

HAND SANITIZER FOAM- ethanol alcohol aerosol, foam
PurCel Labs LLC

Active Ingredient

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Uses

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

- On children less than 2 months of age.
- On open skin wounds.

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 occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a poison control center right away.

Directions

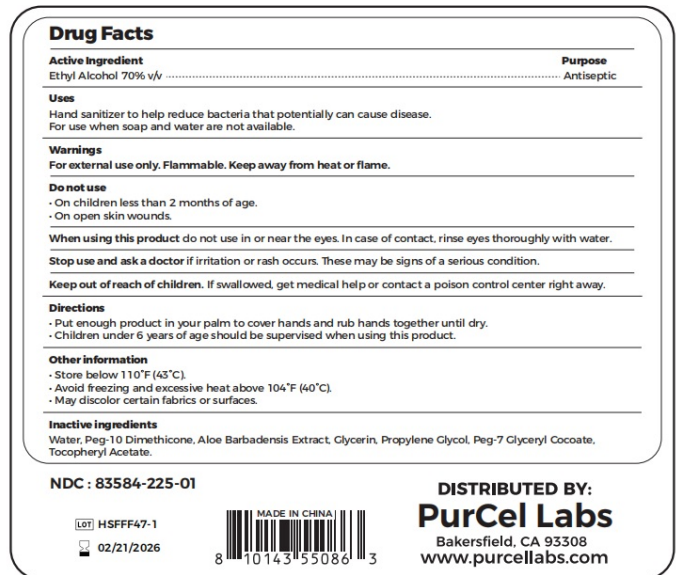
- Put enough product in your palm to cover hands and rub hands together until dry.
- Children under 6 years of age should be supervised when using this product.

