# ANTI-BACTERIAL HAND EUCALYPTUS SPEARMINT- alcohol gel Bath & Body Works, Inc.

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.

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#### **DRUG FACTS**

#### ACTIVE INGREDIENT

Alcohol 71%

#### **PURPOSE**

Antiseptic

#### **USE**

Decrease bacteria on hands.

#### **WARNINGS**

#### For external use only.

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

#### **FLAMMABLE**

Keep away from flame or high heat.

#### 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.

#### INACTIVE INGREDIENTS

Water (Aqua, Eau), Fragrance (Parfum), Carbomer, Lactose, Aminomethyl Propanol, Isopropyl Myristate, Cellulose, Propylene Glycol, Hydroxyethyl Urea, Tocopheryl Acetate, Wheat Amino Acids, Aloe Barbadensis Leaf Juice, Butyrospermum Parkii (Shea) Butter Extract, Hydroxypropyl Methylcellulose, Retinyl Palmitate, Ultramarines (CI 77007), Ext. Violet 2 (CI 60730), Yellow 5 (CI 19140).

#### COMPANY INFORMATION

Bath & Body Works, Distr. Reynoldsburg, Ohio 43068 1-800-395-1001

#### PRODUCT PACKAGING







### .25" Hinge Area

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Bath & Body Works, Distr., 95 West Main Stree New Albany, OH 43054, 1-800-395-1001 pet, www.lb.com/patents NOT TESTED ON ANIMALS

#### ANTI-BACTERIAL HAND EUCALYPTUS SPEARMINT

alcohol gel

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:62670-5892

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

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

#### **Inactive Ingredients**

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

## Packaging

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	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:62670-5892-	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/11/2020		
	NDC:62670-5892-	73 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/11/2020		
	3 NDC:62670-5892-	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/11/2020		

## **Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/11/2020	

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

Revised: 11/2020 Bath & Body Works, Inc.