

SUMMER FRIDAYS HAND SANITIZER- hand sanitizer spray

Naturich Labs

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.

Summer Fridays 2oz Hand Sanitizer

Active Ingredients

Alcohol 65% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

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

In 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. Stop use and ask 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 Poison Control Center right away.

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Directions

Place enough product on hands to cover all surfaces. Rub hands together until dry. Supervise children under 6 years of age when using product to avoid swallowing.

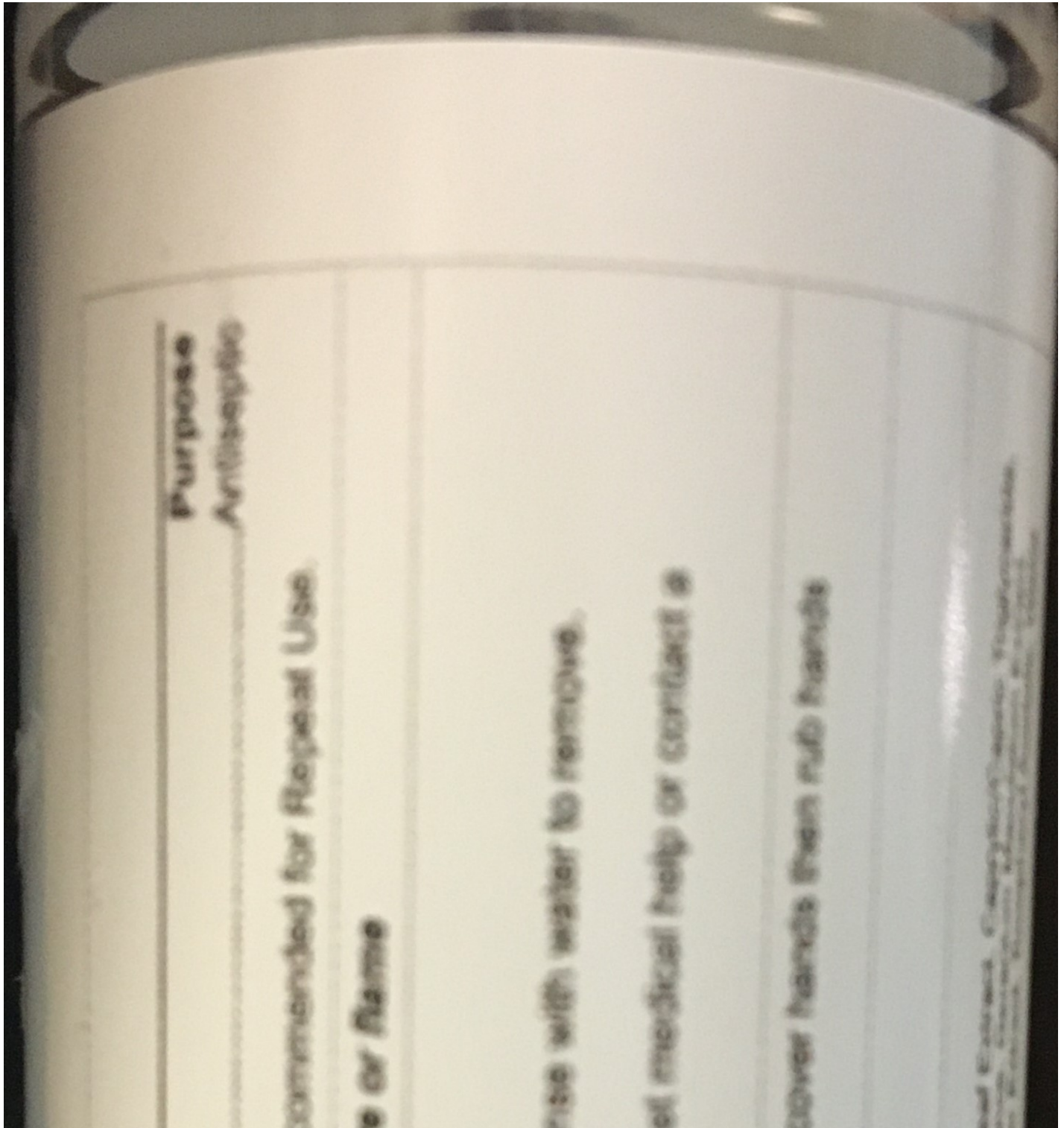
Other information

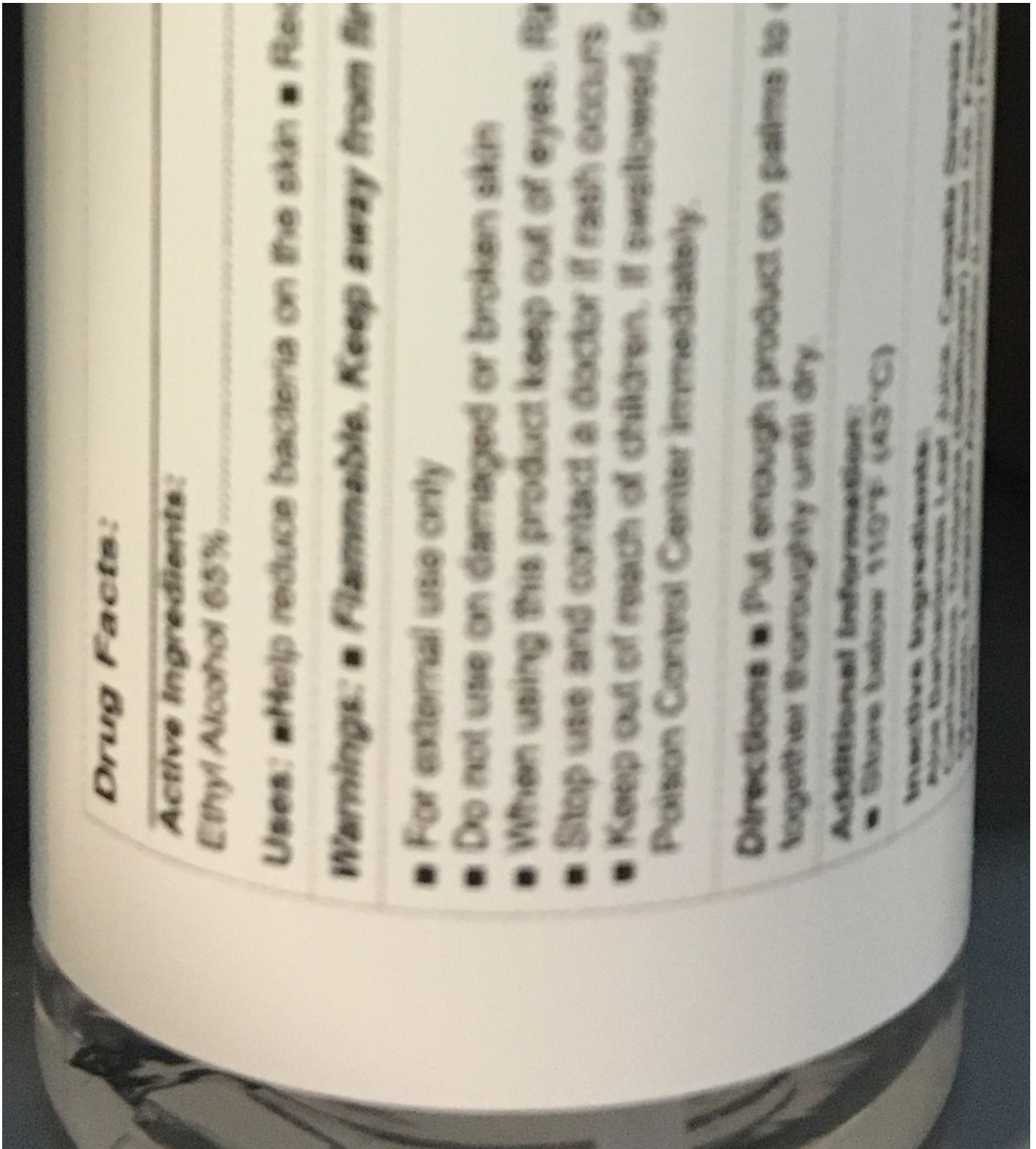
Store between 15-30C (59-86F)

Avoid freezing and excessive heat above 40C (104F)

Inactive Ingredients

Aloe Barbadensis Leaf Juice, Camellia Sinensis Leaf Extract, Caprylic/Capric Triglyceride, Carthamus Tinctorius (Safflower) Seed Oil, Fragrance, Geranium Maculatum Extract, Glycerin, Lavandula Angustifolia (Lavender) Flower Extract, Tocopheryl Acetate, Water (Aqua).





SUMMER FRIDAYS HAND SANITIZER

hand sanitizer spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72496-1127
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	35 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72496-1127-1	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2021	

Labeler - Naturich Labs (124281663)**Registrant** - Naturich Labs (124281663)**Establishment**

Name	Address	ID/FEI	Business Operations
Naturich Labs		124281663	manufacture(72496-1127)

Revised: 4/2023

Naturich Labs