

DIAL PROFESSIONAL ANTIBACTERIAL DEFENSE SPRING WATER- benzalkonium chloride solution
DIAL PROFESSIONAL ANTIBACTERIAL AND SENSITIVE FRAGRANCE FREE- benzalkonium chloride solution
DIAL PROFESSIONAL ANTIBACTERIAL DEFENSE GOLD- benzalkonium chloride solution
Henkel Corporation

Dial Professional Gold Antibacterial Liquid Hand Soap
Dial Professional Antibacterial Gold Liquid Hand Soap Refill for Versa Dispenser
Dial Professional Liquid Antibacterial Defense Hand Soap Spring Water Scent
Dial Professional Liquid Antibacterial Defense Hand Soap Gold
Dial Professional Liquid Antibacterial & Sensitive Hand Soap Fragrance Free

Drug Facts



FREE FROM:
SLS/SLES**
Parabens
Phthalates
Silicones



Cruelty Free
INTERNATIONAL



pH Balanced



Dermatologist
Tested

**Sodium lauryl sulfate/Sodium laureth sulfate

Drug Facts

Active ingredient Purpose

Benzalkonium Chloride 0.13%.....Antibacterial

Use 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. ▶

Drug Facts (continued)

Directions

- Pump into hands, wet as needed
- Lather vigorously for at least 30 seconds
- Wash skin, rinse thoroughly and dry

Inactive ingredients

Aqua (Water, Eau) • Sodium Chloride • Lauramidopropylamine Oxide • Glycerin • Cetrimonium Chloride • Lauramine Oxide • PEG-120 Methyl Glucose Dioleate • Citric Acid • Sodium Benzoate • Myristamidopropylamine Oxide • Parfum (Fragrance) • Aloe Barbadensis Leaf Juice • Zinc Sulfate • Trisodium Ethylenediamine Disuccinate • Dimethyl Lauramine • Alcohol • CI 19140 (Yellow 5) • CI 14700 (Red 4)

*Encountered in away-from-home settings
**Antibacterial Hand Soap Brand

Questions? 1-877-777-3277

Mixed and bottled in the USA.

Discard Seal
Empty &
Replace Cap



PLASTIC
BOTTLE



Scan here to learn more about **Dial Professional**.



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ROCKY HILL, CT 06067
www.dialprofessional.com

CM-5-88047-09/2989834

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

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Directions

- Pump into hands, wet as needed.
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Inactive Ingredients

Variant Gold:

Inactive ingredients: Aqua (Water, Eau) · Sodium Chloride · Lauramidopropylamine Oxide · Glycerin · Cetrimonium Chloride · Lauramine Oxide · PEG-120 Methyl Glucose Dioleate · Citric Acid · Sodium Benzoate · Myristamidopropylamine Oxide · Parfum (Fragrance) · Aloe Barbadensis Leaf Juice · Zinc Sulfate · Trisodium Ethylenediamine Disuccinate · Dimethyl Lauramine. Alcohol. CI 19140 (yellow 5). CI 14700 (Red 4)

Variant AB Fragrance Free:

Inactive ingredients: Aqua (Water, Eau) · Sodium Chloride · Lauramidopropylamine Oxide · Glycerin · Cetrimonium Chloride · Lauramine Oxide · PEG-120 Methyl Glucose Dioleate · Citric Acid · Sodium Benzoate · Myristamidopropylamine Oxide · Aloe Barbadensis Leaf Juice · Zinc Sulfate · Trisodium Ethylenediamine Disuccinate · Dimethyl Lauramine · Alcohol

Variant Spring Water:

Inactive ingredients: Aqua (Water, Eau) · Sodium Chloride · Lauramidopropylamine Oxide · Glycerin · Cetrimonium Chloride · Lauramine Oxide · PEG-120 Methyl Glucose Dioleate · Citric Acid · Sodium Benzoate · Parfum (Fragrance) · Myristamidopropylamine Oxide · Aloe Barbadensis Leaf Juice · Zinc Sulfate · Trisodium Ethylenediamine Disuccinate · Dimethyl Lauramine · Alcohol · CI 42090 (Blue 1) · CI 17200 (Red 33)

Questions

Questions? 1-877-777-3277

Legal Entity

henkel Bean Logo

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Henkel Corporation, Rocky Hill, CT 06067

www.dialsprofessional.com

^Encountered in household settings

#Antibacterial Hand Soap

Indications & Usage

For handwashing to decrease bacteria on the skin.

