# ANTI-BACTERIAL HAND BLUEBERRY CRUMBLE- 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), Blue 1 (CI 42090), Red 33 (CI 17200), Yellow 5(CI 19140), Ext. Violet 2 (CI 60730).

#### COMPANY INFORMATION

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

## PRODUCT PACKAGING







.25" Hinge Area

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Isopropyl Alcohol, Fragrance (Parfum), Carbomer, Lactose, Aminomethyl Propanol, Isopropyl Myristate, Cellulose, Propylene Glycol, Hydroxyethyl Urea, Tocopheryl Acetate, Wheat Amino Acids, Aloe Barbadensis Leaf Juice, Butryospermum Parkii (Shea) Butter Extract, Hydroxypropyl Methylcellulose, Retinyl Palmitate, Ultramarines (C177007), Blue 1 (C142090), Red 33 (C117200), Yellow 5(C119140), Ext. Violet 2 (C160730).

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

## ANTI-BACTERIAL HAND BLUEBERRY CRUMBLE

alcohol gel

## **Product Information**

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:62670-6003

Route of Administration TOPICAL

## **Active Ingredient/Active Moiety**

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)
ALCOHOL (UNII: 3K9958V90M) ALCOHOL 68 mL in 100 mL

## **Inactive Ingredients**

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

ľ	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:62670-6003- 0	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/14/2020		
2	NDC:62670-6003- 1	73 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/14/2020		
3	NDC:62670-6003-3	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/14/2020		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	12/14/2020			

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

Revised: 12/2020 Bath & Body Works, Inc.