

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.

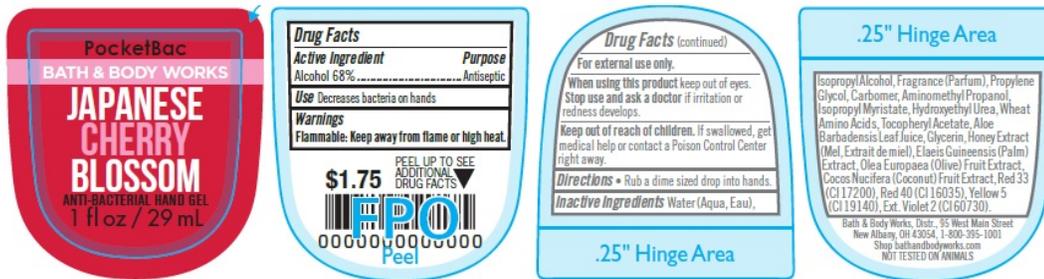
INACTIVE INGREDIENTS

Water (Aqua, Eau), Isopropyl Alcohol, Fragrance (Parfum), Cocos Nucifera (Coconut) Fruit Extract, Olea Europaea (Olive) Fruit Extract, Elaeis Guineensis (Palm) Extract, Honey Extract (Miel, Extrait de miel), Aloe Barbadensis Leaf Juice, Retinyl Palmitate, Tocopheryl Acetate, Wheat Amino Acids, Glycerin, Carbomer, Lactose, Aminomethyl Propanol, Isopropyl Myristate, Cellulose, Propylene Glycol, Hydroxyethyl Urea, Hydroxypropyl Methylcellulose, Ultramarines (CI 77007), Blue 1 (CI 42090), Red 33 (CI 17200).

COMPANY INFORMATION

Bath & Body Works, Distr.
Reynoldsburg, Ohio 43068

PRODUCT PACKAGING



ANTI-BACTERIAL HAND JAPANESE CHERRY BLOSSOM			
alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62670-5269
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			
		Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62670-5269-0	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/21/2017	
2	NDC:62670-5269-1	73 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/21/2017	
3	NDC:62670-5269-3	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/21/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/21/2017	

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

Revised: 11/2017

Bath & Body Works, Inc.