ANTI-BACTERIAL HAND BLACK CHERRY MERLOT- 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, Propylene Glycol, Lactose, Aminomethyl Propanol, Isopropyl Myristate, Cellulose, Hydroxyethyl Urea, Tocopheryl Acetate, Wheat Amino Acids, Aloe Barbadensis Leaf Juice, Butyrospermum Parkii (Shea) Butter Extract, Hydroxypropyl Methylcellulose, Retinyl Palmitate, Ultramarines (CI 77007), Red 33 (CI 17200), Yellow 5 (CI 19140), Blue 1 (CI 42090), 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 BLACK CHERRY MERLOT

alcohol gel

Product Information							
Product T ype	HUMAN OTC DRUG	Item Code (Source)		ND	NDC:62670-5788		
Route of Administration	TOPICAL						
Active Ingredient/Active Moiety							
Ingredient Name			Basis of Strength		Strength		
ALCOHOL (UNII: 3K9958V90M) (A)		ALCOHOL		68 mL in 100 mL			
Inactive Ingredients							
Inactive Ingredients	Ingredient Name				Strength		
Inactive Ingredients WATER (UNII: 059QF0K00R)	Ingredient Name				Strength		
Inactive Ingredients WATER (UNII: 059QF0KO0R)	Ingredient Name				Strength		

#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:62670-5788- 0	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/0 5/20 19				
2	NDC:62670-5788- 1	73 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/0 5/20 19				
3	NDC:62670-5788- 3	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/0 5/20 19				
Marketing Information							
Marketing Category		ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not fina		nal part333E	12/05/2019				

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

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Bath & Body Works, Inc.