DR LIFT ANTIBACTERIAL- benzalkonium chloride lotion 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 Antibacterial Body Lotion

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Warnings

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.

Keep out of reach of children.

Directions

Place enough product in the palm of your hands to thoroughly cover your hands. Rub hands together until dry.

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, Isononyl Isononanoate, Polyacrylamide, C13-14 Isoparaffin, Laureth-7, Palmitoyl Tripeptide-5, Glycerin, *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, Ascorbic Acid (Vitamin C), Retinyl Palmitate (Vitamin A), Tocopheryl Acetate (Vitamin E), Benzyl Alcohol, Dehydroacetic Acid, Fragrance. *CO Certified Organic

DR.LIFT®



ANTIBACTERIAL **BODY LOTION**

Kills 99,99% of Germs and Bacteria

Infused with **Botanical Extracts**









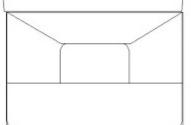








4 fl. oz. | 120 mL



Dr. Lift Antibacterial Body Lotion is specially formulated to kill germs and bacteria, while moisturizing. Infused with Calendula and Chamomile, the skin will look and feel soft and hydrated. Use with Dr. Lift's hygiene regiment collection including Dr. Lift Antibacterial Hand Sanitizer Gel, Hand Sanitizer Spray, Wipes, Body Scrub, Body Wash, Hand Soap, and our all-purpose, natural base surface cleaner for a germ fighting lifestyle!

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: Place enough product on hands to cover all surfaces. Rub hands together until dry.

Inactive Ingredients: Aqua, Isononyl Isononanoate, Polyacrylamide, C13-14 Isoparaffin, Laureth-7, Palmitoyl Tripeptide-5, Glycerin, "CO Glycerin, "CO Humulus Lupulus (Hops) Extract, *CO Lavandula Angustifola (Lavander) 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 Camelia Sinensis Leaf Extract. CO Pyrus Malus (Apple) Fruit Extract, *CO Spirulina Platensis Extract, Ascorbic Acid (Vitamin C), Retinyl Palmitate (Vitamin A), Tocopheryl Acetate (Vitamin E), Benzyl Alcohol, Dehydroacetic Acid, Fragrance. *CO Certified Organic

For questions or comments please call 1-800-266-9506.

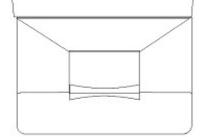
Made in USA DLABL112

Manufactured by: Spa de Soleil, Inc., Sun Valley, CA

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DR LIFT ANTIBACTERIAL

benzalkonium chloride lotion

Product	Inform	nation
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:68062-2243

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM -	BENZALKONIUM	0.156 mg
UNII:7N6JUD5X6Y)	CHLORIDE	in 120 mg

Inactive Ingredients

indetive ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6 A3C0 OX)			
WATER (UNII: 059QF0KO0R)			

Packaging

l	# Item Code Package Description		Marketing Start Date	Marketing End Date	
ı	1	NDC:68062-2243-1	120 mg in 1 BOTTLE; Type 0: Not a Combination Product	09/17/2020	

Marketing Information

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/17/2020	

Labeler - Spa de Soleil (874682867)

Establishment				
Name	Address	ID/FEI	Business Operations	
Spa de Soleil		874682867	manufacture(68062-2243)	

Revised: 9/2020 Spa de Soleil