DR. NATURE HUB- alcohol gel SHIMIZU CO., LTD.

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

Carbomer, Glycerin, Triethanolamine, Allantoin, Dexpanthenol, Dipotassium Glycyrrhizate, Lavandula Angustifolia (Lavender) Extract, Panax Ginseng Root Extract, Orange Oil, Limonene, Water

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 if irritation or redness develops

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

■ Hand sanitizer to help reduce bacteria that potentially can cause disease.

Directions

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

Active ingredient Ethyl Alcohol 70.0%	Purpose Antiseptic
Uses Hand sanitizer to help reduce be cause disease.	acteria that potentially can
Flammable. Keep away from fire a For external use only. When using this product • Do no • If contact occurs, rinse eyes thorous Stop use and ask a doctor if if irr	t get into eyes. July with water.
Keep out of reach of children If so or contact a Poison Control Center righ	wallowed, get medical help
Directions • Place enough product of surfaces. Rub hands together until dry	





ShimizU Co., Ltd. 12,Teheran-ro 63-gil, Gangnam-gu, Seoul, 06160 Republic of Korea





MADE IN KOREA

DR. NATURE HUB

alcohol gel

Net Wt.: 16.90 fl.oz. (500ml)

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:76901-010

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

Ingredient Name

Strength

WATER (UNII: 059QF0KO0R)

CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)

Glycerin (UNII: PDC6A3C0OX)

TROLAMINE (UNII: 9O3K93S3TK)

Allantoin (UNII: 344S277G0Z)

Dexpanthenol (UNII: 106C93RI7Z)

GLYCYRRHIZINATE DIPOTASSIUM (UNII: CA2Y0FE3FX)

LAVENDER OIL (UNII: ZBP1YXW0H8)

ASIAN GINSENG (UNII: CUQ3A77YXI)

Orange Oil (UNII: AKN3KSD11B)

LIMONENE, (+)- (UNII: GFD7C86Q1W)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76901-010- 01	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 - SHIMIZU CO., LTD. (694646122)

Registrant - SHIMIZU CO., LTD. (694646122)

Establishment							
Name	Address	ID/FEI	Business Operations				
Korea Miracle People Corporation Co.,Ltd		693710617	manufacture(76901-010)				

Revised: 5/2020 SHIMIZU CO., LTD.