

DIAL COMPLETE ALOE- antibacterial hand wash aloe solution
DIAL COMPLETE GOLD- antibacterial hand wash gold solution
DIAL COMPLETE SPRING WATER- antibacterial hand wash spring water solution
DIAL COMPLETE WHITE TEA- antibacterial hand wash white tea solution
Henkel Corporation

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

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

For Handwashing to decreases 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 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 in to hands, wet as needed.
- Later vigorously for at least 15 seconds.
- Wash skin, rinse thoroughly and dry.

Inactive Ingredients

Gold

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)

Aloe

Aqua (Water, Eau) · Lauramidopropylamine Oxide · Glycerin · Cetrimonium Chloride · Lauramine Oxide ·
Sodium Chloride ·
PEG-120 Methyl Glucose Dioleate · Citric Acid · Sodium Benzoate · Myristamidopropylamine Oxide ·
Parfum (Fragrance) ·
· Zinc Sulfate · Aloe Barbadensis Leaf Juice · Dimethyl Lauramine · Tetrasodium EDTA · Alcohol ·
Dimethyl Myristamine
· CI 19140 (Yellow 5) · CI 42090 (Blue 1)

Spring Water

Aqua (Water, Eau) · Lauramidopropylamine Oxide · Glycerin · Cetrimonium Chloride · Lauramine Oxide ·
Sodium Chloride ·
PEG-120 Methyl Glucose Dioleate · Citric Acid · Sodium Benzoate · Myristamidopropylamine Oxide ·
Parfum (Fragrance) ·
Zinc Sulfate · Dimethyl Lauramine · Tetrasodium EDTA · Alcohol · Dimethyl Myristamine · CI 42090
(Blue 1) · CI 17200 (Red 33)

White Tea

Aqua (Water, Eau) · Lauramidopropylamine Oxide · Glycerin · Lauramine Oxide · Cetrimonium
Chloride ·
Sodium Chloride · PEG-120 Methyl Glucose Dioleate · Citric Acid · Sodium Benzoate ·
Myristamidopropylamine Oxide ·
Parfum (Fragrance) · Zinc Sulfate · Dimethyl Lauramine · Tetrasodium EDTA · Alcohol · Camellia
Sinensis Leaf Extract ·
Dimethyl Myristamine · CI 42090 (Blue 1) · CI 17200 (Red 33)

Questions

call 1-800-258-DIAL(3425)

Legal Entity

www.dialsoap.com

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^Encountered in household settings

#Antibacterial Hand Soap

For handwashing to decrease bacteria on the skin.

