ANTI-BACTERIAL HAND SWEET AS STRAWBERRIES 2- 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, Lactose, Aminomethyl Propanol, Isopropyl Myristate, Cellulose, Propylene Glycol, Hydroxyethyl Urea, Tocopheryl Acetate, Wheat Amino Acids, Aloe Barbadensis Leaf Juice, Butyrospermum Parkii (Shea) Butter Extract, Glycerin, Hydroxypropyl Methylcellulose, Retinyl Palmitate, 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









ANTI-BACTERIAL HAND SWEET AS STRAWBERRIES 2

alcohol gel

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:62670-5767

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 1	lngredients
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Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:62670-5767- 0	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2019		
2	NDC:62670-5767-	73 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2019		
3	NDC:62670-5767-3	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/06/2019		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	12/06/2019		

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

Revised: 12/2019 Bath & Body Works, Inc.