ANTISEPTIC HAND SANITIZER FOAM- ethyl alcohol liquid US Chemical Corporation

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.

U S Chemical Antiseptic Hand Sanitizer Foam

Drug Facts

Active Ingredient

Ethyl alcohol 62% v/v

Purpose

Antibacterial Agent

Uses

• Hand sanitizer to help reduce bacteria on the skin that could cause disease. Recommended for repeated use.

Warnings

Flammable, keep away from fire or flame. For external use only.

Avoid contact with eyes. In case of contact, flush with plenty of water.

Stop use and ask a doctor if irritation or rash appears and persists.

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

EMERGENCY TELEPHONE: 1-866-923-4913

Directions

- Place enough product in your palm to thoroughly cover your hands. Rub hands together briskly until dry.
- Children under 6 years of age should be supervised when using this product.

Other Information

• Store in a cool, dry place.

Inactive Ingredient

• Water, Glycerin, PEG-10 Dimethicone

Providing Exceptional Quality Since 1962

Distributed by/Distribuido por:

U S Chemical 316 Hart Street · Watertown, WI 53094 www.uschemical.com Emergency Phone 1-866-923-4913

Packaging



Antiseptic Hand Sanitizer Foam

NET CONTENTS/ CONTENIDO NETO: 1 gal. (3.79 L)





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NDC XXXX-XX-XX





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ANTISEPTIC HAND SANITIZER FOAM

ethyl alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:61307-105

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90 M) (ALCOHOL - UNII: 3K9958 V90 M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients Ingredient Name Strength WATER (UNII: 059QF0KO0R) GLYCERIN (UNII: PDC6A3C0OX) PEG-10 DIMETHICONE (600 CST) (UNII: 8 PR7V1SVM0)

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:61307-105-10	3790 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/10/2020		
Marketing Information				
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not f	inal part333A	06/10/2020		

Labeler - US Chemical Corporation (031457842)

Revised: 6/2020 US Chemical Corporation