

LYSOL TOUCH OF FOAM- benzalkonium chloride solution
Reckitt Benckiser LLC

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.

Lysol®
Touch of Foam

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.10%

Purpose

Antibacterial

Uses

for handwashing to decrease 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.

Stop use and ask a doctor if irritation or redness develops.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply product to hands.
- Wash hands.
- Rinse hands with water.

Other Information

store at room temperature

Inactive Ingredients

Water, Glycerin, Lauramine Oxide, PEG-150 Distearate, Cetrimonium Chloride, Propylene Glycol, Fragrance, Citric Acid, Tetrasodium EDTA, Cocamide MEA, Hydrolyzed Collagen, Magnesium Nitrate, Butylene Glycol, Phenoxyethanol, Methylchloroisothiazolinone, Magnesium Chloride, Ethylhexylglycerin, Methylisothiazolinone, D&C Green 5, FD&C Yellow 5.

Questions? Comments?

Call 1-800-228-4722

Distributed by:
Reckitt Benckiser
Parsippany, NJ 07054-0224

PRINCIPAL DISPLAY PANEL - 251 mL Bottle Label

NEW FORMULA!

Lysol[®]
BRAND
KILLS 99.9% OF BACTERIA

Soft on Skin,
HARD ON GERMS[®]

Antibacterial
Foam Hand Wash


with
MOISTURIZING
COLLAGEN
COMPLEX

8.5 FL. OZ.
(251 mL)

Refreshing Citrus Splash

8170171



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INSERT 100% ZERO SUPR. TRUNCATED UPC 0 193362 0	

LYSOL TOUCH OF FOAM

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:638 24-485
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Benzalkonium Chloride (UNII: F5UM2KM3W7) (Benzalkonium - UNII:7N6JUD5X6 Y)	Benzalkonium Chloride	0.1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAURAMINE O XIDE (UNII: 4F6FC4M18 W)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

EDETATE SODIUM (UNII: MP1J8420LU)
CO CO MONOETHANOLAMIDE (UNII: C80684146D)
MAGNESIUM NITRATE (UNII: 77CBG3UN78)
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63824-485-01	251 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	10/08/2015	

Labeler - Reckitt Benckiser LLC (094405024)