Topical Liquid

Liquid Hand Soap

Principal Display Panel

LIQUID

KILLS MORE THAN 99.9% of BACTERIA*
#1 DR. RECOMMENDED BRAND**



LIQUID
antibacterial
defense™

HAND SOAP

+ aloe

spring
water®
scent

1 Gallon (3.78 L)



FREE FROM:
SLS/SLES**
Parabens
Phthalates
Silicones



Cruelty Free
INTERNATIONAL



pH Balanced



Dermatologist
Tested

**Sodium lauryl sulfate/Sodium laureth sulfate

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**Antibacterial Hand Soap Brand

Questions? 1-877-777-3277

Mixed and bottled in the USA.



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PLASTIC BOTTLE



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2989835

KILLS MORE THAN 99.9% of BACTERIA*
#1 DR. RECOMMENDED BRAND**



LIQUID

**antibacterial
defense™**

HAND SOAP

+ aloe 

gold

1 Gallon (3.78 L)

2989834



DIAL PROFESSIONAL ANTIBACTERIAL DEFENSE SPRING WATER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54340-121
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ZINC SULFATE (UNII: 89DS0H96TB)	0.05 g in 100 mL
ALCOHOL (UNII: 3K9958V90M)	0.013 g in 100 mL
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	1.635 g in 100 mL
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	0.19 g in 100 mL
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	0.0185 g in 100 mL
DIMETHYL LAURAMINE (UNII: 6V2OM30I1Z)	0.0162 g in 100 mL
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	0.975 g in 100 mL
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	0.575 g in 100 mL
WATER (UNII: 059QF0KO0R)	89.65 mL in 100 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.0997 g in 100 mL
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	1.5 g in 100 mL
SODIUM BENZOATE (UNII: OJ245FE5EU)	0.4 g in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	1.5 g in 100 mL
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	0.75 g in 100 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X)	2.2 g in 100 mL
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	0.000051 g in 100 mL
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	0.000017 g in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54340-121-02	4 in 1 CARTON	01/01/2024	
1	NDC:54340-121-01	3785 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/01/2024	

DIAL PROFESSIONAL ANTIBACTERIAL AND SENSITIVE FRAGRANCE FREE

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54340-309
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ZINC SULFATE (UNII: 89DS0H96TB)	0.05 g in 100 mL
ALCOHOL (UNII: 3K9958V90M)	0.013 g in 100 mL
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	1.635 g in 100 mL
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	0.19 g in 100 mL
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	0.0185 g in 100 mL
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	0.975 g in 100 mL
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	0.575 g in 100 mL
WATER (UNII: 059QF0KO0R)	89.727 mL in 100 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.0997 g in 100 mL
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	1.5 g in 100 mL
SODIUM BENZOATE (UNII: OJ245FE5EU)	0.4 g in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	1.5 g in 100 mL
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	0.75 g in 100 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X)	2.4 g in 100 mL
DIMETHYL LAURAMINE (UNII: 6V2OM30I1Z)	0.0162 g in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54340-309-04	4 in 1 CARTON	01/01/2024	
1	NDC:54340-309-03	3785 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		
2	NDC:54340-309-02	8 in 1 CARTON	01/01/2024	
2	NDC:54340-309-01	1000 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		
3	NDC:54340-309-06	12 in 1 CARTON	01/01/2024	
3	NDC:54340-309-05	325 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/01/2024	

DIAL PROFESSIONAL ANTIBACTERIAL DEFENSE GOLD

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54340-302
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ZINC SULFATE (UNII: 89DS0H96TB)	0.05 g in 100 mL
ALCOHOL (UNII: 3K9958V90M)	0.013 g in 100 mL
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	1.635 g in 100 mL
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	0.19 g in 100 mL
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	0.0185 g in 100 mL
DIMETHYL LAURAMINE (UNII: 6V2OM30I1Z)	0.0162 g in 100 mL
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	0.975 g in 100 mL
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	0.575 g in 100 mL
WATER (UNII: 059QF0KO0R)	89.8 mL in 100 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.0997 g in 100 mL
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	1.5 g in 100 mL
SODIUM BENZOATE (UNII: OJ245FE5EU)	0.4 g in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	1.5 g in 100 mL
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	0.75 g in 100 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X)	2.2 g in 100 mL
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	0.0056 g in 100 mL
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	0.00029 g in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54340-302-02	12 in 1 CARTON	01/01/2024	

1	NDC:54340-302-01	325 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		
2	NDC:54340-302-06	6 in 1 CARTON	01/01/2024	
2	NDC:54340-302-05	443 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		
3	NDC:54340-302-04	4 in 1 CARTON	01/01/2024	
3	NDC:54340-302-09	3785 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		
4	NDC:54340-302-03	8 in 1 CARTON	01/01/2024	
4		1000 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		
5	NDC:54340-302-08	144 in 1 CARTON	01/01/2024	
5	NDC:54340-302-07	59 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/01/2024	

Labeler - Henkel Corporation (080887708)

Revised: 12/2025

Henkel Corporation