

DR.S CLEAN HAND- 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

Active ingredients: Ethyl Alcohol 70.0%

INACTIVE INGREDIENT

Inactive ingredients:

Purified Water, Aloe Extract, Glycerin, Sodium Hyaluronate, Carbomer, Butylene Glycol, Triethanolamine, Flavor

PURPOSE

Purpose: ANTISEPTIC

WARNINGS

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

Uses:

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

Directions

Directions:

Wet hands thoroughly with product and allow to dry without wiping

For children under 6, use only under adult supervision.

PACKAGE LABEL - Dr.'s Clean Hand Gel 60mL



Dr.'S CLEAN HAND GEL

Simply KILL 99.9 % Germ

HAND SANITIZER

Leaves Hands Feeling Soft
ADVANCED

**70% alcohol
(ethanol) contented**
2.02 FL.OZ. (60ml)

Drug Facts	
Active ingredients Ethyl Alcohol 70.0%	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 or redness develops. 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 • For children under 6, use only under adult supervision.	
Inactive ingredients Purified Water, Aloe Extract, Glycerin, Sodium Hyaluronate, Carbomer, Butylene Glycol, Triethanolamine, Flavoring	
EQMAXON Corp. 6, Gongdo-ro, Gongdo-eup, Anseong-si, Gyeonggi-do, S. Korea	
2.02 FL.OZ. (60ml)	
Questions? +82-31-429-8582 or visit http://www.eqmaxon.com	

Manufactured in Korea (South) by
EQMAXON Corp.



PACKAGE LABEL - Dr.'s Clean Hand Gel 500mL



Dr.'S CLEAN HAND GEL

Simply KILL 99.9 % Germ

HAND SANITIZER



Leaves Hands Feeling Soft
ADVANCED

**70% alcohol
(ethanol) contented**
16.90 FL.OZ. (500ml)

Drug Facts	
Active ingredients Ethyl Alcohol 70.0%	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 or redness develops. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
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DR.S CLEAN HAND

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55526-00 10
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
ALOE (UNII: V5VD430YW9)	
Glycerin (UNII: PDC6A3C0OX)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
Butylene Glycol (UNII: 3XUS85K0RA)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55526-0010-1	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2020	
2	NDC:55526-0010-2	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/01/2020	

Labeler - EQMAXON Corp (557821534)

Registrant - EQMAXON Corp (557821534)

Establishment

Name	Address	ID/FEI	Business Operations
EQMAXON Corp		557821534	manufacture(55526-0010)