APEROL HAND SANITIZER- alcohol gel
REACTION RETAIL LLC

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.

Aperol Hand Sanitizer

Drug Facts

Active Ingredient:

Ethyl Alcohol 75%

Antiseptic

Uses:

Hand sanitizer to help decrease bacteria on the skin.

Warnings:

For external use only.

Flammable. Keep away from fire or flame.

Stop use and ask doctor if

irritation or rash appears and lasts.

Keep out of reach of children.

If swallowed, get medical help or contact a doctor right away.

Directions:

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

Other Information:

Store below 118°F

Inactive Ingredients:

Aqua (Water), Glycerin, Propylene Glycol, Aloe Barbadensis Leaf Juice, Parfum (Fragrance), Acrylates/C 10-30 Alkyl Acrylate Crosspolymer, Benzophenone-3, Triethanolamine, Citric Acid, Potassium Sorbate, Sodium Benzoate, Sucrose, Zea Mays (Corn) Starch, Hydroxypropyl Methylcellulose, Polyvinyl Alcohol, Limonene, Hexyl Cinnamal, Linalool, Butylphenyl Methylpropional, Hydroxycitronello, Citronellol, Coumarin, , CI 77267 (D&C Black No.3), CI 16035(FD&C Red No.40), CI 15985 (FD&C Yellow No.6).

Package Labeling:

MADE IN TURKEY
Distributed by Dillard's
1600 Cantrell road little rock, AR 72201
www.Dillards.com

8 10020 29310 4

APEROL
HAND SANITIZER
30ML / 1.0FL.OZ.

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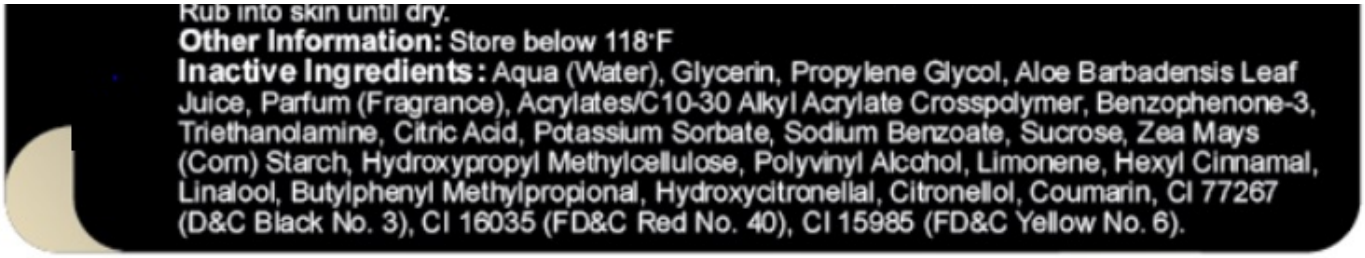

APEROL HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80026-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.75 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
OXYBENZONE (UNII: 95OOS7VE0Y)	
TROLAMINE (UNII: 9O3K93S3TK)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCROSE (UNII: C151H8M554)	
STARCH, CORN (UNII: O8232NY3SJ)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
LIMONENE OXIDE, (+)- (UNII: 278IM94GXB)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
BUTYLPHENYL METHYLPROPIONAL (UNII: T7540GJV69)	
.BETA.-CITRONELLOL, (R)- (UNII: P01OUT964K)	
COUMARIN (UNII: A4VZ22K1WT)	
D&C BLACK NO. 2 (UNII: 4XYU5U00C4)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYDROXYCITRONELLAL (UNII: 8SQ0VA4YUR)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80026-014-30	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/20/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/20/2020	

Labeler - REACTION RETAIL LLC (968085212)

Revised: 10/2020

REACTION RETAIL LLC