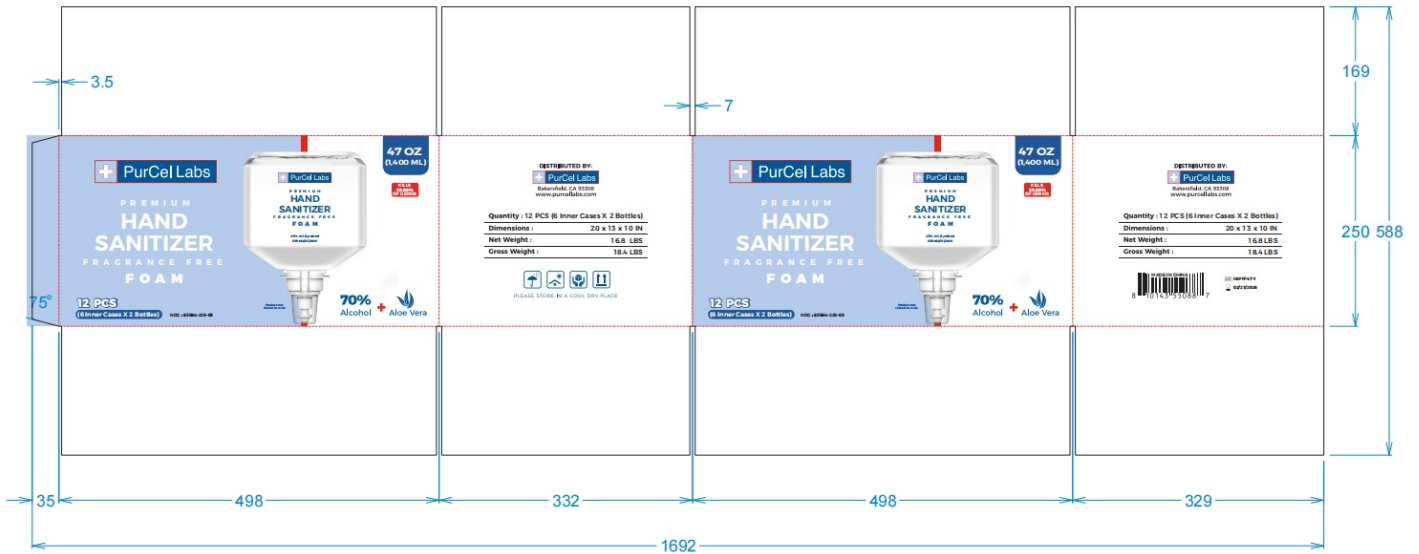
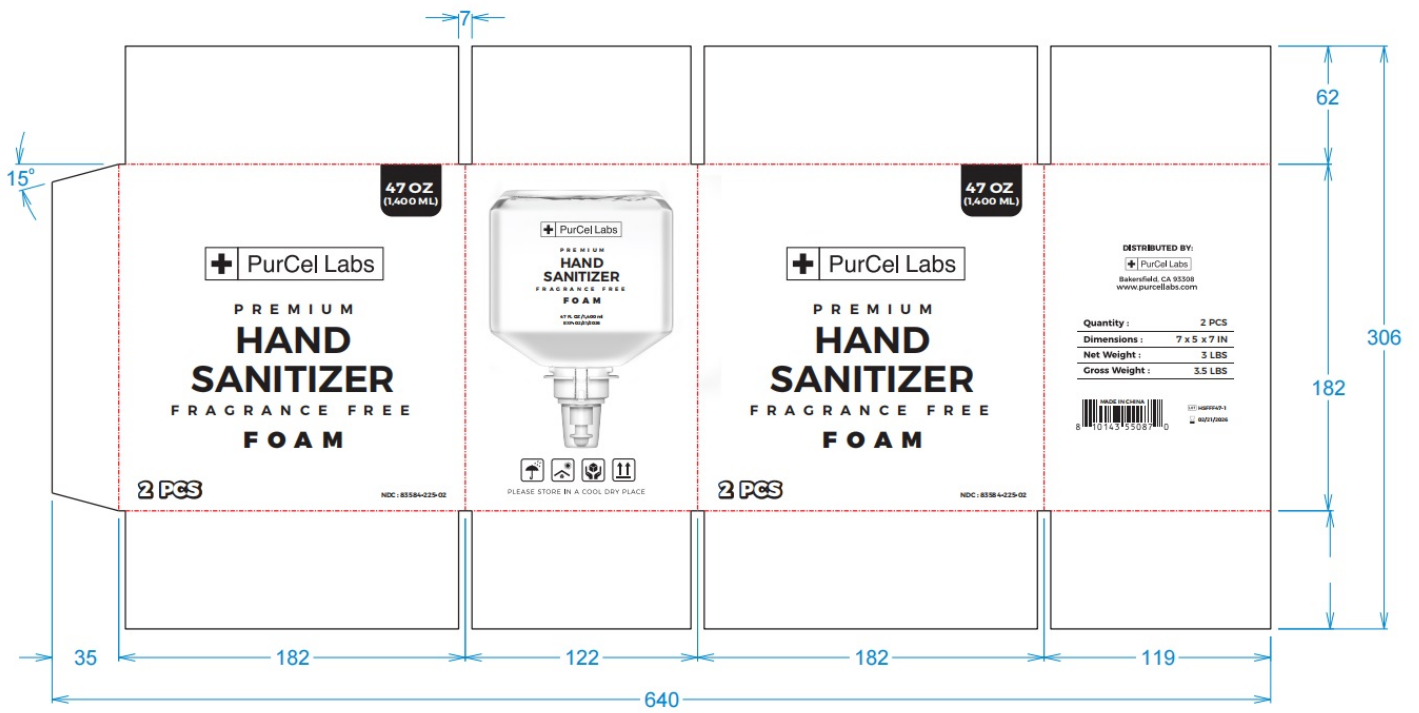
Other information

- Store below 110°F (43°C).
- Avoid freezing and excessive heat above 104°F (40°C).
- May discolor certain fabrics or surfaces.

Inactive ingredients

Water, Peg-10 Dimethicone, Aloe Barbadensis Extract, Glycerin, Propylene Glycol, Peg-7 Glyceryl Cocoate, Tocopheryl Acetate.





HAND SANITIZER FOAM

ethanol alcohol aerosol, foam

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83584-225
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
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83584-225-03	6 in 1 CARTON	02/29/2024	
1	NDC:83584-225-02	2 in 1 BOX		
1	NDC:83584-225-01	1400 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	02/29/2024	

Labeler - PurCel Labs LLC (127672531)

Registrant - PurCel Labs LLC (127672531)

Establishment

Name	Address	ID/FEI	Business Operations
Zhongshan Dermey Commodity Co.,Ltd		550280464	manufacture(83584-225)

Revised: 3/2024

PurCel Labs LLC