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

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.

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.

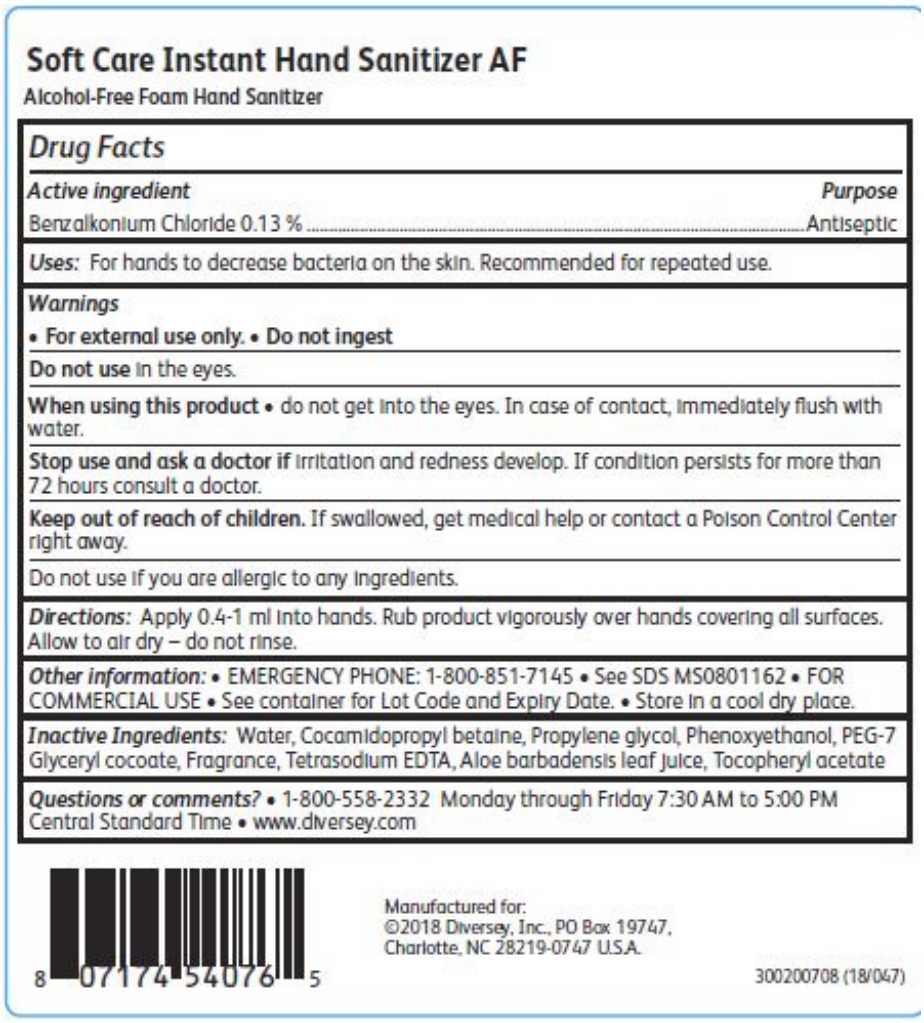
Inactive Ingredients:

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



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 not final	part333A	04/14/2017	

Labeler - Diversey, Inc. (018240817)

Establishment

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