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

- USE AS A REFILL FOR A PUMP DISPENSING BOTTLE. FROM PUMP BOTTLE, 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)



Antibacterial Hand Soap Refill 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%.

intibacterial

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 Use as a refill for a pump dispensing bottle. From pump bottle, apply onto wet hands. Lather and rinse thoroughly.

Other information store at room temperature.

Inactive ingredients water (Aqua), Cetrimonium Chloride, Glycerin, Launyl/Myristy Amidopropyl Amine Oxide, Cocamide MEA, Sodium Chloride, PEG-120 Methyl Glucose Dicleate Fragrance (Parlum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Methylchloroisothiazolinon Methylisothiazolinone, Red 40 (Cl 16035), Yellow 5 (Cl 19140), Red 33 (Cl 17200).

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

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

100% — Satisfaction Guaranteed! (888) 309-9030



36-20547

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
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CETRIMO NIUM CHLO RIDE (UNII: UC9 PE9 5 IBP)	
GLYCERIN (UNII: PDC6A3C0OX)	

LAURAMIDO PRO PYLAMINE O XIDE (UNII: 16 KX 16 0 Q TV)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)	
EDETATE SO DIUM (UNII: MP1J8420 LU)	
SODIUM SULFATE (UNII: 0 YPR6 5R21J)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229 D0 E1QFA)	
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-	1180 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/17/2015	

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 (559 10 - 721)	

Revised: 6/2015 DOLGENCORP INC