HDX HAND SANITIZER- ethyl alcohol liquid Apollo Health and Beauty Care

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

Antiseptic

Use

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

- Put enough hand sanitizer 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, Aloe Barbadensis Leaf Juice, Propylene Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance (Parfum), Ethylhexylglycerin, Yellow 5 (CI 19140), Blue 1 (CI 42090).

Label copy





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MADE IN CANADA

DISTRIBUTED BY: HOME DEPOT 2455 PACES FERRY RD., N.W. ATLANTA, GA 30339

1 HOMEDEPOT.COM/HDX





Drug Facts (continued)

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- Effective at eliminating 99.99% of many common harmful germs and bacteria.
- **Purell® is a registered trademark of GOJO Industries, Inc. HDX™ Hand Sanitizer Gel is not manufactured or distributed by GOJO Industries, Inc. 06

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Continued >

HDX HAND SANITIZER

ethyl alcohol liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:63148-479

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)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
GLYCERIN (UNII: PDC6 A3C0 OX)				
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)				
.ALPHATO COPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)				
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)				
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				

]	Packaging						
#	# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:63148-479- 08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/18/2016				
2	NDC:63148-479- 02	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/18/2016				

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333E	08/18/2016					

Labeler - Apollo Health and Beauty Care (201901209)

Registrant - Apollo Health and Beauty Care (201901209)

Establishment						
Name	Address	ID/FEI	Business Operations			
Apollo Health and Beauty Care		201901209	manufacture(63148-479)			

Revised: 8/2016 Apollo Health and Beauty Care