ANTI BACTERIAL HAND GEL LAVENDER CLOUD- alcohol gel Bath & Body Works, Inc.

ABHG Lavender Cloud

Active Ingredient

Alcohol 71%

Purpose

Antiseptic

Use

Decreases bacteria on hands.

Warnings

Flammable: Keep away from flame or high heat.

For external use only.

When using this product keep out of eyes. Stop use and ask a doctor if irritation or redness develops.

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

Directions

Rub a dime sized drop into hands.

Water (Aqua, Eau), Fragrance (Parfum), Carbomer, Mannitol, Isopropyl Myristate, Cellulose, Hydroxyethyl Urea, Tocopheryl Acetate, Wheat Amino Acids, Aloe Barbadensis Leaf Juice, Butyrospermum Parkii (Shea) Butter Extract, Silica, Kaolin, Retinyl Palmitate, Caprylic/Capric Triglyceride, Hydroxypropyl Methylcellulose, Aminomethyl Propanol, Ultramarines (CI 77007), Red 33 (CI 17200), Blue 1 (CI 42090), Ext. Violet 2 (CI 60730), Yellow 5 (CI 19140).

Bath & Body Works, Distr., 95 West Main Street New Albany, OH 43054, 1-800-395-1001 pat. www.bbwinc.com/patents







.25" Hinge Area

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Fragrance (Parfum), Carbomer, Mannitol, Isopropyl Myristate, Cellulose, Hydroxyethyl Urea, Tocopheryl Acetate, Wheat Amino Acids, Aloe Barbadensis Leaf Julice, Butyrospermum Parkii (Shea) Butter Extract, Silice, Kaolin, Retinyl Palmitate, Caprylic/Capric Triglycaride, Hydroxypropyl Methylcellulose, Aminomethyl Propanol, Ultramarines (Cl 77007), Red 33 (Cl 17200), Blue 1 (Cl 42090), Ext. Violet 2 (Cl 60730), Yellow 5 (Cl 19140).

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ANTI BACTERIAL HAND GEL LAVENDER CLOUD

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:62670-6754

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

ALCOHOL

71 mL in 100 mL

Inactive Ingredients

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

Packaging

	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l		NDC:62670- 6754-0	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/18/2024	11/18/2027

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	505G(a)(3)	11/18/2024	11/18/2027

Labeler - Bath & Body Works, Inc. (878952845)

Establishment								
Name	Address	ID/FEI	Business Operations					
KDC US Holdings, Inc.		080783283	manufacture(62670-6754)					

Establishment							
Name	Address	ID/FEI	Business Operations				
Memphis Contract Packaging		117443103	manufacture(62670-6754)				

Revised: 1/2025 Bath & Body Works, Inc.