

DAYLOGIC ADVANCED HAND SANITIZER REFILL- ethyl alcohol gel

Rite Aid Corporation

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.

Drug Facts

Active ingredient

Ethyl Alcohol 70% (v/v)

Purpose

Antibacterial

Uses

to help reduce bacteria on the skin.

Warnings

For external use only

- Flammable
- Keep away from source of heat or fire

When using this product

avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if

irritation or redness develops and lasts.

Keep out of reach of children.

In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions

- use only to refill a hand sanitizer pump bottle.
- from the pump bottle, put enough product in your palm to cover hands and rub hands together until dry.
- children under 6 years should be supervised when using this product.

Other information

- store at a temperature below 110°F (43°C).
- may discolor certain fabrics or surfaces.

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, Glycerin, Isopropyl Myristate, Aloe Barbadosensis Leaf Juice, Tocopheryl Acetate, Carbomer, Aminomethyl Propanol, Fragrance (Parfum).

Questions or comments?

1-866-695-3030

Label Copy



DAYLOGIC ADVANCED HAND SANITIZER REFILL			
ethyl alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-48 13
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	700 mg in 1 mL
Inactive Ingredients			

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
CARBOMER 934 (UNII: Z135WT9208)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-4813-2	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/27/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/27/2017	

Labeler - Rite Aid Corporation (014578892)

Registrant - Apollo Health and Beauty Care Inc. (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc.		201901209	manufacture(11822-4813)

Revised: 4/2017

Rite Aid Corporation