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.

Pharmacys Prescription Hand Sanitizer

Active Ingredients

Active ingredients - Ethyl Alcool 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 develps 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 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

PHARMACYS PRESCRIPTION HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72197-027

Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	71.5 mL in 100 mL

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72197-027-99	3785.41 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/28/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/28/2020	

Labeler - American Consumer Products Corp (081101181)

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