EXTENDED CARE FOAMING HAND SANITIZER- alcohol liquid
TRIPLE A SUPPLIES, 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.

EXTENDED CARE™ Foaming Hand Sanitizer

Active ingredient

Ethyl Alcohol 70%

Purpose

Antimicrobial

Use

Hand sanitizer to help reduce bacteria on the skin that could cause disease

Warnings

Flammable. Keep away from fire or flame.

For external use only.

When using this product, do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts

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

Directions

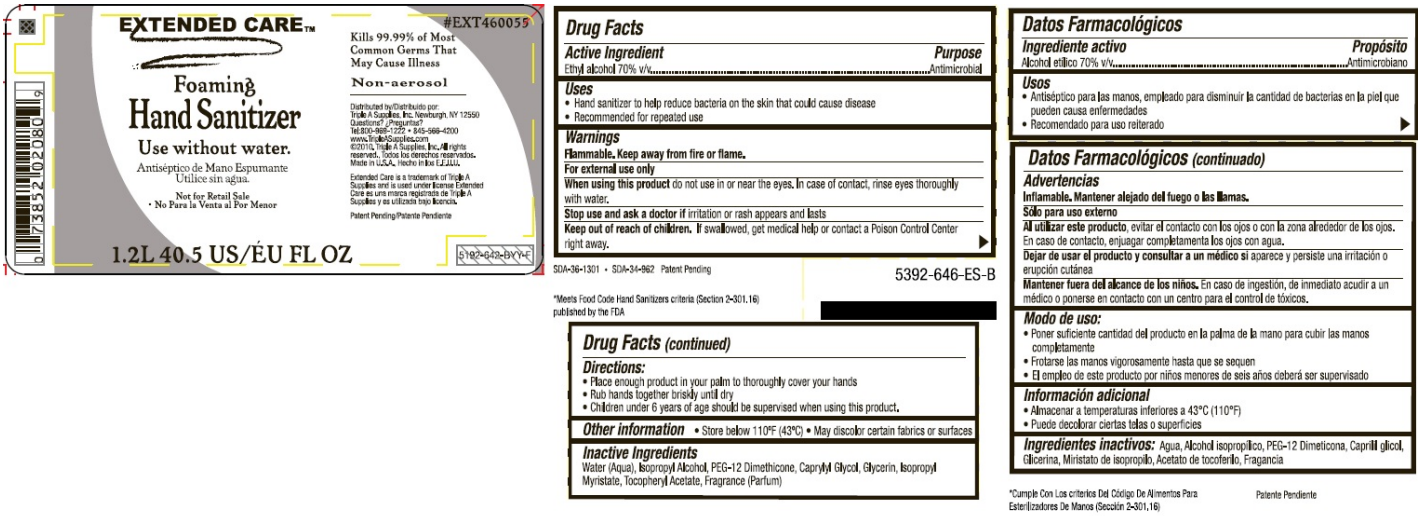
Place enough product in your palm to thoroughly cover your hands.

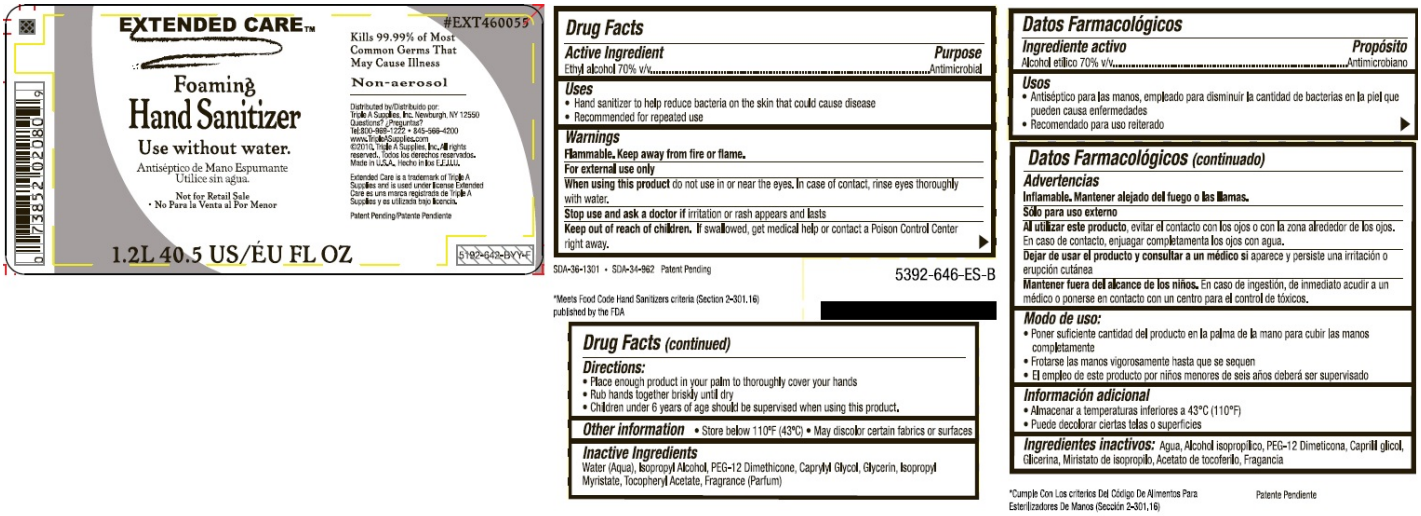
Rub hands together briskly until dry.

Children under 6 years of age should be supervised when using this product.

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, PEG-12 Dimethicone, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Fragrance (Parfum)





EXTENDED CARE FOAMING HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71875-800
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.70 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Isopropyl Alcohol (UNII: ND2M416302)	
Glycerin (UNII: PDC6A3C0OX)	
Isopropyl Myristate (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	

Caprylyl Glycol (UNII: 00YIU5438U)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71875-800-42	1250 mL in 1 PACKAGE; Type 0: Not a Combination Product	05/15/2014	09/30/2021
2	NDC:71875-800-40	1200 mL in 1 PACKAGE; Type 0: Not a Combination Product	05/15/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/15/2014	

Labeler - TRIPLE A SUPPLIES, INC. (872642913)

Revised: 9/2021 TRIPLE A SUPPLIES, INC.