ADORE LAVENDER HAND SANITIZER- alcohol gel K7 Design Group Inc.

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.

Adore Lavender Hand Sanitizer

Drug Facts

Active ingredient

Alcohol 69% v/v

Purpose

Antiseptic

Use

for hand-washing to decrease bacteria on the skin, only when water is not available

Warnings

Flammable, keep away from fire and flames

For external use only

When using this product

- do not get into eyes.
- if contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor if

• irritation and redness develop

Keep out of reach of children.

if swallowed, get medical help or contact a Poison Control Center right away

Directions

wet hands thoroughly with product and allow to dry without wiping

Inactive ingredients

Water, Glycerin, Propylene Glycol, Carbomer, Aloe Barbadensis Leaf Extract, Aminomethyl Propanol, Fragrance, Tocopheryl Acetate, Denatonium Benzoate, Red 33, Blue 1.

Company Information

MANUFACTURED FOR & DISTRIBUTED BY K7 DESIGN GROUP LLC NEW YORK, NY 10016

Product Packaging







LOT: 0Y012021

ORIGIN: CHINA

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ADORE LAVENDER HAND SANITIZER

alcohol gel

Product Information

Inactive Ingredients

Product Type HUMAN OTC DRUG Item Code (Source) NDC:74177-984

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)
ALCOHOL (UNII: 3K9958V90M) ALCOHOL 69 mL in 100 mL

Ingredient Name ALOE VERA LEAF (UNII: ZY81Z83H0 X) AMINOMETHYLPROPANOL (UNII: LU49E6626Q) WATER (UNII: 059QF0KO0R) CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC) FD&C BLUE NO. 1 (UNII: H3R47K3TBD) DENATONIUM BENZO ATE (UNII: 4YK5Z54AT2) GLYCERIN (UNII: PDC6A3C0OX) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) .ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0) D&C RED NO. 33 (UNII: 9DBA0SBB0L)

Packaging					
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:74177-984- 01	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/05/2021		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	01/05/2021		

Labeler - K7 Design Group Inc. (080357784)

Revised: 1/2021 K7 Design Group Inc.