HAND SANITIZER FOAM- alcohol solution
Custom Chemical Solutions

77955-002

Active ingredients (v/v)

Ethyl Alcohol 70%

Purpose

Antimicrobial

Uses

- To decrease bacteria on the skin that could cause disease
- Recommended for repeated use

Warnings

For external use only: hands

Flammable. Keep away from fire or flame.

When using this product avoid contact with eyes.

In case of eye contact, flush eyes with water.

Avoid contact with broken skin.

Do not ingest or inhale.

Stop use and ask a doctor if irritation or redness develops, or if condition persists for more than 72 hours.

Keep out of reach of children.

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

Directions:

Pump enough product into palm of hand to thoroughly cover all surfaces of both hands.

Rub hands together briskly until dry.

For children under 6, use only under adult supervision.

Other information:

Store below 110 F (43 C).

May discolor some fabrics.

May be harmful to wood finishes and plastics.

Inactive ingredients:

Water (Eau), Bis-PEG-10 Dimethicone, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Denatonium Benzoate

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Manufactured for:
Custom Chemical Solutions, LLC.,
167 Commerce Dr., Loveland, OH, 45140 USA

HAND
SANITIZER
FOAM

UniFirst Corporation 88UF 1L (1000 mL, 33.8 US fl. oz.)

HAND SANITIZER FOAM

alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77955-002
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**
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
GLYCERIN (UNII: PDC6A3C0OX)	
BIS-PEG-10 DIMETHICONE/DIMER DILINOLEATE COPOLYMER (UNII: CF5W1YCX11)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:77955-002-02	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/05/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/05/2020	

Labeler - Custom Chemical Solutions (081096319)

Establishment

Name	Address	ID/FEI	Business Operations
Kutol Products Company		004236139	manufacture(77955-002)

Revised: 1/2024 Custom Chemical Solutions