

ANTI-BACTERIAL HAND INTO THE NIGHT- 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 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.

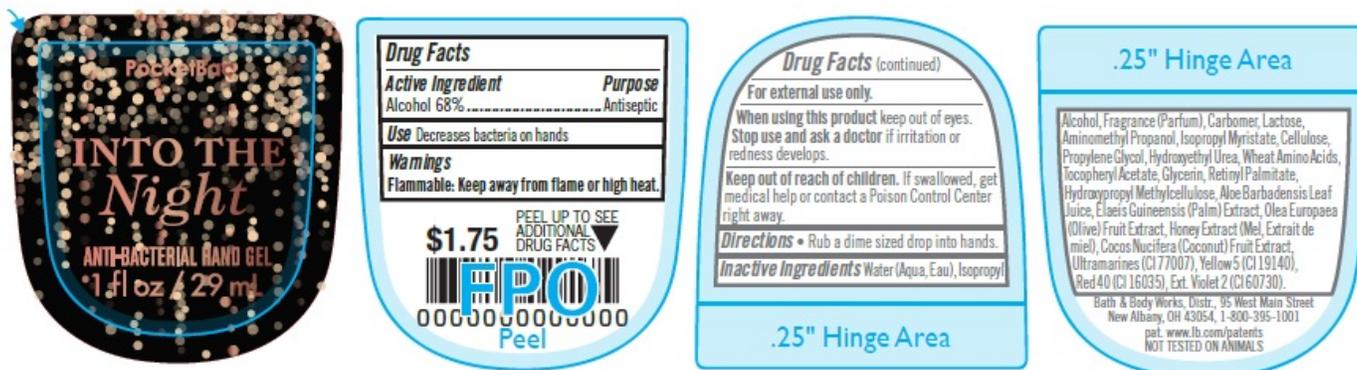
INACTIVE INGREDIENTS

Water (Aqua, Eau), Isopropyl Alcohol, Fragrance (Parfum), Carbomer, Lactose, Aminomethyl Propanol, Isopropyl Myristate, Cellulose, Propylene Glycol, Hydroxyethyl Urea, Wheat Amino Acids, Tocopheryl Acetate, Glycerin, Retinyl Palmitate, Hydroxypropyl Methylcellulose, Aloe Barbadensis Leaf Juice, Elaeis Guineensis (Palm) Extract, Olea Europaea (Olive) Fruit Extract, Honey Extract (Mel, Extrait de miel), Cocos Nucifera (Coconut) Fruit Extract, Ultramarines (CI 77007), Yellow 5 (CI 19140), 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 INTO THE NIGHT

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62670-5752
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: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62670-5752-0	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2019	
2	NDC:62670-5752-1	73 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2019	
3	NDC:62670-5752-3	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/06/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/06/2019	

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

Establishment

Name	Address	ID/FEI	Business Operations
Accel Inc.		838933430	relabel(62670-5752)

Revised: 12/2019

Bath & Body Works, Inc.