DIAL COMPLETE ANTIBACTERIAL LIQUID HAND GOLD- benzalkonium chloride solution
MID-CONTINENT PACKAGING, 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.

Dial Complete® Antibacterial Liquid Hand Soap
Gold

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

- 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

- Pump into hands, wet as needed
- Lather vigorously for at least 15 seconds
- Wash skin, rinse thoroughly and dry

Inactive ingredients

Aqua (Water, Eau) • Lauramidopropylamine Oxide • Glycerin • Cetrimonium Chloride • Lauramine Oxide • Sodium Chloride • PEG-120 Methyl Glucose Dioleate • Citric Acid • Sodium Benzoate • Myristamidopropylamine Oxide • Zinc Sulfate • Parfum (Fragrance) •

Dimethyl Lauramine • Tetrasodium EDTA • Alcohol • Dimethyl Myristamine • CI 19140 (Yellow 5) • CI 14700 (Red 4)

Questions?

1-800-258-DIAL (3425)

Distributed by
Henkel Corporation,
Rocky Hill, CT 06067

PRINCIPAL DISPLAY PANEL - 1.53 L Bottle Label

KILLS 99.9% OF BACTERIA*

NEW
LOOK!

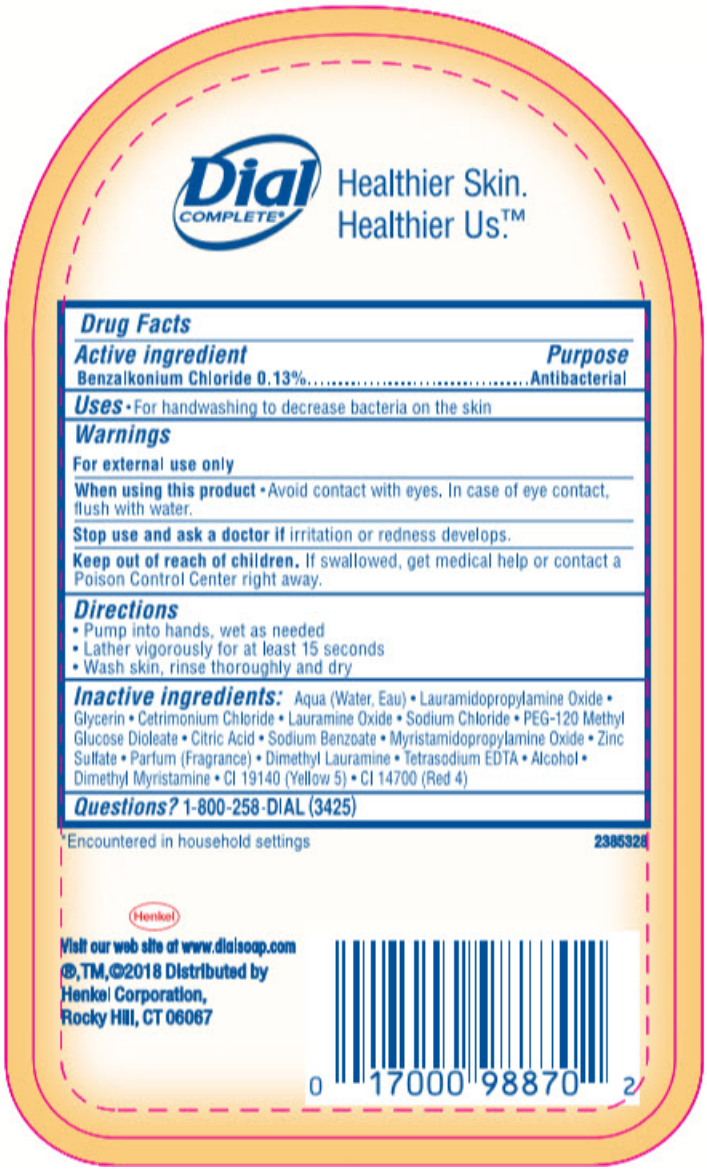
Dial
COMPLETE®

LIQUID
antibacterial
Hand Soap

GOLD

52 FL OZ (1.62 QT) 1.53 L

2370135



DIAL COMPLETE ANTIBACTERIAL LIQUID HAND GOLD

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69560-345
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)	
Lauramidopropylamine Oxide (UNII: I6KX160QTV)	
Lauramine Oxide (UNII: 4F6FC4MI8W)	
Cetrimonium Chloride (UNII: UC9PE95IBP)	
GLYCERIN (UNII: PDC6A3C0OX)	
Sodium Chloride (UNII: 451W47IQ8X)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
Sodium Benzoate (UNII: OJ245FE5EU)	
Edetate Sodium (UNII: MP1J8420LU)	
Myristamidopropylamine Oxide (UNII: 3HSF539C9T)	
Zinc Sulfate Heptahydrate (UNII: N57JI2K7WP)	
Dimethyl Lauramine (UNII: 6V2OM30I1Z)	
Alcohol (UNII: 3K9958V90M)	
Dimethyl Myristamine (UNII: 5E4O85D8T2)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	

Product Characteristics

Color	ORANGE	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69560-345-02	1530 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	05/05/2021	

Labeler - MID-CONTINENT PACKAGING, INC. (798250239)

Revised: 4/2021 MID-CONTINENT PACKAGING, INC.