**PHARMACYS PRESCRIPTION HAND SANITIZER- alcohol gel**
**American Consumer Products Corp**

*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.*

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**Pharmacys Prescription Hand Sanitizer**

**Active Ingredient**

**Active Ingredients -** Ethyl Alcohol 75%

**Purpose**

Antiseptic

**Uses**

**Uses -** Helps reduce bacteria on the skin that could cause disease. Recommended for repeated use.

**Warnings**

**Warnings -** For external use only. Do not ingest or swallow.

Flammable. Keep away from fire or flame.

Do not appy around eyes. Do not use in ears & mouth.

**When using this product,** avoid contact with eyes. In case of contact, flush eyes with water.

**Stop use and ask a doctor**

**Stop use and ask a doctor if** redness or irritation develops and persists for more than 72 hours.

Keep out of reach of children. Do not use on children less than 2 months of age. Supervise use in children under 6 years of age to prevent accidental swallowing. If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

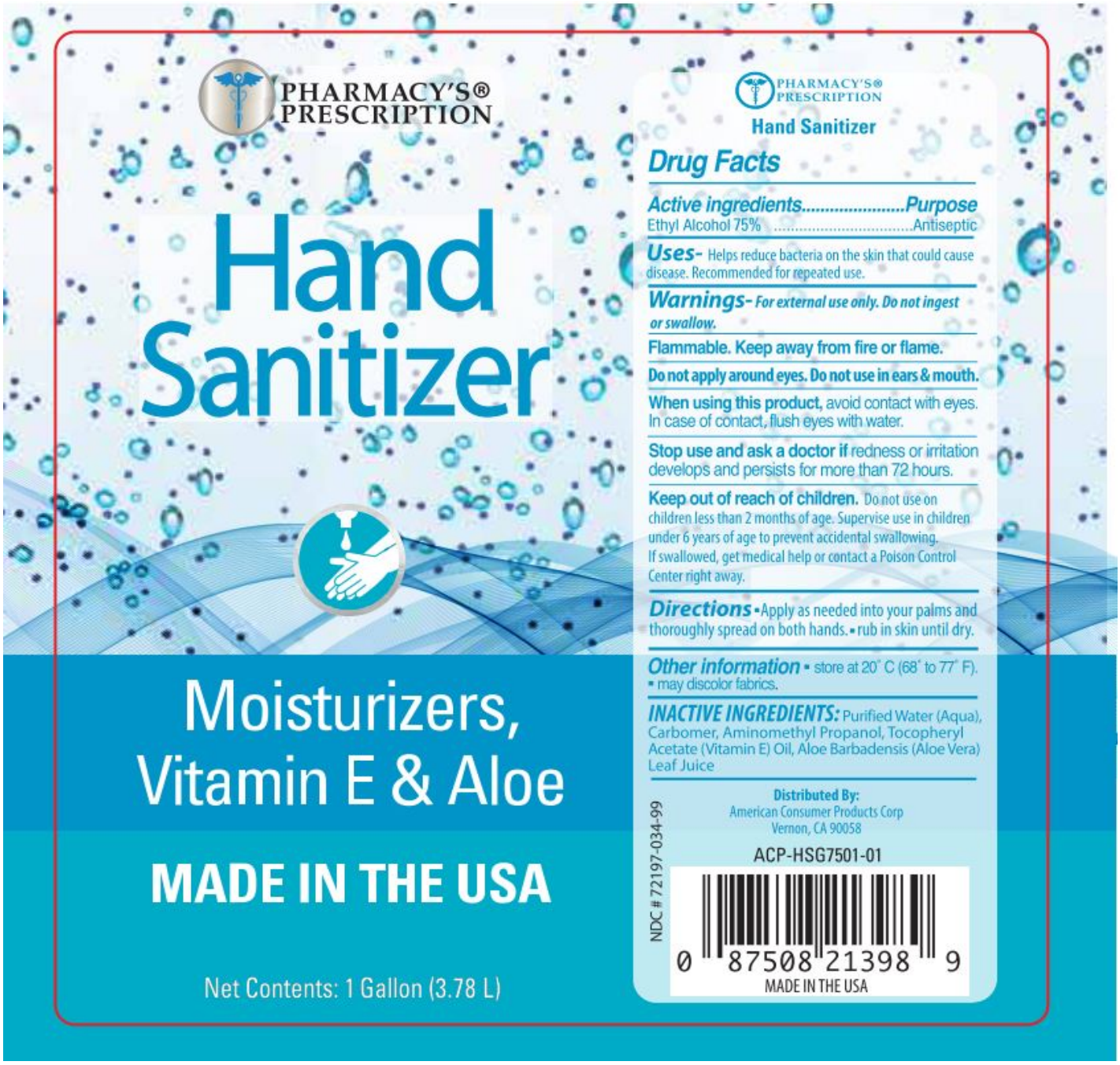
Directions - apply as needed into your palms and thoroughly spread on both hands. Rub into skin until dry.

**Other information**

Other information - store at 20º C (68º to 77º F). May discolor fabrics.

**INACTIVE INGREDIENTS:** Purified Water (Aqua), Carbomer, Aminomethyl Propanol, Tocopheryl Acetate (Vitamin E), Aloe Barbadensis (Aloe Vera) Leaf Juice

**Pharmacys Prescription Hand Sanitizer**




# PHARMACYS PRESCRIPTION HAND SANITIZER

alcohol gel

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:72197-034 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **ALCOHOL** (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 75 mL in 100 mL |

| Inactive Ingredients | |
|---|---|
| Ingredient Name | Strength |
| **CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE** (UNII: 0A5MM307FC) | |
| **ALOE VERA LEAF** (UNII: ZY81Z83H0X) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **AMINOMETHYLPROPANOL** (UNII: LU49E6626Q) | |
| **.ALPHA.-TOCOPHEROL ACETATE** (UNII: 9E8X80D2L0) | |

| Packaging | | | | |
|---|---|---|---|---|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:72197-034-99 | 3785.41 mL in 1 BOTTLE; Type 0: Not a Combination Product | 08/14/2020 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333A | 08/14/2020 | |

## Labeler - American Consumer Products Corp (081101181)

Revised: 8/2020 American Consumer Products Corp