

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.

Keep out of reach of children.

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



**ANTIBACTERIAL
BODY WASH**

Kills 99.9 % of Germs and Bacteria

Dr. Lift's advanced cleansing formula nourishes and refreshes skin while providing long-lasting hydration. This antibacterial body wash works to kill germs while refreshing and moisturizing your skin. **Dr. Lift Body Wash** keeps your skin feeling clean, healthy, and hydrated.


GLUTEN-FREE


LANOLIN-FREE


PARABEN-FREE


FORMALDEHYDE-FREE


VEGAN


MINERAL OIL-FREE


CRUELTY-FREE


PHTHALATE-FREE

8 fl. oz. | 240 ml

Drug Facts:

Active Ingredient.....Purpose
Benzalkonium Chloride 0.13%.....Antiseptic

Uses: For sanitizing to reduce bacteria on the skin.

Warnings: 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.

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.

Inactive Ingredients: 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, Fragrance. *CO Certified Organic

For questions or comments please call 1-800-266-9506. **Made in USA**

Manufactured by: **DLBW104**
Spa de Soleil, Inc., Sun Valley, CA
www.homespacollection.com

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: 5OCF3011KX)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End
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#	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