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

Other information

Other information - store at 20º C (68º to 77º F). May discolor fabrics.

INACTIVE INGREDIENTS: Purified Water (Aqua), Carbomer, Aminomethyl Propanol, Tocopheryl Acetate (Vitamin E), Aloe Barbadensis (Aloe Vera) Leaf Juice

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alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72197-028
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72197-028-99	3785.41 mL in 1 BOTTLE; 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)

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