DIAL COMPLETE ANTIBACTERIAL FOAMING HAND WASH SPRING WATER- 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 Foaming Hand Wash
Spring Water®

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 DRY hands
- Lather vigorously for at least 15 seconds
- Rinse and dry thoroughly

Inactive Ingredients

Aqua (Water, Eau) • Glycerin • Lauramine Oxide • Cetrimonium Chloride • Cocamidopropyl Betaine • Citric Acid • Sodium Benzoate • Hydroxypropyl Methylcellulose • Parfum (Fragrance) • Zinc Sulfate • Sodium Chloride • Dimethyl

Lauramine • Tetrasodium EDTA • Alcohol • Dimethyl Myristamine • CI 42090 (Blue 1) • CI17200 (Red 33)

Distributed by Henkel
Corporation, Rocky Hill, CT 06067

PRINCIPAL DISPLAY PANEL - 1.53 L Bottle Label

KILLS 99.99% OF BACTERIA*

#1 DR. RECOMMENDED#

Dial
COMPLETE®

FOAMING
antibacterial
HAND WASH

spring
water®

REFILL
USE ONLY
IN DIAL
COMPLETE®
FOAM PUMP

SKIN
SMART

moisturizing conditioners
gentle cleansers
free from parabens,
phthalates, silicones

52 FL OZ (1.62 QT) 1.53 L REFILL

2632116

DIAL COMPLETE ANTIBACTERIAL FOAMING HAND WASH SPRING WATER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69560-456
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)	
Lauramine Oxide (UNII: 4F6FC4MI8W)	
Cetrimonium Chloride (UNII: UC9PE95IBP)	
Glycerin (UNII: PDC6A3C0OX)	
Cocamidopropyl Betaine (UNII: 5OCF3O11KX)	
Anhydrous Citric Acid (UNII: XF417D3PSL)	
Hypromellose, Unspecified (UNII: 3NXW29V3WO)	
Sodium Benzoate (UNII: OJ245FE5EU)	
Zinc Sulfate Heptahydrate (UNII: N57JI2K7WP)	
Edetate Sodium (UNII: MP1J8420LU)	
Dimethyl Lauramine (UNII: 6V2OM30I1Z)	
Alcohol (UNII: 3K9958V90M)	
Dimethyl Myristamine (UNII: 5E4O85D8T2)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69560-456-02	1530 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2021	
2	NDC:69560-456-03	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	05/01/2021	

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

Revised: 12/2021 MID-CONTINENT PACKAGING, INC.