DR LIFT UNSCENTED ANTIBACTERIAL BODY WASH- benzalkonium chloride liquid
Spa de Soleil

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.

Dr Lift Unscented Antibacterial Body Wash

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Warnings

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For external use only. If product gets in eyes, rinse promptly and thoroughly with water. Discontinue use if irritation and redness occur. Stop use and ask a doctor if irritation or redness lasts for more than 72 hours. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Keep out of reach of children.

Directions

Directions: Wet your skin in the shower or bath. Lather body wash on your hands and use a loofah or a washcloth to apply it to your skin. Rinse it off and pat dry with a towel.

Place enough product in the palm of your hands to thoroughly cover you hands. Rub hands together briskly until product is completely absorbed and hands are dry.

Inactive Ingredients:

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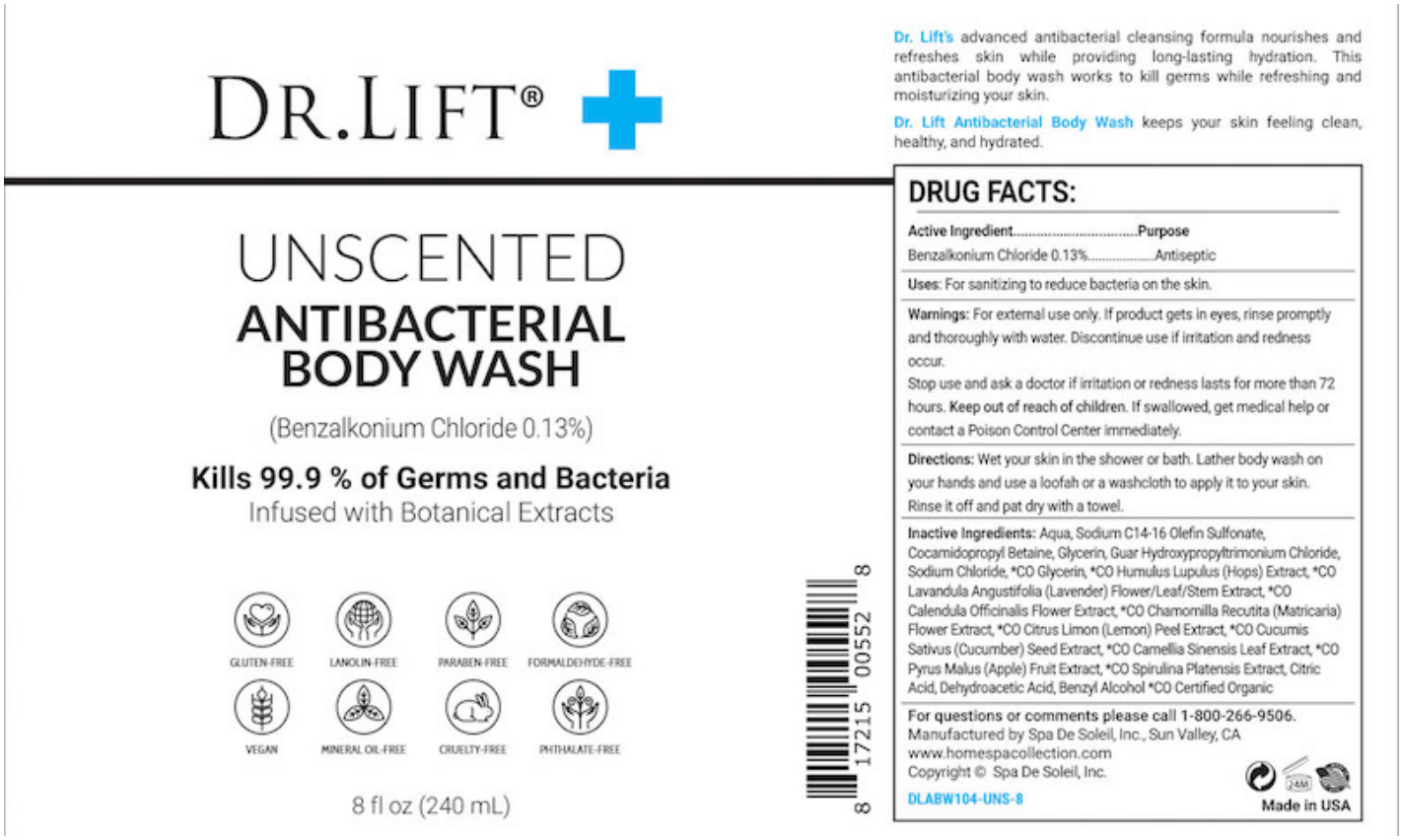
Aqua, Sodium C14-16 Olefin Sulfonate, Cocamidopropyl Betaine, Glycerin, Guar Hydroxypropyltrimonium Chloride, Sodium Chloride, *CO Glycerin, *CO Humulus Lupulus (Hops) Extract, *CO Lavandula Angustifolia (Lavender) Flower/Leaf/Stem Extract, *CO Calendula Officinalis Flower Extract, *CO Chamomilla Recutita (Matricaria) Flower Extract, *CO Citrus Limon (Lemon) Peel Extract, *CO Cucumis Sativus (Cucumber) Seed Extract, *CO Camellia Sinensis Leaf Extract, *CO Pyrus Malus (Apple) Fruit Extract, *CO Spirulina

Platensis Extract, Citric Acid, Dehydroacetic Acid, Benzyl Alcohol.

*CO Certified Organic



DR LIFT UNSCENTED ANTIBACTERIAL BODY WASH

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68062-2248
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.312 mg in 240 mL

Inactive Ingredients	
Ingredient Name	**Strength**
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68062-2248-1	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/15/2022	

Labeler - Spa de Soleil (874682867)

Establishment

Name	Address	ID/FEI	Business Operations
Spa de Soleil		874682867	manufacture(68062-2248)

Revised: 3/2022 Spa de Soleil