ANTI BACTERIAL HAND GEL VAMPIRE BLOOD- alcohol gel Bath & Body Works, Inc.

ABHG Vampire Blood

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

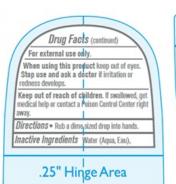
Rub a dime sized drop into hands.

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, Hydroxypropyl Methylcellulose, Caprylic/Capric Triglyceride, Retinyl Palmitate, Aminomethyl Propanol, Ultramarines (CI 77007), Red 33 (CI 17200), Red 4 (CI 14700), Yellow 5 (CI 19140), Ext. Violet 2 (CI 60730)

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







.25" Hinge Area

Fragrance (Parfum), Carbomer, Mannitol, Isopropyl Myristale, Cellulose, Hydrokyethyl Urea, Dicopheryl Acetate, Wheat Amino Aciks, Aloe Barbadensis Leaf Juice, Butynespermum Palrid (Shea) Buther Extract, Silica, Kaolin, Hydroxypropyl Methylcellulose, Caprylic/Capric Trighyceride, Retinyl Palmitate, Aminomethyl Propanol, Ultramarines (Cl 77007), Red 33 (Cl 17200), Red 4 (Cl 14700), Yellow 5 (Cl 19140), Ext. Violet 2 (Cl 60730).

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ANTI BACTERIAL HAND GEL VAMPIRE BLOOD

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:62670-6846

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958V90M) (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
I	NDC:62670-	29 mL in 1 BOTTLE; Type 0: Not a Combination	06/30/2025	06/30/2028

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/30/2025	06/30/2028

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

Establishment

Locabilonicite			
Name	Address	ID/FEI	Business Operations

KDC US Holdings, Inc.	080783283	manufacture(62670-6846)	
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Establishment				
Name	Address	ID/FEI	Business Operations	
Memphis Contract Packaging		117443103	manufacture(62670-6846)	

Revised: 7/2025 Bath & Body Works, Inc.