

CLEANSE- benzalkonium chloride 0.13% solution
Medline Industries, LP

177 Antibacterial 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 or 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, disteareth-75 IPDI, fragrance, glycerin, lauramidopropylamine oxide, lauramine oxide, myristamidopropylamine oxide, PEG-150 distearate, red 40, red 33, sodium benzoate, sodium chloride, tetrasodium EDTA, water, yellow 5

Manufacturing Information

Manufactured for:

Medline Industries, LP

Three Lakes Drive, Northfield, IL 60093, USA

Made in USA with domestic and foreign materials

www.medline.com

1-800-MEDLINE (633-5463)

REF: HHABSP1200G

V2 RB23VJO

Package Label





L0020798BA

REF HHABSP1200G

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Drug Facts

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CLEANSE

benzalkonium chloride 0.13% solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53329-177
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
SODIUM BENZOATE (UNII: OJ245FE5EU)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
SULISOBENZONE (UNII: 1W6L629B4K)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
EDETATE SODIUM (UNII: MP1J8420LU)	
GLYCERIN (UNII: PDC6A3C0OX)	
DISTEARETH-75 ISOPHORONE DIISOCYANATE (UNII: 5365FJ30SC)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
WATER (UNII: 059QF0KO0R)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53329-177-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/19/2016	
2	NDC:53329-177-08	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/19/2016	
3	NDC:53329-177-84	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/19/2016	
4	NDC:53329-177-25	3790 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/19/2016	
5	NDC:53329-177-06	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/19/2016	
6	NDC:53329-177-74	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)	08/19/2016	

Labeler - Medline Industries, LP (025460908)

Registrant - Medline Industries, LP (025460908)

Revised: 5/2024

Medline Industries, LP