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 Green No. 5, D&C Red No. 33.

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

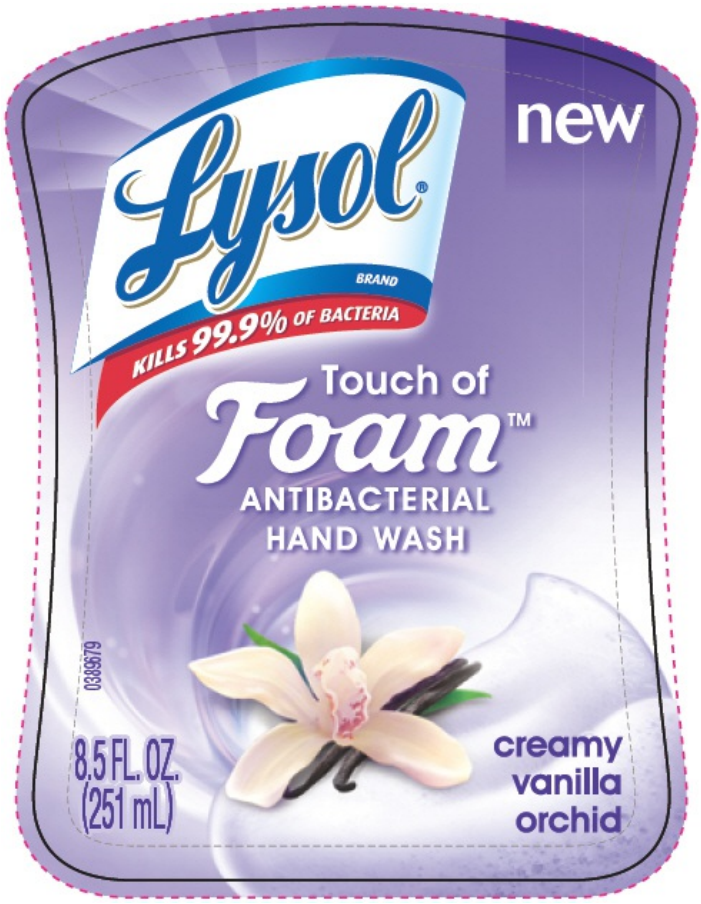
***Lysol*®**
BRAND
KILLS 99.9% OF BACTERIA

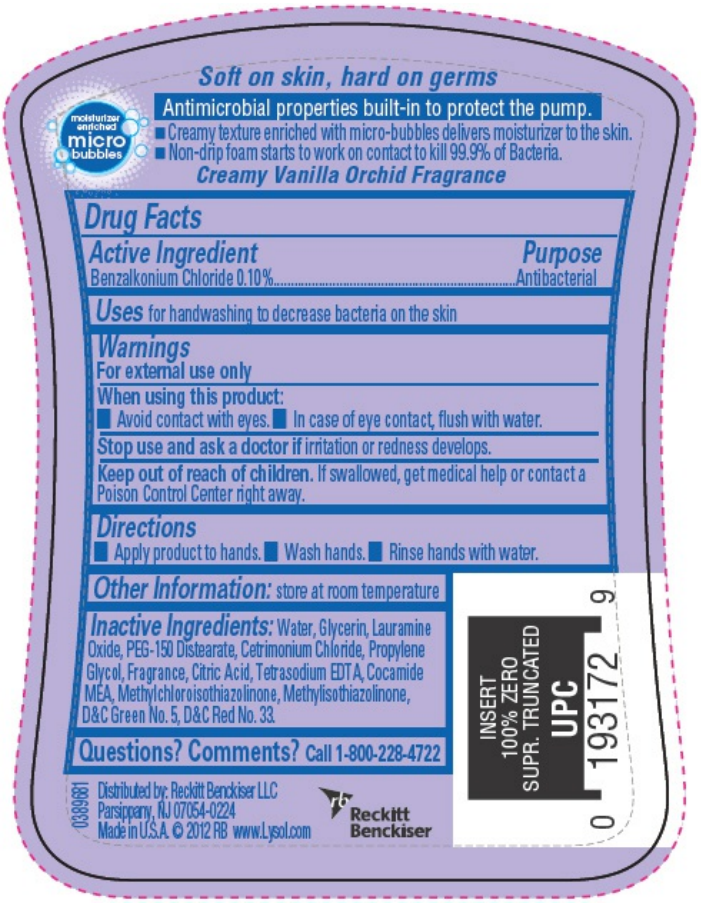
Touch of
***Foam*™**
ANTIBACTERIAL
HAND WASH

8.5 FL. OZ.
(251 mL)

creamy
vanilla
orchid

0389679





LYSOL TOUCH OF FOAM

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63824-466
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: 4F6FC4MI8W)	
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 Green No. 5 (UNII: 8J6RDU8L9X)	
D&C Red No. 33 (UNII: 9DBA0SBB0L)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63824-466-08	251 mL in 1 BOTTLE, PUMP		
2	NDC:63824-466-11	325 mL in 1 BOTTLE, PUMP		
3	NDC:63824-466-25	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)

Revised: 11/2012 Reckitt Benckiser LLC