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.PRINCIPAL DISPLAY PANEL

DR.S CLEAN HAND

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55526-0011
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 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:55526-0011-1	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph not final	part333E	04/01/2020	

Labeler - EQMAXON Corp (557821534)

Registrant - EQMAXON Corp (557821534)

Establishment

Name	Address	ID/FEI	Business Operations
EQMAXON Corp		557821534	manufacture(55526-0011)

Revised: 4/2020 EQMAXON Corp