

**ZUUM KLIN ANTIBACTERIAL- alcohol gel**  
**U.S. Cotton Mexico, S. de R.L. de C.V.**

*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.*

-----

**Zuum Klin Gel Antibacterial**

***Drug Facts***

***Active ingredient***

Ethyl alcohol 63%

***Purpose***

Antiseptic

***Use***

For hand washing to decrease bacteria on the skin

***Warnings***

**For external use only**

**Flammable, keep away from fire or flame**

**Do not use**

in the eyes

**Stop use and ask a doctor**

- irritation and redness develop
- condition persists for more than 72 hours

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away.

***Directions***

wet hands thoroughly with product and allow to dry without wiping

***Other information***

Keep this product away from heat, fire or sunlight.

***Inactive ingredients***

Isopropyl alcohol, deionized water, aloe vera (aloe barbadensis) extract, propylene glycol, carbomer, aminomethyl propanol, fragrance, EDTA, vitamin E, vitamin A, ceramides

***Questions or Comments?***

**Package Labeling:**



**ZUUM KLIN ANTIBACTERIAL**

alcohol gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:77388-001
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.63 mL in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>CARBOMER HOMO POLYMER, UNSPECIFIED TYPE</b> (UNII: 0A5MM307FC)	
<b>AMINOMETHYLPROPANOL</b> (UNII: LU49E6626Q)	
<b>EDETIC ACID</b> (UNII: 9G34HU7RV0)	
<b>.ALPHA.-TOCOPHEROL</b> (UNII: H4N855PNZ1)	
<b>VITAMIN A</b> (UNII: 81G40H8B0T)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77388-001-01	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/01/2020	

**Labeler** - U.S. Cotton Mexico, S. de R.L. de C.V. (816167605)

### Establishment

Name	Address	ID/FEI	Business Operations
U.S. Cotton Mexico, S. de R.L. de C.V.		816167605	manufacture(77388-001)

Revised: 11/2020

U.S. Cotton Mexico, S. de R.L. de C.V.