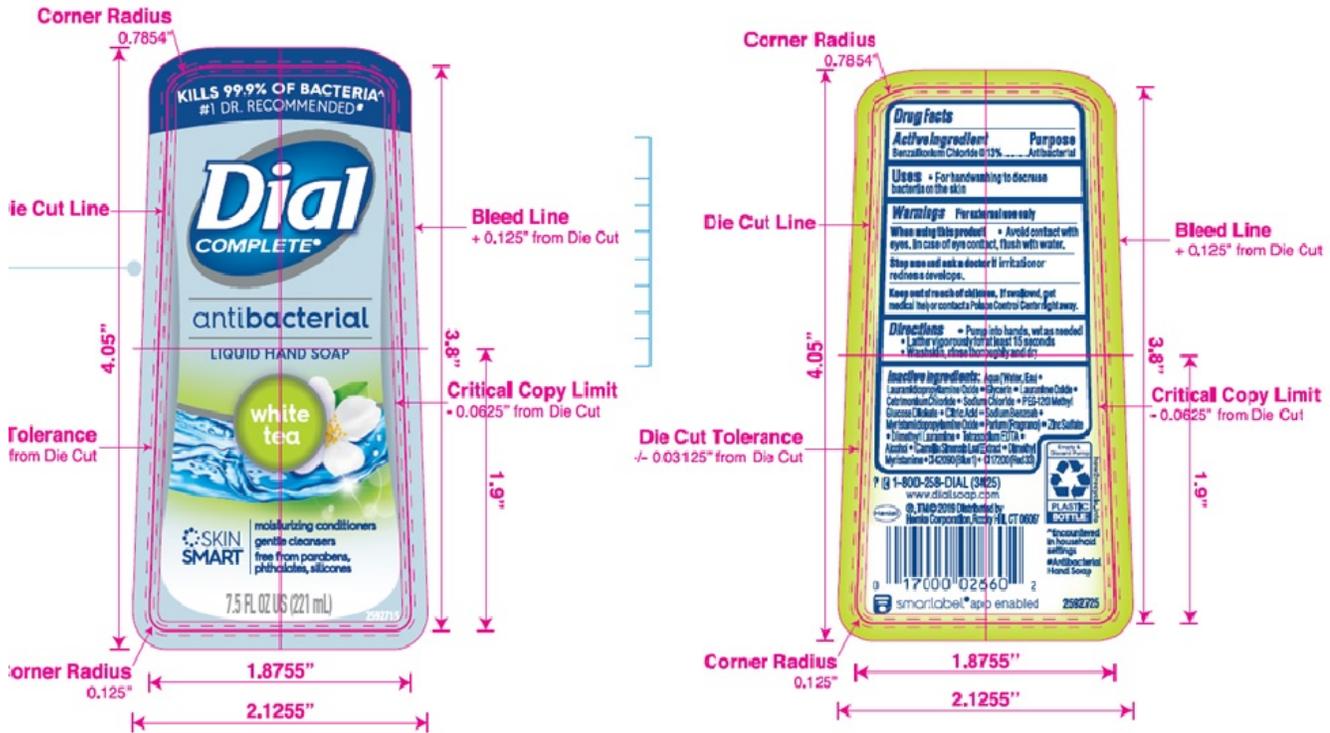
Topical Liquid

Liquid Hand Soap

Front and the back of the pack







DIAL COMPLETE ALOE

antibacterial hand wash aloe solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54340-146
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
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
EDETATE SODIUM (UNII: MPLJ8420LU)	0.02 mL in 100 mL
WATER (UNII: 059QF0K00R)	90.27 mL in 100 mL
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
DIMETHYL LAURAMINE (UNII: 6V2OM301IZ)	
ZINC SULFATE (UNII: 89DS0H96TB)	0.05 mL in 100 mL
MYRISTAMIDO PROPYLAMINE OXIDE (UNII: 3HSF539C9T)	

LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHYL MYRISTAMINE (UNII: 5E4085D8T2)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	0.4 mL in 100 mL
ALCOHOL (UNII: 3K9958V90M)	0.013 mL in 100 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54340-146-01	221 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/01/2020	
2	NDC:54340-146-02	277 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/01/2020	
3	NDC:54340-146-03	1180 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

DIAL COMPLETE GOLD

antibacterial hand wash gold solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54340-143
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
EDETATE SODIUM (UNII: MP1J8420LU)	0.02 mL in 100 mL
WATER (UNII: 059QF0KO0R)	90.43 mL in 100 mL
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	0.005624 mL in 100 mL
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
PEG-120 METHYL GLUCOSE DIOLATE (UNII: YM0K64F20V)	
DIMETHYL LAURAMINE (UNII: 6V2OM30IIZ)	
ZINC SULFATE (UNII: 89DS0H96TB)	0.05 mL in 100 mL

MYRISTAMIDO PROPYLAMINE O XIDE (UNII: 3HSF539C9T)	
LAURAMIDO PROPYLAMINE O XIDE (UNII: I6KX160QTV)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHYL MYRISTAMINE (UNII: 5E4O85D8T2)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	0.4 mL in 100 mL
ALCOHOL (UNII: 3K9958V90M)	0.013 mL in 100 mL
FD&C RED NO. 4 FREE ACID (UNII: WJE3T5596E)	0.000288 mL in 100 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X)	0.9503 mL in 100 mL
LAURAMINE O XIDE (UNII: 4F6FC4M18W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54340-143-01	221 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/01/2020	
2	NDC:54340-143-02	1530 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

DIAL COMPLETE SPRING WATER

antibacterial hand wash spring water solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54340-142
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
EDETATE SODIUM (UNII: MP1J8420LU)	
WATER (UNII: 059QF0KO0R)	90.34 mL in 100 mL
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
LAURAMINE O XIDE (UNII: 4F6FC4M18W)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	

DIMETHYL LAURAMINE (UNII: 6V2OM3011Z)	
ZINC SULFATE (UNII: 89DS0H96TB)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHYL MYRISTAMINE (UNII: 5E4O85D8T2)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
ALCOHOL (UNII: 3K9958V90M)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54340-142-02	221 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/01/2020	
2	NDC:54340-142-04	946 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/01/2020	
3	NDC:54340-142-05	1530 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

DIAL COMPLETE WHITE TEA

antibacterial hand wash white tea solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54340-145
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
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
LAURAMINE OXIDE (UNII: 4F6FC4M18W)	

EDETATE SODIUM (UNII: MP1J8420LU)	
WATER (UNII: 059QF0K00R)	90.46 mL in 100 mL
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
DIMETHYL LAURAMINE (UNII: 6V2OM301IZ)	
ZINC SULFATE (UNII: 89DS0H96TB)	
MYRISTAMIDO PROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHYL MYRISTAMINE (UNII: 5E4O85D8T2)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
ALCOHOL (UNII: 3K9958V90M)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54340-145-01	221 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

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