ESTHETE HAND SANITIZER- ethyl alcohol gel
EQMAXON Corp

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 70.0%

INACTIVE INGREDIENTS

PURIFIED WATER, ALOE EXTRACT, GLYCERIN, SODIUM HYALURONATE, CARBOMER, BUTYLENE GLYCOL, AMINOMETHYL PROPANOL, FRAGRANCE

PURPOSE

ANTISEPTIC

WARNINGS

▪ FOR EXTERNAL USE ONLY ▪ FLAMMABLE, KEEP AWAY FROM FIRE AND FLAME

When using this product
▪Keep out of eyes. In case of contact with eyes, flush thoroughly with water. ▪ Avoid contact with broken skin ▪ Do not inhale or ingest.

Stop use and ask a doctor if irritation or redness develops.

KEEP OUT OF REACH OF CHILDREN

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

Uses

HAND SANITIZER TO HELP REDUCE BACTERIA ON THE SKIN

Directions

▪ DISPENSE ONE TO TWO PUMPS OF SANITIZER ONTO HANDS AND RUB THOROUGHLY UNTIL DRY.

▪ USE AS NEEDED ▪ FOR CHILDREN UNDER 6, USE WITH ADULT SUPERVISION

▪ NOT RECOMMENDED FOR INFANTS

Other Information

▪ Store below 110 °F(43°C)

▪ May discolor some fabrics

▪ Harmful to wood finishes and plastics

Questions or Comments?

Email : gajoany@gmail.com

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

ESTHETE HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55526-0014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	350 mL in 500 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
ALOE (UNII: V5VD430YW9)	
Glycerin (UNII: PDC6A3C0OX)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
Butylene Glycol (UNII: 3XUS85K0RA)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging				
#	Item Code	Package Description	Marketing Start	Marketing End

#	Item Code	Package Description	Date	Date
1	NDC:55526-0014-1	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/01/2020	

Labeler - EQMAXON Corp (557821534)

Registrant - EQMAXON Corp (557821534)

Establishment

Name	Address	ID/FEI	Business Operations
EQMAXON Corp		557821534	manufacture(55526-0014)

Revised: 5/2020 EQMAXON Corp