DEB MED ANTIMICROBIAL FOAMING HAND WASH FRAGRANCE FREE DYE FREE- 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 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 foaming cleanser to hands

Rub hands together to spread lather

Wash for 15-20 seconds

Rinse and dry hands thoroughly

Inactive ingredients

Water, Propyl Alcohol, Butylene Glycol, Sodium Laureth Sulfate, Disodium Laureth Sulfosuccinate, Cocamidopropyl Betaine, Ethylhexylglycerin, Benzyl Alcohol, Phenoxyethanol, Polyglycerin-6, Citric Acid, DMDM Hydantoin, Cocamidopropyl PG-Dimonium Chloride Phosphate, Dihydroxypropyl PEG-5 Linoleammonium Chloride, Tetrasodium EDTA, Allantoin, Methylchloroisothiazolinone, Methylisothiazolinone.

deb med Engineering Hand Hygiene Compliance

AntiMicrobial Foaming Hand Wash with Microhydration

Meets Protocol For Healthcare Personnel Handwash

400 mL

13.5 fl. oz.

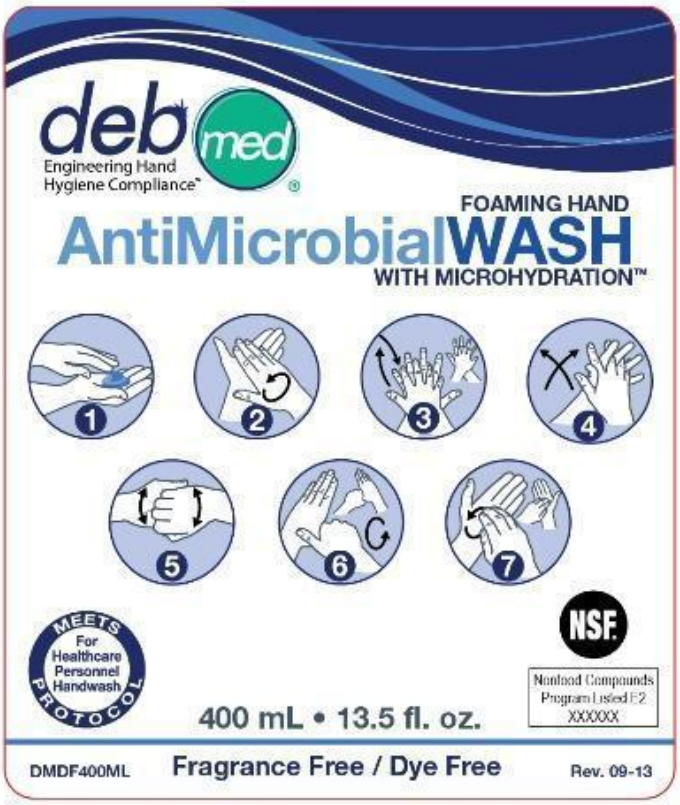
Made in USA

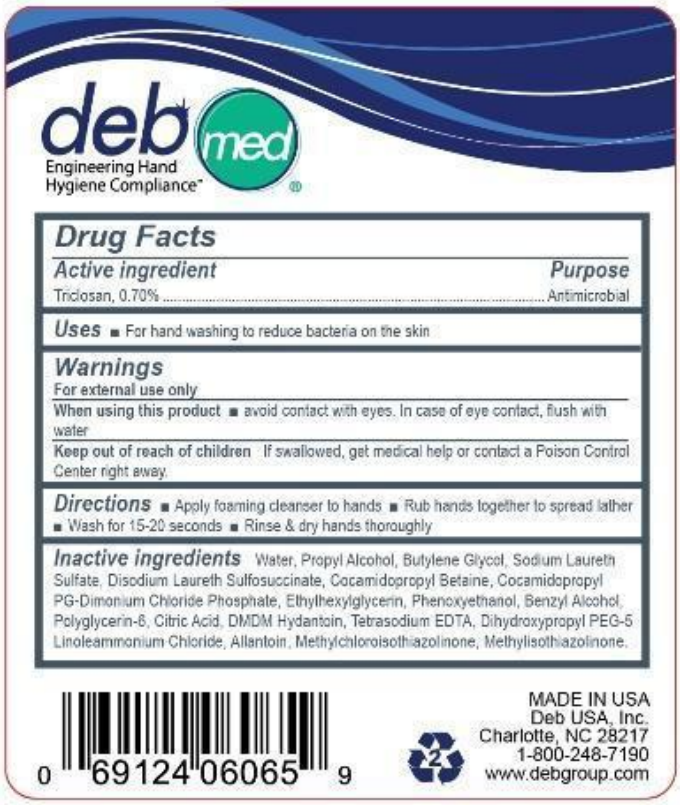
www.debgroup.com

DMDF400ML

Fragrance Free Dye Free

Rev. 09-13





DEB MED ANTIMICROBIAL FOAMING HAND WASH FRAGRANCE FREE DYE FREE

triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-266
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	7 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYL ALCOHOL (UNII: 96F264O9SV)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
DISODIUM LAURETH SULFOSUCCINATE (UNII: D6DH1DTN7E)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
POLYGLYCERIN-6 (UNII: M51422LRAM)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
EDETATE SODIUM (UNII: MP1J8420LU)	
DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	
ALLANTOIN (UNII: 344S277G0Z)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-266-40	400 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/15/2013	

Labeler - Deb USA, Inc. (607378015)

Establishment

Name	Address	ID/FEI	Business Operations
Deb USA, Inc.		607378015	manufacture(11084-266)

Revised: 9/2013 Deb USA, Inc.