# PHARMACYS PRESCRIPTION HAND SANITIZER- alcohol liquid 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 Hand Sanitizer**

## **Active Ingredient**

**Active Ingredients:** Ethyl Alcohol 75%

#### **Purpose**

Purpose: Antiseptic

Uses - Reduces bacteria on skin that could cause disease.

#### Warnings

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

Flammable. Keep away from fire or flame.

# 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.** Do not use on children less than 2 months of age. If swallowed, get medical help or contact a Poison Control Center right away.

# **Inactive Ingredients**

**Inactive ingredients -** Glycerin, propylene glycol, triethanolamine, carbomer, fragrance, disodium edta, tocopheryl acetate, deionized water.

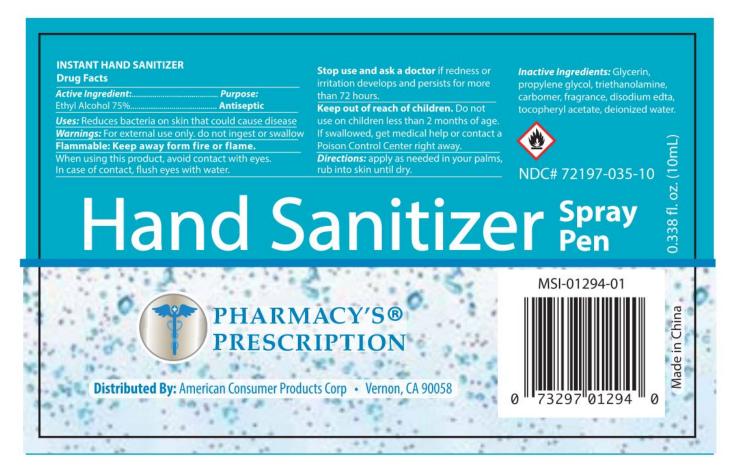
#### **Directions**

**Directions:** apply as needed in your palms, rub into skin until dry.

# **Indications & Usage Section**

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

## **Pharmacy's Prescription Label**



# PHARMACYS PRESCRIPTION HAND SANITIZER

alcohol liquid

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:72197-035 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	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL 2-(2-METHYLBUTYRATE) (UNII: 3Z73C506A9)		

CARBOMER HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)	
TRIETHANOLAMINE 2-CYCLOHEXYL-4,6-DINITROPHENOLATE (UNII: N2TK31JIAH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:72197- 035-10	10 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/15/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/15/2021	

Labeler - American Consumer Products Corp (081101181)

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