

ALPET E3 HAND SANITIZER FOAM- alcohol liquid
Best Sanitizers, Inc.

Drug Facts 5900-125

Active ingredient

Alcohol 62.0%

Purpose

Hand Sanitizer

Uses

- sanitizes hands to help reduce bacteria that potentially can cause disease
- helps to prevent cross-contamination
- recommended for repeated use
- helps prevent drying of skin

Warnings

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

When using this product

- do not use in or near eyes
- use in a well-ventilated area

Stop use and consult a doctor if irritation or redness develop.

Keep out of reach of children.

If swallowed, seek medical attention or contact Poison Control Center immediately

Directions

- receive an adequate amount of foam into palm of hand
- rub thoroughly over all surfaces of both hands
- rub hands together until dry

Other Information

store in a cool, dry place below 104 °F (40 °C)

Inactive Ingredients

Water, DEA-C8-18 perfluoroalkylethyl phosphate, glycereth-2 cocoate, denatonium benzoate, tert-butyl alcohol

Questions?

Contact Best Sanitizers at **888-225-3267** Mon-Fri 9am - 4 pm PST

Maximize the Clean

BEST SANITIZERS INC.

ALPET E3

Hand Sanitizer Foam

For Food Industry

Professionals

Meets all FDA (GRAS) and

USDA requirements for

food handling

Enhanced with moisturizers

to keep skin soft and healthy

NSF certified E3 classification

Kosher certified and Pareve

Kills

99.999%

of tested pathogens

in 15 seconds

BEST

SANITIZERS

INC.

NSF

Nonfood Compounds

Program Listed E3

Registration Number 143167

Net Contents:

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Meets all FDA (GRAS) and USDA requirements for food handling

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- NSF certified E3 classification
- Kosher certified and Pareve

Kills **99.999%** of tested pathogens in 15 seconds



Nonfood Compounds Program Listed E3 Registration #143167

Manufactured by:
Best Sanitizers, Inc.
P.O. Box 1360
Penn Valley, CA 95946
www.bestsanitizers.com



Net Contents:

ALPET E3 HAND SANITIZER FOAM

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59900-125
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DIETHANOLAMINE BIS(C8-C18 PERFLUOROALKYLETHYL)PHOSPHATE (UNII: 4J55VM509S)	
GLYCERETH-2 COCOATE (UNII: JWM00VS7HC)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	

TERT-BUTYL ALCOHOL (UNII: MD83SFE959)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59900-125-02	208197 mL in 1 DRUM; Type 0: Not a Combination Product	05/01/2019	
2	NDC:59900-125-05	1250 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2019	
3	NDC:59900-125-08	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2019	
4	NDC:59900-125-03	3785 mL in 1 JUG; Type 0: Not a Combination Product	05/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/01/2019	

Labeler - Best Sanitizers, Inc. (957473614)

Establishment

Name	Address	ID/FEI	Business Operations
Best Sanitizers, Inc.		627278224	manufacture(59900-125)

Revised: 12/2024

Best Sanitizers, Inc.