

**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 65%

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

**Other information**

Other information - store at 20° C (68° to 77° 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**

PHARMACY'S<sup>®</sup>  
PRESCRIPTION

# Hand Sanitizer

Distributed By:  
American Consumer Products Corp  
Vernon, CA 90058

ACP-HSG16M-01

NDC # 72197-028-16

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MADE IN THE USA

PHARMACY'S<sup>®</sup>  
PRESCRIPTION

### Hand Sanitizer

**Drug Facts**

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Moisturizers,  
Vitamin E & Aloe

**MADE IN THE USA**

Net Contents: 16 fl. oz (473 mL)

## PHARMACYS PRESCRIPTION HAND SANITIZER

alcohol gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:72197-028
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	65 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0K00R)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72197-028-16	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/05/2020	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/05/2020	

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

Revised: 5/2020

American Consumer Products Corp