BIOMAX USA- is opropyl 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.

Bio max FIRST AID ANTISEPTIC

DRUG FACTS

Active Ingredient: 70% Isopropyl Alcohol

Purpose: Antiseptic

USES

- hand sanitizer to decrease bacteria on the skin
- recommended for repeated use
- for use when soap and water are not available

Inactive Ingredients: IPurified Water/aqua

WARNINGS

For external use only. If taken internally serious gastric disturbances will result.

Flammable - keep away from heat, spark, electrical, fire or flame. Ask a doctor before use for deep or puncture wounds, animal bites or serious burns.

WHEN USING THIS PRODUCT:

- Do not get into eyes.
- Do not inhale.
- Do not apply over large areas of the body.
- Do not use longer than 1 week.

STOP USE AND ASK A DOCTOR IF CONDITION PERSISTS OR GETS WORSE.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

IDIRECTIONS

• supervise children under 6 years of age when using this product to avoid swallowing

HEALTHCARE

ISOPROPYL RUBBING ALCOHOL 70%

WARNING: FLAMMABLE,

KEEP AWAY FROM FIRE OR FLAME

ANTISEPTIC

- Topical antiseptic and sanitizer
- Antibacterial Cleansing Agent

CAUTION: Fumes can be acutely irritating to skin, eyes and the respiratory system. Do not apply to irritated skin or if excessive irritation develops. Avoid getting into the eyes or on mucous membranes. Avoid inhaling this product.

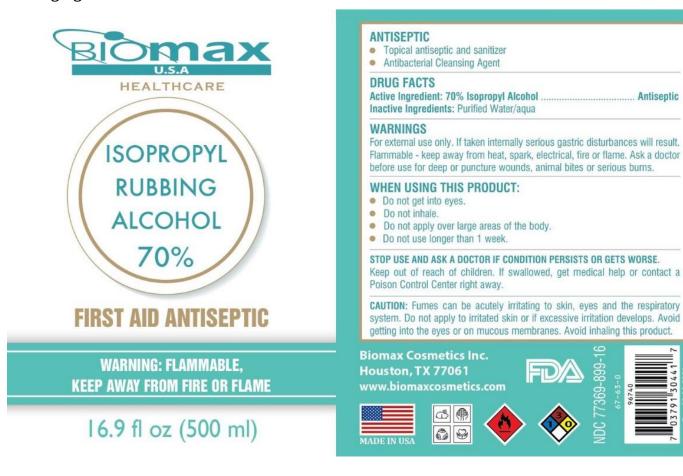
Biomax Cosmetics Inc.

Houston, TX 77061

www.biomaxcosmetics.com

MADE IN USA

Packaging



BIOMAX USA isopropyl alcohol liquid Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:77369-899 Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 mL in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77369-899-16	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/05/2020	
2	NDC:77369-899-01	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/05/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	08/05/2020		

Labeler - Biomax Cosmetics (084032031)

Establishment			
Name	Address	ID/FEI	Business Operations
Biomax Cosmetics		084032031	manufacture(77369-899)

Revised: 8/2020 Biomax Cosmetics