DOVE- men skin defense antibacterial 3n1 hand body shave bar soap
Conopco Inc. d/b/a/ Unilever

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.

Dove Men +Care Skin Defense Antibacterial 3-N-1 Hand + Body + Shave Bar

DOVE MEN +CARE SKIN DEFENSE ANTIBACTERIAL 3-N-1 HAND + BODY + SHAVE BAR - Benzalkonium Chloride soap

Dove Men +Care Skin Defense Antibacterial 3-N-1 Hand + Body + Shave Bar

Drug Facts

Active ingredient

Benzalkonium Chloride (0.13%)

Purpose

Antibacterial

Uses

For washing to decrease bacteria on the skin

Warnings

- **For external use only**
- **When using this product** do not use in or near the eyes. In case of contact, rinse eyes thoroughly with wate **r**
- **Stop use and ask doctor** if irritation or redness develop.
- **Keep out of reach of children except under adult supervision.**

If swallowed, get medical help or contact a poison control Center right away.

Directions

Wet bar with water, lather vigorously and use all over body. Rinse clean.

Inactive ingredients

Sodium Lauroyl Isethionate, Stearic Acid, Sodium Oleate, Sodium Stearate, Water (Aqua), Sodium Isethionate, Lauric Acid, Sodium C14-16 Olefin Sulfonate, Sodium Laurate, Fragrance (Parfum), Dipropylene Glycol, Sodium Chloride, Tetrasodium

Etidronate, Tetrasodium EDTA, Kaolin*, Titanium Dioxide*, Yellow 5 (CI 19140), Yellow 6 (CI 15985). *Contains one or more of these ingredients

Questions or Comments?

Call toll-free 1-800-761-3683

Packaging

DOVE

men skin defense antibacterial 3n1 hand body shave bar soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:64942-1846
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 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
SODIUM LAUROYL ISETHIONATE (UNII: M590021Z02)	
SODIUM OLEATE (UNII: 399SL044HN)	
SODIUM STEARATE (UNII: QU7E2XA9TG)	
SODIUM ISETHIONATE (UNII: 3R36J71C17)	
LAURIC ACID (UNII: 1160N9NU9U)	
SODIUM LAURATE (UNII: K146MR5EXO)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
ETIDRONATE TETRASODIUM (UNII: CZZ9T1T1X4)	
KAOLIN (UNII: 24H4NWX5CO)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	
EDETATE SODIUM (UNII: MP1J8420LU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-1846-1	212 g in 1 CARTON; Type 0: Not a Combination Product	01/15/2021	
2	NDC:64942-1846-2	425 g in 1 CARTON; Type 0: Not a Combination Product	01/15/2021	
3	NDC:64942-1846-3	637 g in 1 CARTON; Type 0: Not a Combination Product	01/15/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/15/2021	

Labeler - Conopco Inc. d/b/a/ Unilever (001375088)

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