ANTI-BACTERIAL HAND JAPANESE CHERRY BLOSSOM- 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.

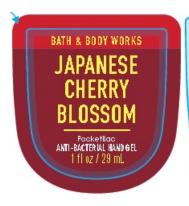
INACTIVE INGREDIENTS

Water (Aqua, Eau), Isopropyl Alcohol, Fragrance (Parfum), Propylene Glycol, Carbomer, Lactose, Aminomethyl Propanol, Isopropyl Myristate, Cellulose, Hydroxyethyl Urea, Wheat Amino Acids, Tocopheryl Acetate, Retinyl Palmitate, Glycerin, 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), Red 4 (CI 14700), 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

Bath & Body Works, Distr, 95 West Main Stree New Albany OH (\$054, 1-800-395-100) pat, www.lb.com/patents NOT TESTED ON ANIMALS

ANTI-BACTERIAL HAND JAPANESE CHERRY BLOSSOM

alcohol gel

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:62670-5651

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

68 mL in 100 mL

Inactive Ingredients

Ingredient Name Strength
WATER (UNII: 059QF0KO0R)

Packaging

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Item Code	Package Description	Marketing Start Date	Marketing End Date				
NDC:62670-5651- 0	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/18/2019					
NDC:62670-5651- 1	73 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/18/2019					
NDC:62670-5651- 3	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/18/2019					
0 N	NDC:62670-5651-	NDC:62670-5651- 29 mL in 1 BOTTLE; Type 0: Not a Combination Product NDC:62670-5651- 73 mL in 1 BOTTLE; Type 0: Not a Combination Product NDC:62670-5651- 236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination	NDC:62670-5651- 29 mL in 1 BOTTLE; Type 0: Not a Combination Product 06/18/2019 NDC:62670-5651- 73 mL in 1 BOTTLE; Type 0: Not a Combination Product 06/18/2019 NDC:62670-5651- 236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination 06/18/2019				

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	06/18/2019			

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

Revised: 6/2019 Bath & Body Works, Inc.