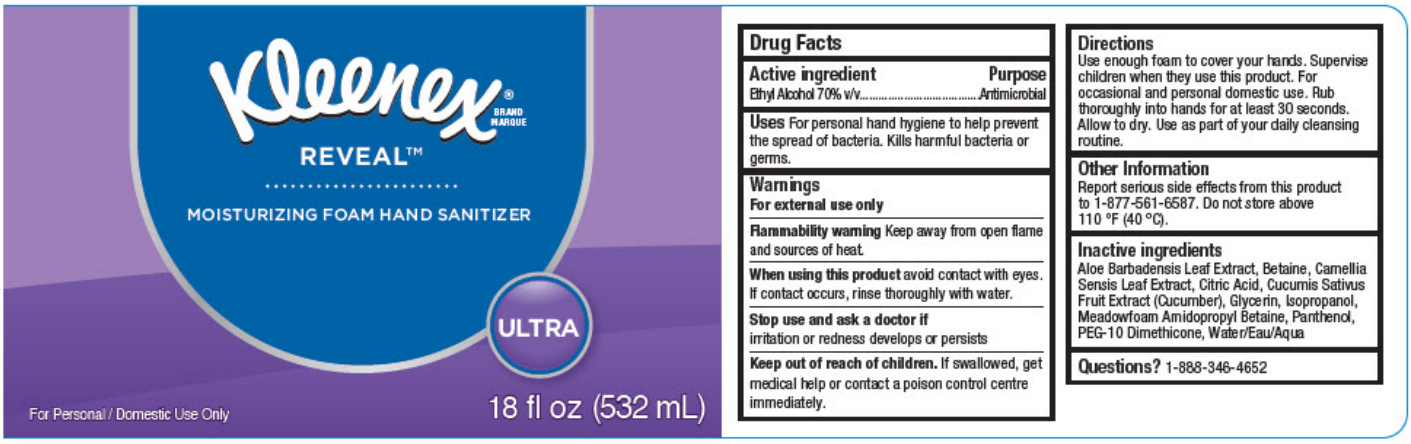
Kleenex®
BRAND
MARQUE

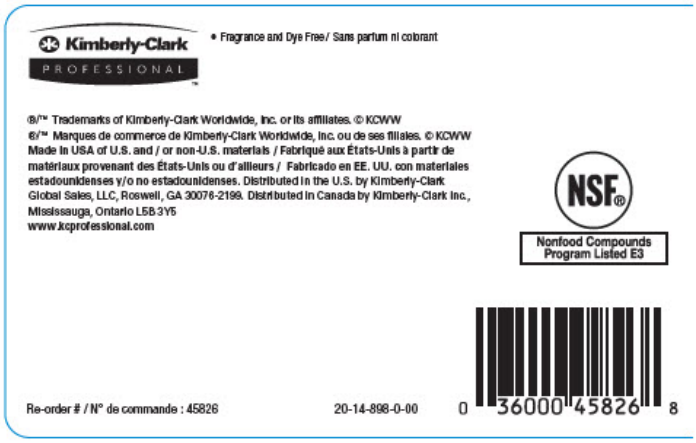
REVEAL™

MOISTURIZING FOAM HAND SANITIZER

ULTRA

For Personal / Domestic Use Only
18 fl oz (532 mL)





KLEENEX REVEAL MOISTURIZING FOAM HAND SANITIZER

alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55118-531
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**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
Betaine (UNII: 3SCV180C9W)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CUCUMBER (UNII: YY7C30VXJT)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
Glycerin (UNII: PDC6A3C0OX)	
Meadowfoamamidopropyl Betaine (UNII: HNV0L650LG)	
Panthenol (UNII: WV9CM0O67Z)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
Water (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55118-531-64	24 in 1 CARTON	05/15/2018	
1	NDC:55118-531-11	45 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:55118-531-67	4 in 1 CARTON	05/15/2018	
2	NDC:55118-531-18	532 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:55118-531-10	6 in 1 CARTON	06/01/2022	
3		1000 mL in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:55118-531-12	2 in 1 CARTON	06/01/2022	
4		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 drug	M003	05/15/2018	

Labeler - Kimberly-Clark (830997032)

Revised: 12/2024 Kimberly-Clark