

## **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 Ingredients**

**Active ingredients** - Ethyl Alcohol 71.5%

#### **Purpose**

**Purpose** - Antiseptic

#### **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.**

#### **Indications & Usage**

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

#### **Stop Use**

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

#### **Keep out of reach of children**

**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 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 in skin until dry.

#### **Other Information**

Other information - store at 20° C (68° to 77° F).

-may discolor fabrics.

#### **Inactive Ingredients**

**INACTIVE INGREDIENTS:** Purified Water (Aqua), Isopropanol, Propylene Glycol, Isopropyl Myristate.

**Distributed By:**

**Distributed By:**

American Consumer Products Corp

Vernon, CA 90058

**Pharmacys Prescription Hand Sanitizer**



**Hand Sanitizer**

**Drug Facts**

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**Active ingredients.....Purpose**  
Ethyl Alcohol 71.5% .....Antiseptic

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

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ACP-HSG05M-01



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NDC # 72197-024-05

**PHARMACYS PRESCRIPTION HAND SANITIZER**

alcohol gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:72197-024
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
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ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	71.5 mL in 100 mL
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### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72197-024-05	117 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/17/2020	

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