DOCTORS KLINE AND GREEN BOARD CERTIFIED DERMATOLOGIST HAND SANITIZER ANTISEPTIC- alcohol aerosol, foam
Formulated Solutions, LLC

Doctors Kline & Green Board Certified Dermatologist Hand Sanitizer Antiseptic

Drug Facts

Active ingredient

Ethyl Alcohol 62% v/v

Purpose

Antiseptic

Uses(s)

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. Extermely flammable, do not use near heat or flame or while smoking.

Do not use

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

When using this product

- keep out of eyes, ears, and mouth
- In case of contact with eyes, rinse eyes thoroughly with water.
- Do not puncture or incinerate
- Contents under pressure
- Do not expose to heat or store at temperature above 120°F (49°C)

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

- Hold can upside down and dispense into palm. Place enough product on hands to cover all surfaces. Rub hands together until dry
- Supervise children under 6 years of age when using this product to avoid swallowing.

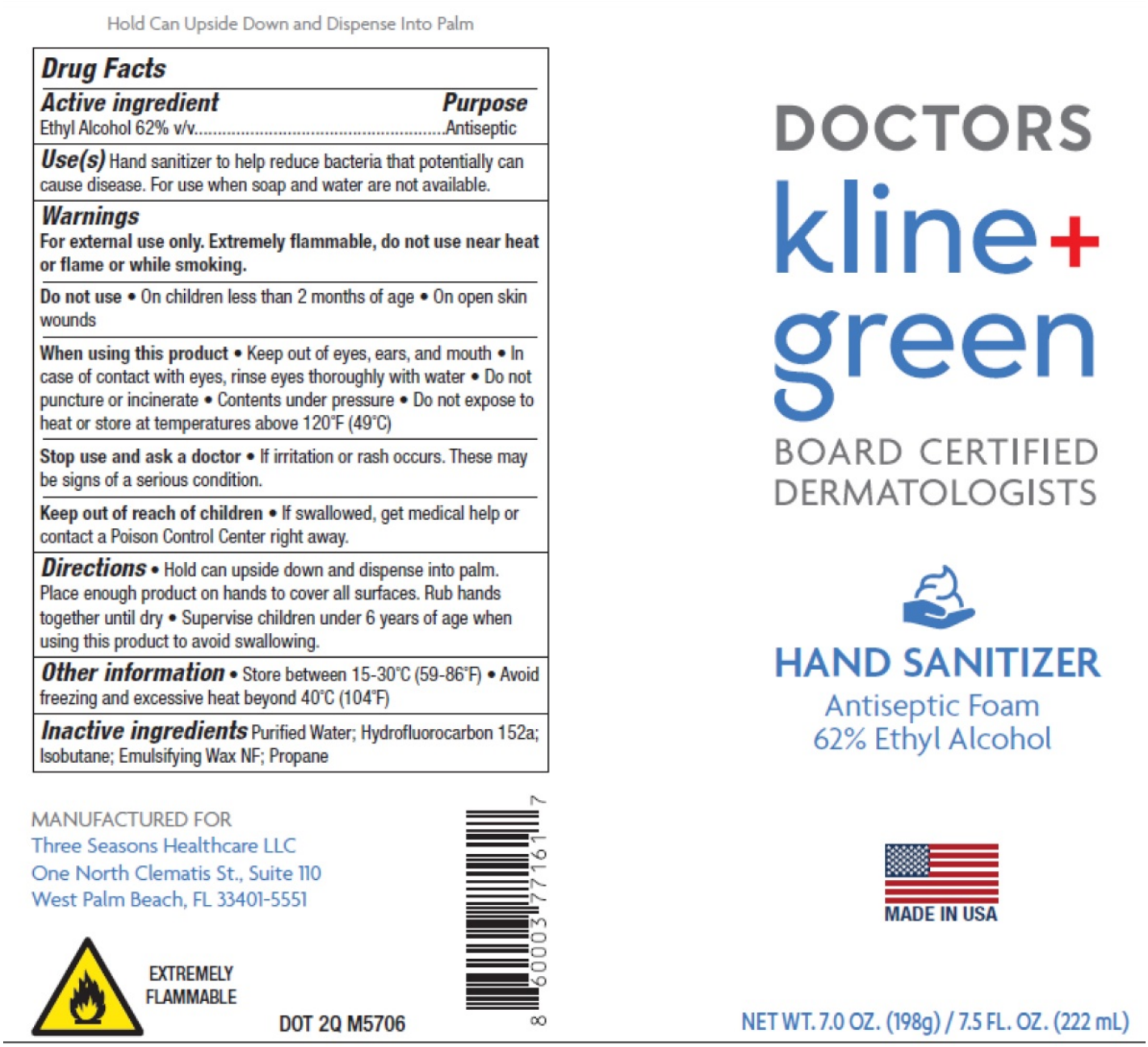
Other information

- Store between 15-30°C (59-86°F)
- Avoid freezing and excessive heat beyond 40°C (104°F)

Inactive ingredients

Purified Water, Hydrofluorocarbon 152a; Isobutane; Emulsifying Wax NF; Propane

Package Labeling:



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alcohol aerosol, foam

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:23667-104
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)	
1,1-DIFLUOROETHANE (UNII: 0B1U8K2ME0)	
ISOBUTANE (UNII: BXR49TP611)	
PROPANE (UNII: T75W9911L6)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:23667-104-00	222 mL in 1 CAN; Type 0: Not a Combination Product	02/09/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/09/2021	

Labeler - Formulated Solutions, LLC (143266687)

Revised: 11/2023 Formulated Solutions, LLC