ANTI-BACTERIAL HAND MERRY HOLIDAY SPARKLE- 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.

ACTIVE INGREDIENT

Alcohol 68%

PURPOSE

Antiseptic

USE

Decrease bacteria on hands.

WARNINGS

For external use only.

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 or increase.

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 immediately.

DIRECTIONS

• Rub a dime sized drop into hands.

INACTIVE INGREDIENTS

Water (Aqua, Eau), Isopropyl Alcohol, Fragrance (Parfum), Honey Extract (Mel, Extrait de miel), Elaeis Guineensis (Palm) Extract, Cocos Nucifera (Coconut) Fruit Extract, Olea Europaea (Olive) Fruit Extract, Retinyl Palmitate, Tocopheryl Acetate, Wheat Amino Acids, Carbomer, Lactose, Cellulose, Hydroxyethyl Urea, Hydroxypropyl Methylcellulose, Glycerin, Isopropyl Myristate, Propylene Glycol, Aminomethyl Propanol, Ultramarines (CI 77007), Yellow 5 (CI 19140), Blue 1 (CI 42090), Red 40 (CI 16035).

COMPANY INFORMATION

Bath & Body Works, Distr. Reynoldsburg, Ohio 43068

1-800-395-1001

PRODUCT PACKAGING







Honey Extract (Mel. Extrait de miel), Elaeis Guineensis (Palm) Extract, Cocos Nucifera (Coconut) Fruit Extract, Olea Europaea (Olive) Fruit Extract, Retinyl Palmitate, Tocopheryl Acetate, Wheat Amino Acids, Carbomer, Lactose, Cellulose, Hydroxyethyl Urea, Hydroxypropyl Methylcellulose, Glycerin, Isopropyl Myristate, Propylene Glycol, Aminomethyl Propanol, Ultramarines (C177007), Yellow 5 (C119140), Blue 1 (C142990), Red 40 (C116035).

NOT TESTED ON ANIMALS Bath & Body Works, Distr. Reynoldsburg, Ohio 43068, 1-800-395-1001 Shop 24/7 at www.bathandbodyworks.com

ANTI-BACTERIAL HAND MERRY HOLIDAY SPARKLE

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:62670-4429
Route of Administration	TOPICAL	DEA Schedule	

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (ALCOHOL)	ALCOHOL	68 mL in 100 mL	

Inactive Ingredients	
Ingredient Name	Strength
WATER	

I	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62670-4429-0	29 mL in 1 BOTTLE		

Marketing Inform	mation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph not final	part333E	0 1/18/20 13	

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

Establishment				
Name	Address	ID/FEI	Business Operations	
Tri-Tech Laboratories, Inc.		792844680	manufacture(62670-4429)	

Revised: 1/2013 Bath & Body Works, Inc.