DEBMED ANTIMICROBIAL BODY WASH AND SHAMPOO - triclosan liquid
Deb USA, 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

Triclosan, 0.70%

Purpose

Antimicrobial

Uses

For hand and body washing to reduce bacteria on the skin

Warnings

For external use only

When using this product avoid contact with eyes. In case of eye contact, flush with water

Keep out of reach of children

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

Directions

Apply product onto washcloth or directly onto skin

Lather and wash for 15-20 seconds

Rinse and dry thoroughly

Inactive ingredients

Water, Disodium Laureth Sulfosuccinate, Sodium Laureth Sulfate, TEA-Dodecylbenzene Sulfonate, Citric Acid, Phenoxyethanol, Hyroxypropyl Guar Hydroxypropyltrimonium Chloride, Aloe Barbadensis Leaf Juice, Sodium Benzoate, Allantoin, Sodium Chloride, Tetrasodium EDTA, Potassium Sorbate, Fragrance, Yellow 5 (CI19140), Red 40 (CI 16035), Blue 1 (CI 42090).

DebMed Engineering Hand Hygiene Compliance

AntiMicrobial Body Wash and Shampoo

1L

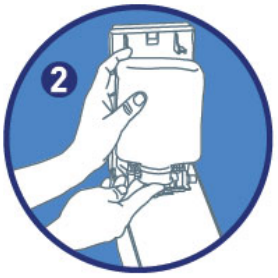
33.8 fl oz

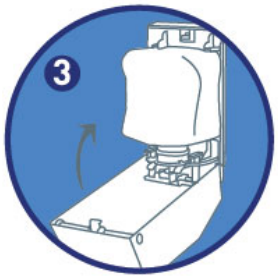
Made in USA

DMABHB1L

deb med
Engineering Hand Hygiene Compliance™

AntiMicrobial BODY WASH & SHAMPOO™





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2

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MADE IN USA
Deb,
Stanley, NC 28164
1-800-248-7190
www.debgroup.com

DMABHB1L AntiMicrobial Body Wash & Shampoo Rev. 07-11

DEBMED ANTIMICROBIAL BODY WASH AND SHAMPOO

triclosan liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-061
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Triclosan (UNII: 4NM5039Y5X) (Triclosan - UNII:4NM5039Y5X)	Triclosan	0.70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DISODIUM LAURETH SULFOSUCCINATE (UNII: D6DH1DTN7E)	
TEA-DODECYLBENZENESULFONATE (UNII: 8HM7ZD48HN)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE SODIUM (UNII: MP1J8420LU)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GUAR GUM (UNII: E89I1637KE)	
ALLANTOIN (UNII: 344S277G0Z)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-061-27	1000 mL in 1 BOTTLE, PLASTIC		
2	NDC:11084-061-38	222 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/01/2011	

Labeler - Deb USA, Inc. (607378015)

Establishment

Name	Address	ID/FEI	Business Operations
Deb USA, Inc.		607378015	manufacture

Establishment

Name	Address	ID/FEI	Business Operations
Deb Worldwide Healthcare Inc.		205662831	manufacture

Revised: 7/2011

Deb USA, Inc.