

**PHARMACYS PRESCRIPTION 8OZ 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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**Pharmacy's Prescription 8OZ Hand Sanitizer**

**Active Ingredient**

**Active Ingredients:** Ethyl Alcohol 62%

**Purpose**

**Purpose:** Antiseptic

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

**Warnings**

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

**Flammable. Keep away from fire or flame.**

**Do not apply around eyes. Do not use in ears & mouth.**

**Stop Use**

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

**Keep out of reach of children**

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

**Inactive Ingredients**

**Inactive ingredients** - water, triethanolamine, glycerin, propylene glycol, tocopheryl acetate, aloe barbadensis gel, carbomer, fragrance.

**Directions**

**Directions** - pump as needed into your palms and thoroughly spread on both hands.- rub into skin until dry.

**Indications & Usage Section**

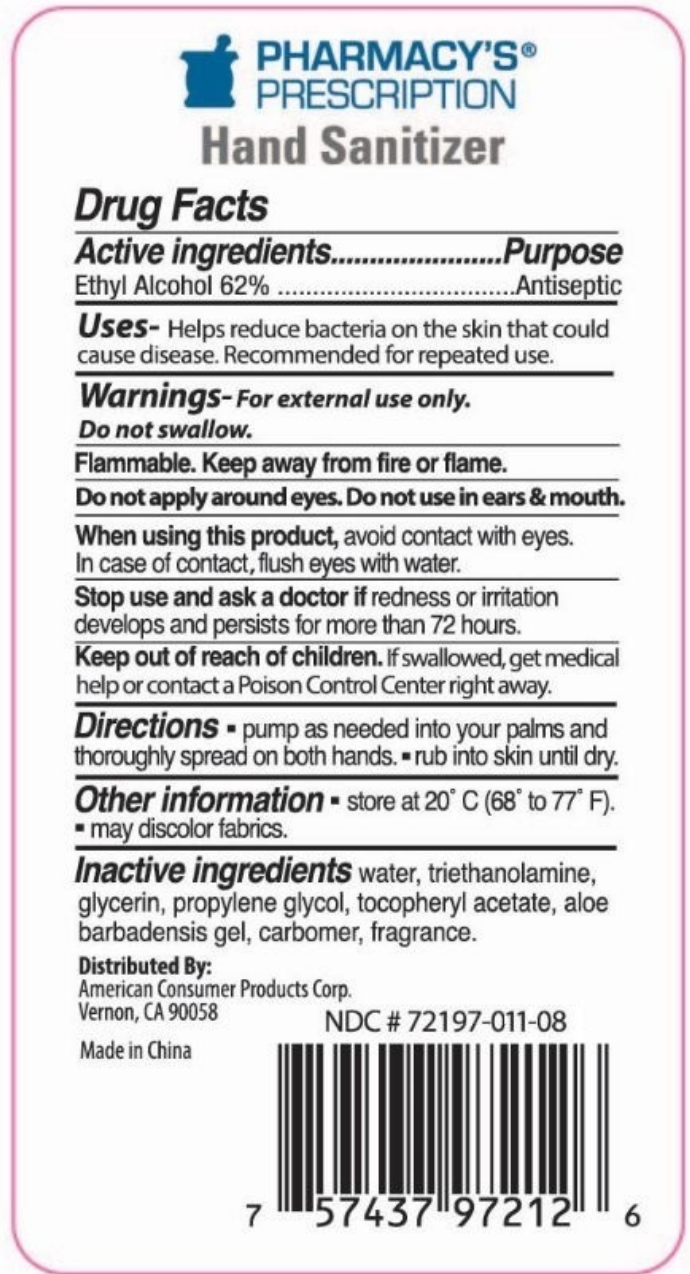
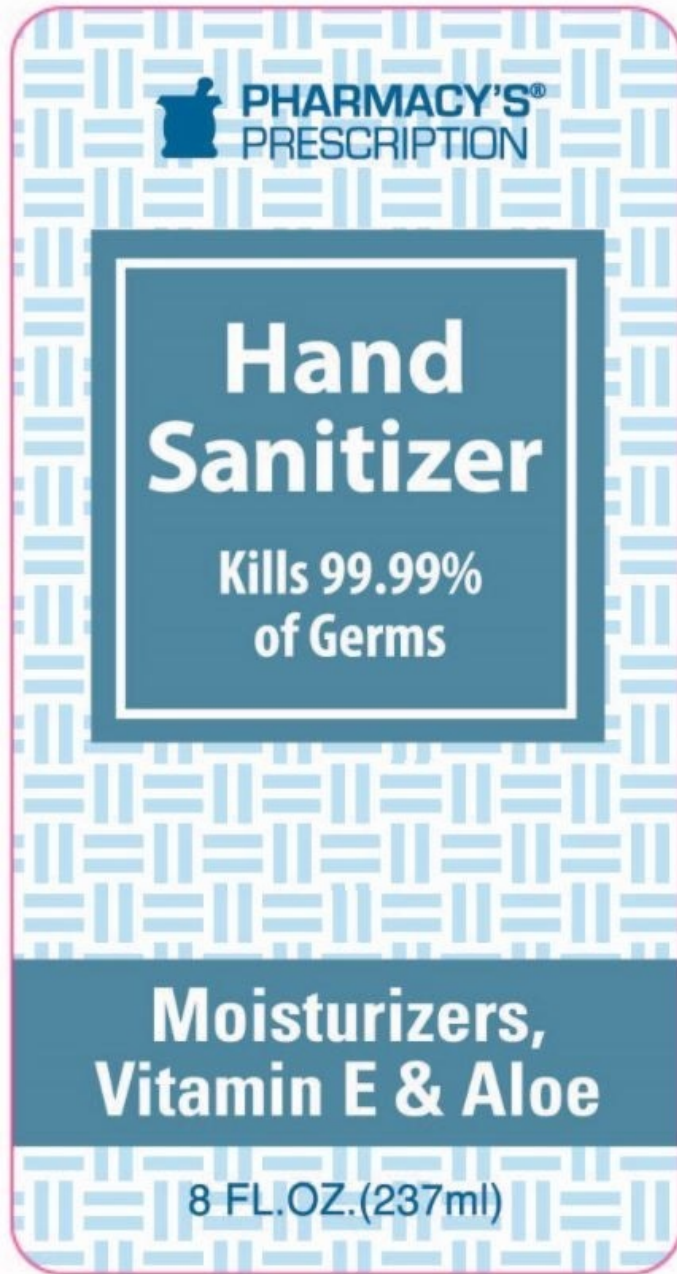
**When using this product,** avoid contact with eyes. In case of contact, flush eyes with water.

**Other Information**

**Other Information** - store at 20o C (68o to 77o F).

- may discolor fabrics.

Pharmacy's Prescription 8 OZ Label



**PHARMACYS PRESCRIPTION 8OZ HAND SANITIZER**

alcohol gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:72197-011
<b>Route of Administration</b>	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)	
TRIETHANOLAMINE 2-CYCLOHEXYL-4,6-DINITROPHENOLATE (UNII: N2TK31JIAH)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
ALOE ARBORESCENS WHOLE (UNII: 0EF00WD7LU)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL 2-(2-METHYLBUTYRATE) (UNII: 3Z73C506A9)	
CARBOMER HOMO POLYMER TYPE C (UNII: 4Q93RCW27E)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72197-011-08	237 mL in 1 BOTTLE, PUMP; 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 not final	part333A	05/01/2019	

**Labeler** - American Consumer Products Corp (081101181)

Revised: 5/2019

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