

**DIAL ANTIBACTERIAL BAR- dial gold antibacterial bar soap soap
Henkel Corporation**

**Dial Antibacterial Bar Soap - Gold
Dial Gold Antibacterial Deodorant Bar
Dial Advanced Clean Deodorant Bar Soap Antibacterial Gold**

Active ingredient

Benzalkonium Chloride 0.10%

Purpose

Antibacterial

#1 ANTIBACTERIAL BAR SOAP BRAND*
ELIMINATES 99.9% OF BACTERIA**

Dial
3
LARGE BARS

advanced clean™
DEODORANT BAR SOAP
ANTIBACTERIAL
gold

ROUND THE CLOCK®
ODOR PROTECTION

CLEANRINSE
TECHNOLOGY™

Dial advanced clean™
DEODORANT BAR SOAP
ANTIBACTERIAL
gold

Enjoy the crisp, iconic fragrance of Dial® Gold with the confidence of round the clock odor protection.
Dial® Clean Rinse Technology cleanses deep while being gentle on your skin, leaving it feeling soft & healthy.

FREE FROM:
SLS/SLES**
Parabens
Phthalates
Silicones

Cruelty Free
INTERNATIONAL

Dermatologist
Tested

*Based on IRI MULO unit sales data for the 52 week period ending 12/31/23
**Sodium Lauryl Sulfate / Sodium Laureth Sulfate
***Encountered in household settings Scan here to learn more about Dial.

Made in Colombia

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Uses

- for washing to decrease the bacterial 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 using 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

- wet bar with water
- lather vigorously and wash skin
- rinse and dry thoroughly

Inactive ingredients:

Soap [Sodium Palmate* · Sodium Cocoate* · Sodium Palm Kernelate*] · Water · Glycerin · Coconut Acid* · Palm Acid* · Palm Kernel Acid* · Stearic Acid* · Fragrance · Sorbitol · Sodium Chloride · Tetrasodium Glutamate Diacetate · Titanium Dioxide · Yellow 5 · Alcohol · Yellow 8 · Red 4

*Contains one or more of these ingredients



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3-4 OZ SOAP BARS (113 g), TOTAL NET WT 12 OZ (339 g)

OPEN HERE

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Drug Facts

Active Ingredient	Purpose
Benzalkonium Chloride 0.10%	Antibacterial

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Questions? 1-800-258-DIAL (3425)

Visit our website at www.dialsoap.com



2904695



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54340-272
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
EXT. D&C YELLOW NO. 7 (UNII: 08F8S9O3I5)	
GLYCERIN (UNII: PDC6A3C0OX)	8.1 g in 100 g
WATER (UNII: 059QF0KO0R)	14.086 g in 100 g

SODIUM CHLORIDE (UNII: 451W47IQ8X)	0.393 g in 100 g
SODIUM PALM KERNELATE (UNII: 6H91L1NXTW)	11.07 g in 100 g
SORBITOL (UNII: 506T60A25R)	0.652 g in 100 g
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	0.0182 g in 100 g
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	0.0588 g in 100 g
STEARIC ACID (UNII: 4ELV7Z65AP)	2.45 g in 100 g
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	0.035 g in 100 g
ALCOHOL (UNII: 3K9958V90M)	0.01 g in 100 g
CI 14700 (UNII: X3W0AM1JLX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54340-272-22	181 g in 1 CELLO PACK; Type 0: Not a Combination Product	01/01/2023	01/01/2024
2	NDC:54340-272-02	8 in 1 CARTON	02/01/2023	
2	NDC:54340-272-39	452 g in 1 CELLO PACK; Type 0: Not a Combination Product		
3	NDC:54340-272-20	12 in 1 CARTON	02/01/2023	
3	NDC:54340-272-38	339 g in 1 CELLO PACK; Type 0: Not a Combination Product		
4	NDC:54340-272-05	4 in 1 CARTON	02/01/2023	
4	NDC:54340-272-36	904 g in 1 CELLO PACK; Type 0: Not a Combination Product		
5	NDC:54340-272-41	1356 g in 1 CASE; Type 0: Not a Combination Product	02/01/2023	
6	NDC:54340-272-28	63 g in 1 CARTON; Type 0: Not a Combination Product	03/01/2023	01/01/2024
7	NDC:54340-272-21	678 g in 1 CELLO PACK; Type 0: Not a Combination Product	01/31/2023	01/01/2024
8	NDC:54340-272-01	36 in 1 CARTON	02/01/2023	
8	NDC:54340-272-37	113 g in 1 CELLO PACK; 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)	02/01/2010	

Labeler - Henkel Corporation (080887708)