HAND SANITIZER- ethyl alcohol gel Medustry 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.

Active Ingredient

Ethyl Alcohol 75%

Purpose

Antiseptic

Uses

For hand washing to decrease bacteria on the skin.

Recommended for repeated use.

Warnings

Flammable. Keep away from fire or flame.

For external use only.

Avoid contact with broken skin.

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts.

Keep out of reach of children. If swallowed, get medical help.

Directions

Not recommended for infants.

Wet hands thoroughly with product and allow to dry without wiping.

Children under 6 years of age should be supervised when using this product.

Other Information

Do not store above 110°F. May discolor some fabrics.

Inactive Ingredients

Water, Aloe Barbadensis Leaf Juice, Maltodextrin, Glycerin, Propylene Glycol, Acrylates/C1O-30 Alkyl Acrylate crosspolymer, Triethanoamine, Fragrance, Tocopheryl Acetate.



Drug Facts

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Distributed by Medustry LLC 1-866-880-7908 614 Cranbury Road, #273 East Brunswick, NJ 08816

www.medustry.com Made in China

HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type HUMA	N OTC DRUG Item Code	(Source) NDC:74418-208
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Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients Ingredient Name Strength MALTO DEXTRIN (UNII: 7CVR7L4A2D) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) WATER (UNII: 059QF0KO0R) ALOE VERA LEAF (UNII: ZY81Z83H0X) GLYCERIN (UNII: PDC6A3C0OX) CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO) TROLAMINE (UNII: 903K93S3TK)

.ALPHA.-TO COPHEROL ACETATE (UNII: 9E8X80D2L0)

Packaging						
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC •74418 -208-01	236 mL in 1 BOTTLE: Type 0: Not a Combination Product	0.4/24/20.20			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	04/24/2020		

Labeler - Medustry LLC (117306610)

Revised: 4/2020 Medustry LLC