

BIOMAX USA HAND SANITIZER- ethyl alcohol liquid

Biomax Cosmetics

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.

Biomax ADVANCED Hand Sanitizer

▣Drug Facts

Active Ingredient

Ethyl alcohol 70% v/v

▣Purpose

Antiseptic

▣Inactive Ingredient

Water(Aqua), SDA Alcohol, Triethanolamine, Carbomer, Aloe Barbadensis, Fragrance, Glycerin, Propylene Glycol, Tocopheryl Acetate (Vitamin E), CI 42090, CI 19140.

▣Uses:

For hand washing to help reduce bacteria on the skin.

▣Directions

- Put a small amount in your palm and rub hands together until dry.
- Children under 6 years of age should be supervised when using this product. Recommended for repeated use.

Warnings

For external use only-hands.

Flammable. Keep away from heat and flame.

Stop use and ask a doctor if irritation or redness develops and lasts.

▣When using this product

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- If Swallowed, get medical help or contact a Poison Control Center immediately.

▣ADVANCED Hand Sanitizer

MOISTURIZING WITH VITAMIN E & ALOE

KILLS 99.99% OF MOST GERMS

MADE IN USA

Manufactured by:

Biomax Cosmetics Inc.

Houston, TX 77061

www.biomaxcosmetics.com

BENEFITS:

- With Vitamin E.
- Leaves hands feeling soft & refreshed.
- Kills more than 99.99% of germs and bacteria.

Packaging

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77369-927
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE (UNII: V5VD430YW9)	

GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77369-927-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/08/2020	
2	NDC:77369-927-06	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/08/2020	
3	NDC:77369-927-08	236.6 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/08/2020	
4	NDC:77369-927-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/08/2020	
5	NDC:77369-927-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/08/2020	
6	NDC:77369-927-01	3785 mL in 1 JUG; Type 0: Not a Combination Product	07/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/08/2020	

Labeler - Biomax Cosmetics (084032031)

Establishment

Name	Address	ID/FEI	Business Operations
Biomax Cosmetics		084032031	manufacture(77369-927)

Revised: 7/2020

Biomax Cosmetics