DR LIFT ANTIBACTERIAL BODY WASH- benzalkonium chloride gel
Spa de Soleil

Dr Lift Antibacterial Body Wash

Active Ingredient

Benzalkonium Chloride 0.13 %

Purpose

Antiseptic

Warnings

Warnings

For external use only.

When using this product: If in eyes, rinse promptly and thoroughly with water. Discontinue use if irritation or redness develop.

Stop use and ask a doctor if irritation or redness occur for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Directions

Apply to wet skin. Scrub thoroughly. Rinse and dry. Use as needed.

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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Sodium Chloride, Cocamidopropyl betaine, Aqua, Sodium Laureth Sulfate, Carbomer, 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, Xanthan Gum.

*CO Certified Organic

DR LIFT ANTIBACTERIAL BODY WASH

benzalkonium chloride gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68062-2241
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.46 mg in 240 mg

Inactive Ingredients

Ingredient Name	Strength
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End

#	Item Code	Package Description	Date	Date
1	NDC:68062-2241-1	240 mg in 1 TUBE; Type 0: Not a Combination Product	07/29/2020	
2	NDC:68062-2241-2	60 mg in 1 TUBE; Type 0: Not a Combination Product	10/10/2025	
3	NDC:68062-2241-3	100 mg in 1 TUBE; Type 0: Not a Combination Product	10/10/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	07/29/2020	

Labeler - Spa de Soleil (874682867)

Establishment

Name	Address	ID/FEI	Business Operations
Spa de Soleil		874682867	manufacture(68062-2241)

Revised: 10/2025 Spa de Soleil