

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.

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

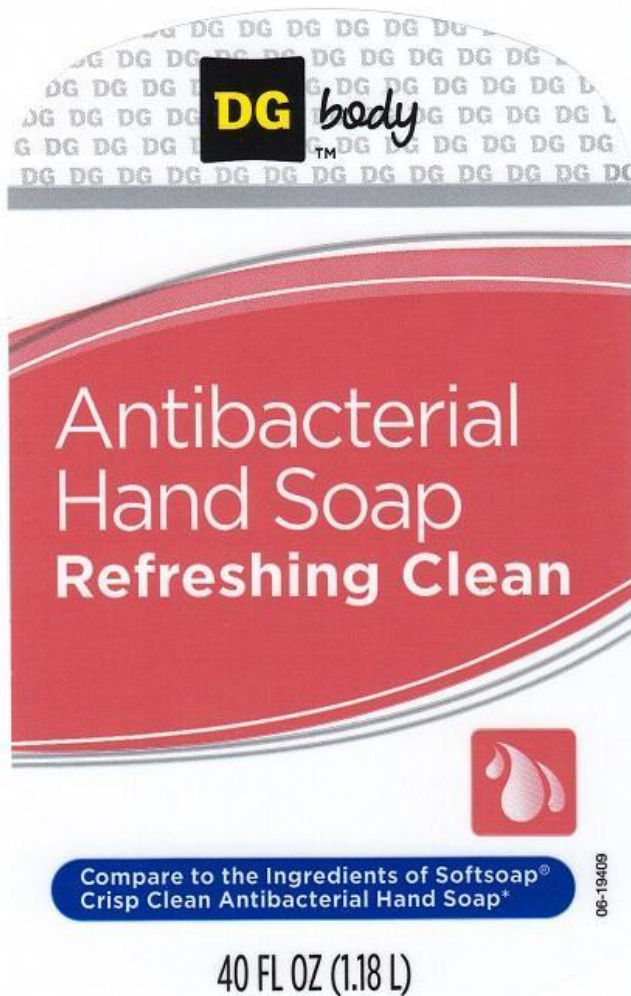
OTHER INFORMATION

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

DG™ Body Antibacterial Hand Soap Refreshing Clean is an antibacterial formula that eliminates the dirt you see and the germs you don't. DG™ Body Antibacterial Hand Soap Refreshing Clean actually kills germs and bacteria so it's perfect for use at both the bathroom and kitchen sinks. DG™ Body Antibacterial Hand Soap Refreshing Clean is effective, yet gentle and mild, so it's great for the entire family.

Drug Facts	
Active ingredient	Purpose
Benzalkonium Chloride 0.13%	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 Dioleate, Fragrance (Parfum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Methylchloroisothiazolinone, Methylisothiazolinone, Red 40 (CI 16035), Yellow 5 (CI 19140), Red 33 (CI 17200).	

*This product is not manufactured or distributed by the Colgate-Palmolive Company, owner of the registered trademark Softsoap®.

DISTRIBUTED BY DOLGENCORP, LLC
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GOODLETTSVILLE, TN 37072
MADE IN CANADA

B0204

100% Quality Guaranteed
(888) 309-9030



06-19410

DG BODY REFRESHING CLEAN

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-721
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
GLYCERIN (UNII: PDC6A3C0OX)	

LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PEG-120 METHYL GLUCOSE DIOLATE (UNII: YM0K64F20V)	
CITRIC ACID MONOHYDRATE (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:55910-721-40	1180 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)