ANTI-BACTERIAL HAND STRESS RELIEF 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.

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

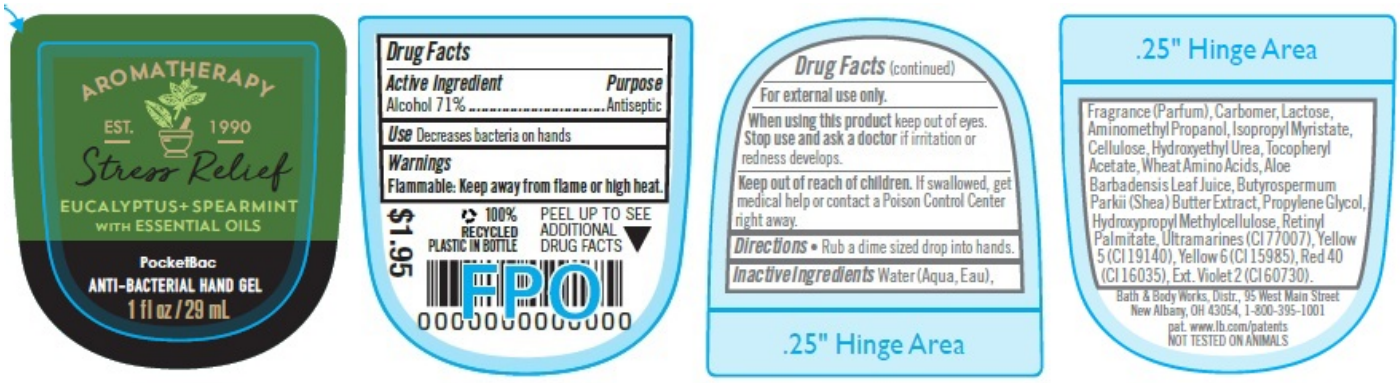
INACTIVE INGREDIENTS: Water (Aqua, Eau), Fragrance (Parfum), Carbomer, Lactose, Aminomethyl Propanol, Isopropyl Myristate, Cellulose, Hydroxyethyl Urea, Tocopheryl Acetate, Wheat Amino Acids, Aloe Barbadensis Leaf Juice, Butyrospermum Parkii (Shea) Butter Extract, Propylene Glycol, Hydroxypropyl Methylcellulose, Retinyl Palmitate, Ultramarines (CI 77007), Yellow 5 (CI 19140), Yellow 6 (CI 15985), Red 40 (CI 16035), 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 STRESS RELIEF EUCALYPTUS SPEARMINT

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62670-5933
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (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:62670-5933-0	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2020	
2	NDC:62670-5933-1	73 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2020	
3	NDC:62670-5933-3	225 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/09/2020	

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC monograph not final	part333E	12/09/2020	

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

Establishment

Name	Address	ID/FEI	Business Operations
Accel Inc.		838933430	relabel(62670-5933)

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