ECOPURE HAND SANITIZER- alcohol gel
Adrian Rivera Maynez Enterprises 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.

ECOPURE HAND SANITIZER

Drug Facts

Active Ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Uses

to help reduce germs and bacteria on the skin

Warnings

For external use only

- Flammable
- Keep away from sources of fire or heat
- Do not ingest

When using this product

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

STOP AND CONSULT YOUR DOCTOR

If irritation or redness occurs

Keep out of Reach of Children.

In case of accidental ingestion contact the your local or national poison control center immediately 1-800-222-1222

Directions

- Wet hands thoroughly with product and rub together vigorously until dry.

Other information

Store at a temperature below 106° degrees F (41° C)

Questions or comments:

888-735-3621

Inactive ingredients:

D-Water, Aloe Vera (Leaf), Isopropyl Alcohol, Glycerin, Propylene Glycol, hydrogen Peroxide, Carbomer, Triethanolamine, Fragrance, FD&C Blue #1 and Yellow #5

Package Labeling:

ECOPURE HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79241-000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	

CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79241-000-00	429 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/01/2020	

Labeler - Adrian Rivera Maynez Enterprises Inc (013690640)

Revised: 8/2020 Adrian Rivera Maynez Enterprises Inc