

ANTIBACTERIAL HAND SANITIZER- alcohol gel

Maketa,LLC

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 hand washing 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 persist 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 20-25° C (68 to 77°F)
- may discolor certain fabrics

Inactive Ingredients

Water, Propylene Glycol, Acrylates/C10-3- Alkyl Acrylate Crosspolymer, Peg-60, Almond Glycerides, Aloe Barbadosensis Leaf Juice, Triisopropanolamine, Fragrance

Questions?

(800) 638-8149

PRINCIPAL DISPLAY PANEL - 59.15 ml Bottle Label

Makéta™

Made in the
U.S.A.

antibacterial
HAND
SANITIZER

kills 99.99%
of all germs

2 fl. oz. (59.15 ml)



ANTIBACTERIAL HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72102-829
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: 059QF0K00R)	
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:72102-829-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/16/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	04/16/2020	

Labeler - Maketa,LLC (081031516)

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(72102-829) , LABEL(72102-829) , PACK(72102-829)

Revised: 4/2020

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