DR.S CLEAN ADVANCED 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, Triethanolamine, Flavoring

PURPOSE

ANTISEPTIC

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 or redness develops.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Uses

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

Directions

Wet hands thoroughly with product and allow to dry without wiping

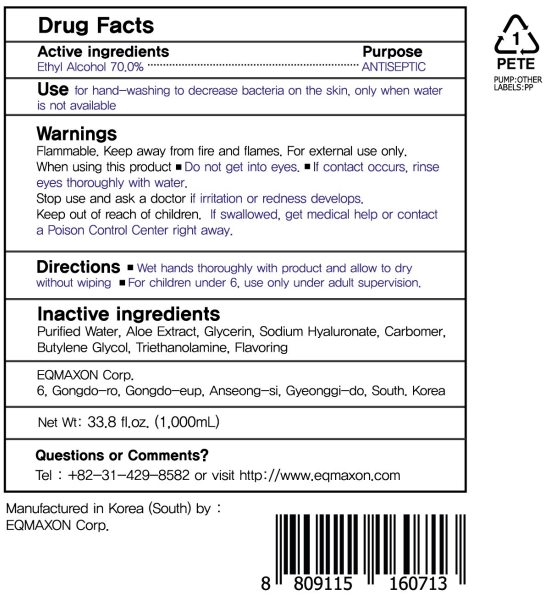
For children under 6, use only under adult supervision.

Questions

Questions?

+82-31-429-8582 or visit http://www.eqmaxon.com

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



DR.S CLEAN ADVANCED HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55526-0017
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	700 mL in 1000 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)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55526-0017-1	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/01/2021	

Labeler - EQMAXON Corp. (557821534)

Registrant - EQMAXON Corp. (557821534)

Establishment

Name	Address	ID/FEI	Business Operations
EQMAXON Corp.		557821534	manufacture(55526-0017)

Revised: 6/2021 EQMAXON Corp.