

**SEALUXE HAND SANITIZER- ethyl alcohol gel**  
**APR Beauty Group Inc.**

*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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**Naturaluxe Hand Sanitizer**

**Drug Facts**

**Active Ingredient**

Ethyl Alcohol, 70% w/w

**Purpose**

Antiseptic

**Uses**

Kills harmful bacteria/germs

**Warnings**

**Flammable, keep away from open flame and sources of heat.**

**For occasional and personal domestic use.**

**Stop use and consult a health care practitioner if irritation develops.**

Avoid contact with eyes.

If contact occurs, rinse thoroughly with water.

**If swallowed, call a poison control centre.**

**Directions**

Rub thoroughly into hands for at least 30 seconds until dry.

For children 2 years and over supervise when product is used

**Other Information**

**Recommended storage conditions:**

20 Degree Celsius

**Inactive Ingredients**

Aloe barbadensis leaf juice, Aminomethyl Propanol, Glycerin, Propylene Glycol, Water, Isopropyl Myristate, Carbomer







## SEALUXE HAND SANITIZER

ethyl alcohol gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:79959-001
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
TOCOPHEROL (UNII: R0ZB2556P8)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79959-001-24	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/28/2020	
2	NDC:79959-001-50	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/28/2020	
3	NDC:79959-001-35	315 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/28/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/28/2020	

**Labeler** - APR Beauty Group Inc. (204198006)

Revised: 8/2020

APR Beauty Group Inc.