# ANTI-BACTERIAL HAND FRENCH LAVENDER- 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 68%

#### **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), Isopropyl Alcohol, 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), Red 33 (CI 17200), Yellow 5 (CI 19140), Blue 1 (CI 42090), Ext. Violet 2 (CI 60730).

#### **COMPANY INFORMATION**

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

#### PRODUCT PACKAGING





# ANTI-BACTERIAL HAND FRENCH LAVENDER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62670-5787
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	68 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

## **Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62670-5787- 0	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/05/2019	
2	NDC:62670-5787- 1	73 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/05/2019	
3	NDC:62670-5787-3	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/05/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/05/2019	

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

Revised: 12/2019 Bath & Body Works, Inc.