

LYSOL NO-TOUCH- 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®
No-Touch™ Antibacterial Hand Soap

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.10%

Purpose

Antibacterial

Use

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

- Wet hands
- Place hands under dispenser until soap is dispensed
- Wash hands
- Rinse hands with water
- Dry hands after rinsing

Other Information

store at room temperature

Inactive Ingredients

Water, Cetrimonium Chloride, Glycerin, PEG-150 Distearate, Lauramine Oxide, Cocamide MEA, Citric Acid, Fragrance, Tetrasodium EDTA, Sodium Chloride, Methylchloroisoithiazolinone/ Methylisothiazolinone, PPG-12-Buteth-16, FD&C Blue #1

Questions? Comments?

Call 1-800-228-4722

Made in U.S.A.

PRINCIPAL DISPLAY PANEL - 251 mL Bottle Label

Limited Edition

**Sensitive
Skin**

**Dermatologically
Tested**

Lysol[®]

BRAND

KILLS 99.9% OF BACTERIA

No-Touch[™]

Refill

Antibacterial Hand Soap

Paraben Free • Triclosan Free

8.5 FL. OZ. (251 mL)

8105848

Limited Edition

Sensitive
Skin



No-Touch™
Refill



Antibacterial Hand Soap

Paraben Free • Triclosan Free

8105848

8.5 FL. OZ. (251 mL)

LYSOL® No-Touch®
Antibacterial Hand Soap Refill

- Kills 99.9% of harmful bacteria
- Is enriched with moisturizing ingredient
- Suitable for everyday use
- Dermatologically tested

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Reckitt Benckiser LLC
Parsippany, NJ 07054-0224
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www.Lysol.com



LYSOL NO-TOUCH

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63824-479
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)	
Cetrimonium Chloride (UNII: UC9PE95IBP)	
Glycerin (UNII: PDC6A3C0OX)	
PEG-150 Distearate (UNII: 6F36Q0I0AC)	
Lauramine Oxide (UNII: 4F6FC4M8W)	
Coco Monoethanolamide (UNII: C80684146D)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	

Edetate Sodium (UNII: MP1J8420LU)	
Sodium Chloride (UNII: 451W47IQ8X)	
Methylchloroisothiazolinone (UNII: DEL7T5QRPN)	
Methylisothiazolinone (UNII: 229D0E1QFA)	
PPG-12-Buteth-16 (UNII: 58CG7042J1)	
FD&C Blue No. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63824-479-02	251 mL in 1 BOTTLE		
2	NDC:63824-479-01	1 in 1 BLISTER PACK		
2		251 mL in 1 BOTTLE, PLASTIC		
3	NDC:63824-479-03	3 in 1 BLISTER PACK		
3		251 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	12/01/2013	

Labeler - Reckitt Benckiser LLC (094405024)

Revised: 12/2013

Reckitt Benckiser LLC