CLEANSE- benzalkonium chloride solution
Medline Industries, LP

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184 Antibacterial Foaming Hand Soap

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antibacterial

Use

for handwashing to decrease bacteria on the skin

Warnings

For external use only: hands only

When using this product
- avoid contact with the eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if
- irritation and redness develops
- condition persists for more than 72 hours

Keep out of reach of children.

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

Directions
- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

Inactive ingredients

benzophenone-4, citric acid, cocamidopropyl betaine, fragrance, glycerin, lauramidopropylamine oxide, lauramine oxide, myristamidopropylamine oxide, red 4, sodium benzoate, tetrasodium EDTA, water, yellow 5

Manufacturing Information

www.medline.com

Made in USA of foreign and domestic materials

Manufactured for Medline Industries, LP

Three Lakes Drive, Northfield, IL 60093 USA

1-800-MEDLINE

REF: HHABSP1200F

Package Label

CLEANSE

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53329-184
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)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
EDETATE SODIUM (UNII: MP1J8420LU)	
GLYCERIN (UNII: PDC6A3C0OX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
SULISOBENZONE (UNII: 1W6L629B4K)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:53329-184-84	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/26/2016	
2	NDC:53329-184-08	221 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2021	
3	NDC:53329-184-74	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/26/2016	

Labeler - Medline Industries, LP (025460908)

Registrant - Medline Industries, LP (025460908)

Revised: 2/2026 Medline Industries, LP