

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, Methylchloroisothiazolinone, Methylisothiazolinone, D&C Red No. 33, FD&C Yellow No. 5.

Questions? Comments?

Call 1-800-228-4722

Distributed by: Reckitt Benckiser LLC
Parsippany, NJ 07054-0224
Made in U.S.A.

PRINCIPAL DISPLAY PANEL - 251 mL Bottle Label

new

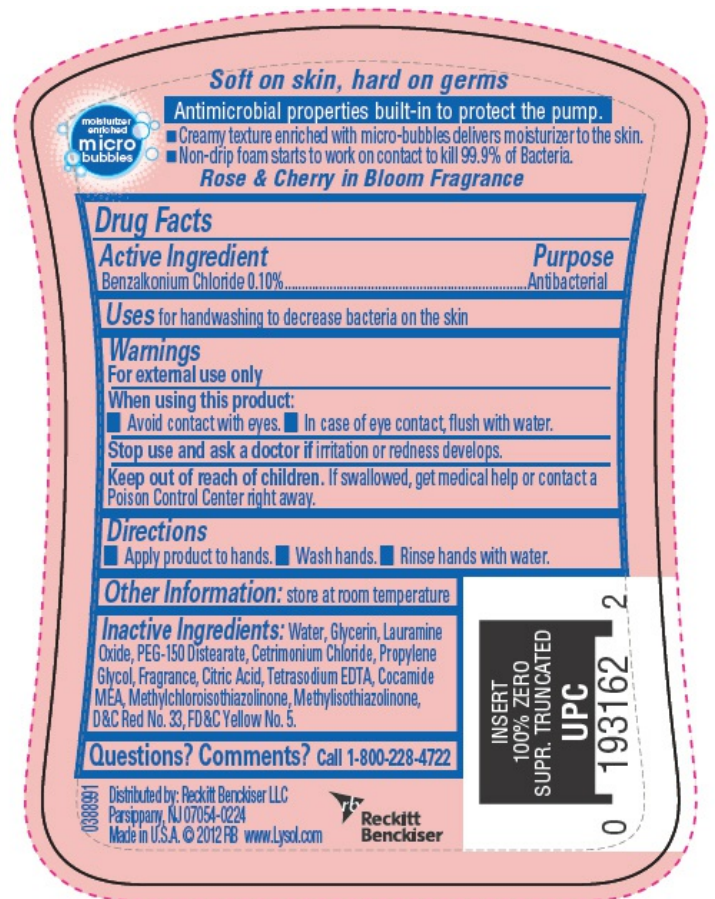
Lysol®
BRAND
KILLS 99.9% OF BACTERIA

Touch of
Foam™
ANTIBACTERIAL
HAND WASH

8.5 FL. OZ.
(251 mL)

rose &
cherry
in bloom

0388990



benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63824-465
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Benzalkonium Chloride (UNII: F5UM2KM3W7) (Benzalkonium - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	0.1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	
Lauramine Oxide (UNII: 4F6FC4M8W)	
PEG-150 Distearate (UNII: 6F36Q0I0AC)	
Cetrimonium Chloride (UNII: UC9PE95IBP)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
Edetate Sodium (UNII: MP1J8420LU)	
Coco Monoethanolamide (UNII: C80684146D)	
Methylchloroisothiazolinone (UNII: DEL7T5QRPN)	
Methylisothiazolinone (UNII: 229D0E1QFA)	
D&C Red No. 33 (UNII: 9DBA0SBB0L)	
FD&C Yellow No. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63824-465-08	251 mL in 1 BOTTLE, PUMP		
2	NDC:63824-465-11	325 mL in 1 BOTTLE, PUMP		
3	NDC:63824-465-25	739 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	12/26/2012	

Labeler - Reckitt Benckiser LLC (094405024)