

**SOFT CARE INSTANT HAND SANITIZER AF- benzalkonium chloride solution
Diversey, Inc.**

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Uses:

For hands to decrease bacteria on the skin.

Recommended for repeated use.

Warnings

For external use only.

Do not ingest

Do not use in the eyes.

When using this product do not get into the eyes.

In case of contact, immediately flush with water.

Stop use and ask a doctor if irritation and redness develop.

If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Do not use if you are allergic to any ingredients.

Directions:

Apply 0.4-1 ml into hands.

Rub product vigorously over hands covering all surfaces.

Allow to air dry - do not rinse.

Other information:

EMERGENCY PHONE: 1-800-851-7145

See SDS MS0881162

FOR COMMERCIAL USE

See container for Lot Code and Expiry Date.

Store in a cool dry place.

Inactive Ingredients:

Water, Cocamidopropyl betaine, Propylene glycol, Phenoxyethanol, PEG-7 Glyceryl cocoate, Fragrance, Tetrasodium EDTA, Aloe barbadensis leaf juice, Tocopheryl acetate

Questions or comments?

1-800-558-2332 Monday through Friday 7:30 AM to 5:00 PM Central Standard Time

www.sealedair.com



Soft Care Instant Hand Sanitizer AF

Alcohol-Free Foam Hand Sanitizer

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Manufactured for:
©2018 Diversey, Inc., PO Box 19747,
Charlotte, NC 28219-0747 U.S.A.

300200708 (18/047)

NDC 64536-3628-3

Diversey

SoftCare

Instant Hand

Sanitizer AF

Alcohol-Free Foam Hand Sanitizer

(L) Exp.

NSF

Nonfood Compounds

Program Listed E3

Registration # 156292

Net Contents

1.3 L/ 1.37 U.S. Qt,

SKU: 100961733

SOFT CARE INSTANT HAND SANITIZER AF

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64536-3628
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
EDETATE SODIUM (UNII: MP1J8420LU)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64536-3628-3	1300 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/14/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/14/2017	

Labeler - Diversey, Inc. (018240817)

Establishment

Name	Address	ID/FEI	Business Operations
KUTOL PRODUCTS COMPANY		004236139	manufacture(64536-3628)

Revised: 12/2024

Diversey, Inc.