

**ADVANCED HAND SANITIZER- hand sanitizer liquid**  
**Flex Beauty Labs, 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.*

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**Drug Facts**

**Active Ingredient**

Ethyl Alcohol 62.0%

**Purpose**

Antimicrobial

**Uses**

Hand Sanitizer to help reduce bacteria on skin that may cause disease.

**Warnings**

Flammable. Keep away from heat or flame.

For external use only.

**Keep out of reach of children.**

**When using this product,**

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water. Stop use and ask doctor if irritation or rash appears and lasts. Keep out of reach for children.

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

**Directions:**

- . Place enough Product in your palm to thoroughly spread on both hands and rub into the skin until dry.
- . Children under 6 years of age should be supervised when using this product.

**Other Information:**

- . Store below 106°F. (41°C)

**Inactive Ingredients:**

Water (Aqua), Aloe Vera Leaf , Carbomer, Fragrance, Glycerin, Propylene Glycol, Tocopheryl Acetate (Vitamin E), Triethanolamine,

FRONT LABEL



INSIDE OF BACK LABEL



BACK LABEL



## ADVANCED HAND SANITIZER

hand sanitizer liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72308-001
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0K00R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72308-001-01	295 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/07/2018	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/02/2017	

**Labeler** - Flex Beauty Labs, LLC (080858917)

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Flex Beauty Labs, LLC