

ALLURE CHEMICALS INSTANT FOAM HAND SANITIZER- alcohol liquid
Allure Chemicals LP

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.

ALLURE CHEMICALS INSTANT FOAM HAND SANITIZER

Drug Facts

Active ingredient

Ethanol Alcohol 70%

Purpose

Antiseptic

Uses [s]

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame.

Do not use: In children less than 2 months of age and open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs.

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

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30°C (59-86°F)
- Avoid Freezing and excessive heat above 40°C (104°F)

Inactive Ingredients:

Cetaryl alcohol, Propylene glycol, Sodium Lauryl Ether Sulfate

ALCOHOL LIQUID - MEDICAL GRADE

KILLS 99.9% OF GERMS, COLD AND FLU VIRUSES

Manufactured and Packaged by

ALLURE Chemicals

9650-A Railroad Drive

El Paso Texas 79924

www.allurechemicals.com

Packaging

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alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78804-777
Route of Administration	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
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78804-777-08	3800 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/21/2021	
2	NDC:78804-777-09	250 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/21/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/13/2020	

Labeler - Allure Chemicals LP (080442496)**Establishment**

Name	Address	ID/FEI	Business Operations
Allure Chemicals LP		080442496	manufacture(78804-777)

Revised: 10/2020

Allure Chemicals LP