

**KROGER SPRINGWATER ANTIBACTERIAL- benzalkonium chloride liquid**  
**Kroger Company**

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**Kroger Moisturizing spring water antibacterial Hand Soap**

**Active Ingredient**

Benzalkonium Chloride 0.13%

**Purpose**

Antibacterial

**Uses**

Helps decrease bacteria on hands.

**Warnings**

For external use only.

**When using the product**

Avoid contact with eyes. In case of Contact, Flush with Water.

**Keep out of reach of children.**

In case of accidental ingestion, seek medical attention or contact a Poison Control Center immediately.

**Directions**

- From pump bottle, apply onto wet hands.
- Lather and rinse thoroughly.

**Other Information**

Store at room temperature.

**Inactive Ingredients**

Water (Aqua), Lauramidopropylamine Oxide, Glycerin, Cetrimonium Chloride, Sodium Chloride, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Fragrance (Parfum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Methylchloroisoethiazolinone, Methylisothiazolinone, Blue 1 (CI42090), Red 33 (CI 17200).

**PDP-1, NDC- 41226-510-11**



## Drug Facts

### Active ingredient

Benzalkonium Chloride

**Uses** helps decrease

### Warnings

For external use only

When using this product  
In case of contact, flush

Stop use and ask a doctor  
develops and lasts.

Keep out of reach of children  
accidental ingestion, or  
contact a Poison Control

**Directions** ■ From  
wet hands. ■ Lather a

### Other information

### Inactive ingredients

Lauramidopropylamine  
Cetrimonium Chloride  
MEA, PEG-120 Methyl  
(Parfum), Citric Acid, T  
Sulfate, Methylchloro  
Methylisothiazolinone  
(CI 17200).

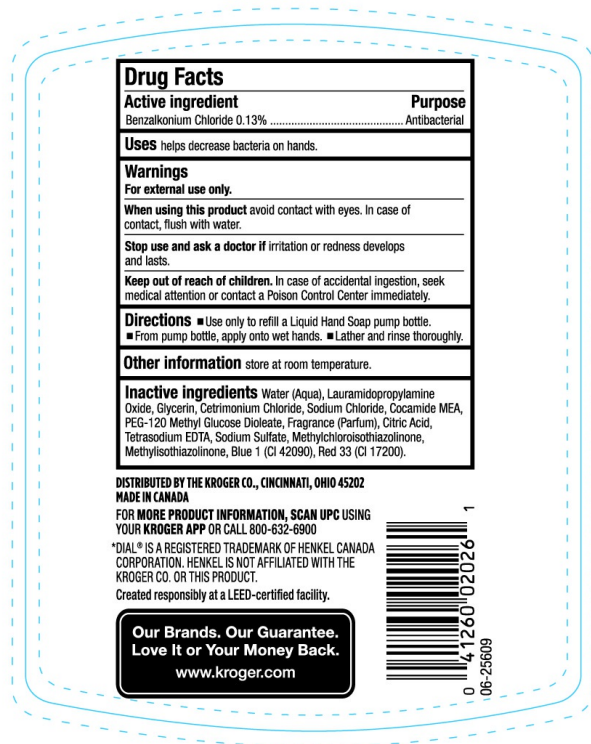
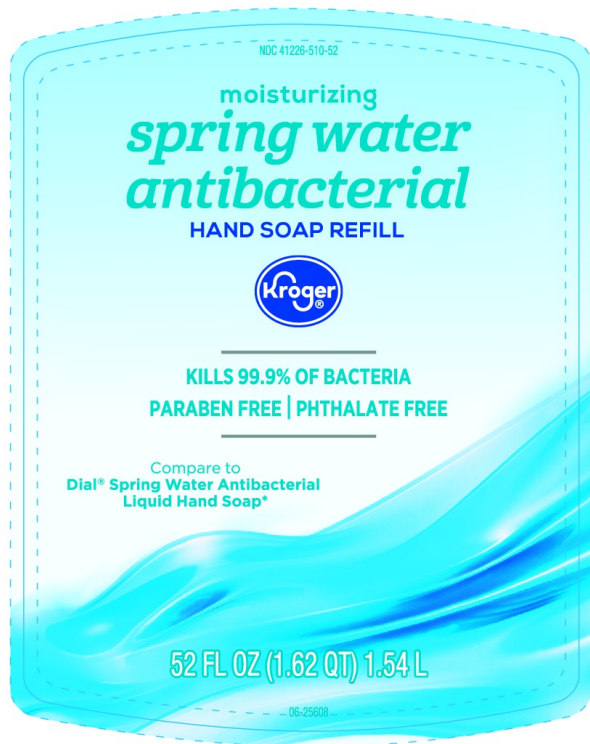
DISTRIBUTED BY THE KROGER COMPANY  
CINCINNATI, OHIO 45202  
MADE IN CANADA

FOR MORE PRODUCT INFORMATION  
SCAN UPC USING YOUR SMARTPHONE  
OR CALL 800-632-6900

DIAL® IS A REGISTERED TRADEMARK  
OF HENKEL CANADA COMPANY  
HENKEL IS NOT AFFILIATED WITH  
KROGER CO. OR THIS PRODUCT

Created responsibly at a LEED-certified facility.

**PDP-2, NDC- 41226-510-52**



## KROGER SPRINGWATER ANTIBACTERIAL

benzalkonium chloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:41226-510
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	130 mg in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>METHYLCHLOROISOTHIAZOLINONE</b> (UNII: DEL7T5QRPN)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>EDETATE SODIUM</b> (UNII: MP1J8420LU)	
<b>FRAGRANCE CLEAN ORC0600327</b> (UNII: 329LCV5BTF)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>CETRIMONIUM CHLORIDE</b> (UNII: UC9PE95IBP)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>LAURAMIDOPROPYLAMINE OXIDE</b> (UNII: I6KX160QTV)	
<b>COCO MONOETHANOLAMIDE</b> (UNII: C80684146D)	

<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>PEG-120 METHYL GLUCOSE DIOLEATE</b> (UNII: YM0K64F20V)	
<b>SODIUM SULFATE</b> (UNII: 0YPR65R21J)	
<b>METHYLISOTHIAZOLINONE</b> (UNII: 229D0E1QFA)	
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41226-510-11	325 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/05/2024	
2	NDC:41226-510-52	1540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/05/2024	

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/05/2024	

**Labeler** - Kroger Company (006999528)

**Registrant** - Apollo Health and Beauty Care (201901209)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care		201901209	manufacture(41226-510)

Revised: 2/2024

Kroger Company