

**CLEAN IS IN HAND SANITIZER ANTISEPTIC FOAM- hand sanitizer antiseptic foam aerosol, foam**

**Beauty-Lab 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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**Ikoo Care clean is in Hand Sanitizer Antiseptic Foam**

**Warnings**

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For external use only.

Flammable. Keep away from fire or flame.

Contents under pressure.

- Do not puncture or incinerate.
- Do not store above 104°F (40°C)

When using this product

- Avoid contact with eyes.If contact occurs, rinse immediately and thoroughly with water.
- Do not use on children under 2 months of age.
- Do not use on open wounds.
- Discontinue use if irritation/redness occurs.

Stop use and ask a doctor if irritation occurs. This may be a sign of a serious condition.

Keep out of reach of children. •In case of ingestion, get medical help or contact a Poison Control Center immediately.

**Inactive Ingredients**

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Water, Hydrofluorocarbon 152a, Isobutane, Cetearyl alcohol, Polysorbate 60, Cetyl Lactate, Steareth-2, Sodium Benzoate, Sodium Sesquicarbonate, Fragrance

**Active Ingredients**

Active Ingredient.....	Purpose
Alcohol 62% v/v.....	Antiseptic

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**Keep out of Reach**

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**Uses**

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•To help decrease bacteria/germs on the skin. •For use when soap and water are not available. •Recommended for repeated use.

## Dosage

•Spray product on hands, enough to cover all surfaces. Rub hands together until all surfaces are wet and fully covered. Continue rubbing until hands feel dry. Do not rinse or wipe off sanitizer.

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## PACKAGE LABEL. PRINCIPAL DISPLAY PANEL



## CLEAN IS IN HAND SANITIZER ANTISEPTIC FOAM

hand sanitizer antiseptic foam aerosol, foam

Product Information		
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:80473-030
Route of Administration	Topical	
Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 g in 100 g
Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
1,1-DIFLUOROETHANE (UNII: 0B1U8K2ME0)		
Isobutane (UNII: BXR49TP611)		

<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>Polysorbate 60</b> (UNII: CAL22UVI4M)	
<b>Cetyl Lactate</b> (UNII: A7EVH2RK4O)	
<b>Steareth-2</b> (UNII: V56DFE46J5)	
<b>Sodium Benzoate</b> (UNII: OJ245FE5EU)	
<b>Sodium Sesquicarbonate</b> (UNII: Y1X815621J)	

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80473-030-01	200 g in 1 CONTAINER; Type 0: Not a Combination Product	09/10/2020	

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/10/2020	

**Labeler** - Beauty-Lab LLC (053878473)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Accra-Pac, Inc. (DBA KIK Custom Products)		024213616	manufacture(80473-030)

Revised: 9/2020

Beauty-Lab LLC