ANTIBACTERIAL HAND SANITIZER- alcohol gel
Body One Products , 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.

Antibacterial Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Uses

- for handwashing to decrease bacteria on the skin
- recommended for repeated use.

Warnings

For external use only. Flammable, keep away from fire or flame.

Do not use in the eyes. In case of contact, rinse eyes with water.

Stop use and ask a doctor if irritation and redness develop and presist for more than 72 hours.

Keep out of reach of children. If swallowed get medical help or contact poison control center right away.

Directions

- wet hands thoroughly with product
- briskly rub hands together until dry
- supervise children in the use of this product

Other information

- store at 20° to 20°C (68 to 77° F)
- may discolor certain fabrics

Inactive ingredients

Water, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer,PEG 60 Almond Glycerides, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate,Triisopropanolamine, Fragrance

PRINCIPAL DISPLAY PANEL - 1064 ML Bottle Label

BodyOne
Products

ANTIBACTERIAL

HAND SANITIZER

kills 99.99%
of all germs

(32 FL.OZ. 1064ML)

ANTIBACTERIAL HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73563-080
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**

Propylene Glycol (UNII: 6DC9Q167V3)	
water (UNII: 059QF0KO0R)	
PEG-60 ALMOND GLYCERIDES (UNII: 4Y0E651N0F)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TRIISOPROPANOLAMINE (UNII: W9EN9DLM98)	
CARBOMER INTERPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 132584PQMO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73563-080-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/10/2020	
2	NDC:73563-080-18	532 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/10/2020	
3	NDC:73563-080-32	1064 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/10/2020	07/31/2020
4	NDC:73563-080-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020	
5	NDC:73563-080-04	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020	
6	NDC:73563-080-01	29.5 mL in 1 TUBE; Type 0: Not a Combination Product	08/05/2020	
7	NDC:73563-080-06	66 mL in 1 TUBE; Type 0: Not a Combination Product	07/20/2020	
8	NDC:73563-080-07	7 mL in 1 PACKET; Type 0: Not a Combination Product	05/15/2020	
9	NDC:73563-080-12	355 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	03/10/2020	

Labeler - Body One Products , Inc. (117376115)

Registrant - BMC 1092, Inc. dba Solo Laboratories, Inc. (078831987)

Establishment

Name	Address	ID/FEI	Business Operations
BMC 1092, Inc. dba Solo Laboratories, Inc.		078831987	MANUFACTURE(73563-080) , LABEL(73563-080) , PACK(73563-080)

Revised: 8/2020 Body One Products , Inc.