

**DG BODY REFRESHING CLEAN- benzalkonium chloride liquid**  
**DOLGENCORP INC**

*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.*

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**DRUG FACTS**

**ACTIVE INGREDIENT**

BENZALKONIUM CHLORIDE 0.13%

**PURPOSE**

ANTIBACTERIAL

**USES**

HELPS ELIMINATE BACTERIA ON HANDS

**WARNINGS**

FOR EXTERNAL USE ONLY

**WHEN USING THIS PRODUCT**

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE EYES THOROUGHLY WITH WATER

STOP USE AND ASK A DOCTOR IF

IRRITATION OR REDNESS DEVELOPS AND LASTS

*KEEP OUT OF REACH OF CHILDREN*

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY

**DIRECTIONS**

APPLY ONTO WET HANDS. LATHER AND RINSE THOROUGHLY

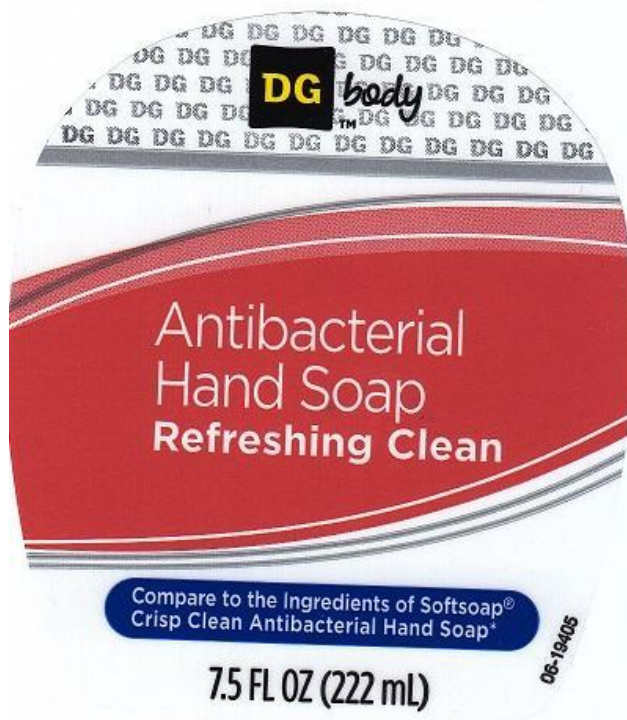
**OTHER INFORMATION**

STORE AT ROOM TEMPERATURE

**INACTIVE INGREDIENTS**

WATER (AQUA), CETRIMONIUM CHLORIDE, GLYCERIN, LAURYL/MYRISTYL AMIDOPROPYL AMINE OXIDE, COCAMIDE MEA, SODIUM CHLORIDE, PEG-120 METHYL GLUCOSE DIOLATE, FRAGRANCE (PARFUM), CITRIC ACID, TETRASODIUM EDTA, SODIUM SULFATE, METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE, RED 40 (CI 16035), YELLOW 5 (CI 19140), RED 33 (CI 17200)

**LABEL COPY**



**Antibacterial Hand Soap Refreshing Clean**

<b>Drug Facts</b>		<b>Drug Facts (continued)</b>	
<b>Active ingredient</b> Benzalkonium Chloride 0.13%	<b>Purpose</b> Antibacterial	<b>Directions</b> ■ Apply onto wet hands. Lather and rinse thoroughly.	
<b>Uses</b> ■ Helps eliminate bacteria on hands.		<b>Other information</b> ■ Store at room temperature.	
<b>Warnings</b> <b>For external use only</b> When using this product ■ avoid contact with eyes. ■ If contact occurs, rinse eyes thoroughly with water.		<b>Inactive Ingredients:</b> Water (Aqua), Cetrimonium Chloride, Glycerin, Lauryl/Myristyl Amidopropyl Amine Oxide, Cocamide MEA, Sodium Chloride, PEG-120 Methyl Glucose Dioleate, Fragrance (Parfum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Methylchloroisothiazolinone, Methylisothiazolinone, Red 40 (CI 16035), Yellow 5 (CI 19140), Red 33 (CI 17200).	
<b>Stop use and ask a doctor if</b> ■ irritation or redness develops and lasts.		*This product is not manufactured or distributed by the Colgate-Palmolive Company, owner of the registered trademark Softsoap®.	
<b>Keep out of reach of children.</b> ■ In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.			

B0310

100% QUALITY GUARANTEED (888) 309-9030

0 67153 94641 8

06-19406

DISTRIBUTED BY DOLGENCORP, LLC  
100 MISSION RIDGE, GOODLETTSVILLE, TN 37072  
MADE IN CANADA

**DG BODY REFRESHING CLEAN**

benzalkonium chloride liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:559 10-721
<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	1.3 mg in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CETRIMONIUM CHLORIDE</b> (UNII: UC9PE95IBP)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>LAURAMIDOPROPYLAMINE OXIDE</b> (UNII: I6KX160QTV)	
<b>COCO MONOETHANOLAMIDE</b> (UNII: C80684146D)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>PEG-120 METHYL GLUCOSE DIOLEATE</b> (UNII: YM0K64F20V)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	

EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-721-08	222 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	02/25/2014	

**Labeler** - DOLGENCORP INC (068331990)

**Registrant** - APOLLO HEALTH AND BEAUTY CARE (201901209)

### Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(55910-721)

Revised: 2/2014

DOLGENCORP INC