ANTI-BACTERIAL HAND HONOLULU SUN- alcohol spray
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.

ACTIVE INGREDIENT

Alcohol (72%)

PURPOSE

Antiseptic

USE

To decrease bacteria on skin.

WARNINGS

For external use only.

FLAMMABLE

Keep away from flame or high heat.

WHEN USING THIS PRODUCT

avoid contact with eyes. If contact occurs, rinse thoroughly with water.

STOP USE AND ASK A DOCTOR

if irritation and redness develop.

KEEP OUT OF THE REACH OF CHILDREN

If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

- Spray hands thoroughly.
- Rub until dry.

INACTIVE INGREDIENTS

Water (Aqua, Eau), Fragrance (Parfum), PEG-12 Dimethicone, Tocopheryl Acetate, Butyrospermum Parkii (Shea) Butter Extract, Aloe Barbadensis Leaf Juice, Propylene

Glycol, 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 HONOLULU SUN

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62670-6117
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	72 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:62670-6117-2	88 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/15/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/15/2021	

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

Revised: 6/2021 Bath & Body Works, Inc.