# PHARMACYS PRESCRIPTION ALCOHOL WIPES- alcohol swab 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 Wipes**

# **Active Ingredient**

**Active Ingredients** - Ethyl Alcohol 75%

# Purpose

Disinfectant

#### Uses

Helps reduce bacteria on the skin that could cause disease.

# Warnings

**Warnings** - For external use only.

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

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

## When using this product

When using this product, avoid contact with eyes.

In case of contact, flush eyes thoroughly with water.

# Stop use and ask a doctor

**Strop use and ask 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 accidental swallowing. If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

**Directions** - Remove alcohol wipe from package. Thouroughly wipe skin, hands, fingers and wrists. Allow to dry. Discard after single use. Dispose of used alcohol wipes in trash receptacle. Do not flush.

## Other Information

Other information - this is not a baby wipe - store at 20°C (68° to 77°F).

- may discolor fabrics

## **INACTIVE INGREDIENTS**

# **INACTIVE INGREDIENTS:**

Water (Aqua), Aloe Essence Extract, Vitamin E, Phenoxyethanol, Sodium PCA, Benzalkonium Chloride, Sodium Carboxymethl Laurl Glucoside, Ethyl Glycerin, Cocamine DEA, Citric Acid

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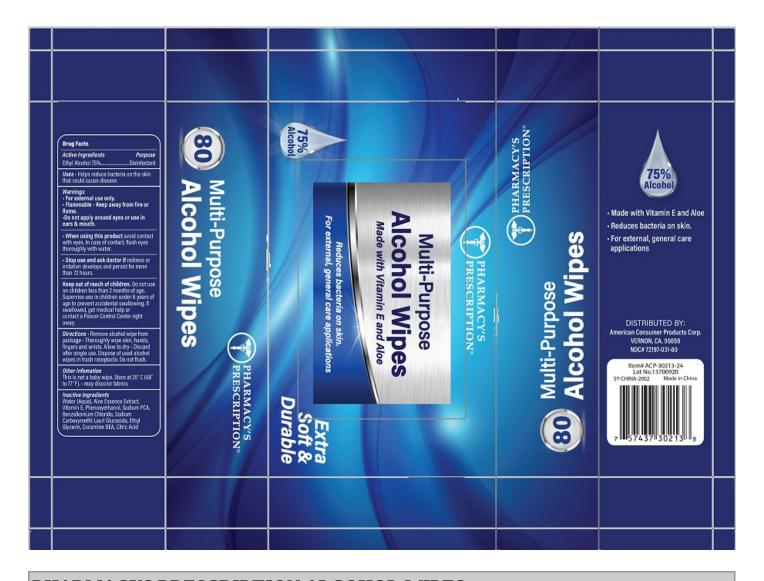
American Consumer Products Corp.

VERNON, CA 90058

Made in China

# **Pharmacys Prescription Alcohol Wipes**





# PHARMACYS PRESCRIPTION ALCOHOL WIPES

alcohol swab

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72197-031
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	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
ETHYL GLYCERIN ACETAL LEVULINATE (UNII: 7233JH4G65)		
COCO DIETHANOLAMIDE (UNII: 92005F972D)		

CITRIC ACID MONO HYDRATE (UNII: 2968 PHW8 QP)	
WATER (UNII: 059QF0KO0R)	
SO DIUM PYRRO LIDO NE CARBO XYLATE (UNII: 469 OTG57A2)	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)	
LAURYL GLUCO SIDE (UNII: 76LN7P7UCU)	

Packaging			
# Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
1 NDC:72197-031-50	50 mL in 1 BAG; Type 0: Not a Combination Product	08/14/2020	
2 NDC:72197-031-80	80 mL in 1 BAG; 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)

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