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

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 if irritation or redness develops

KEEP OUT OF REACH OF CHILDREN

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

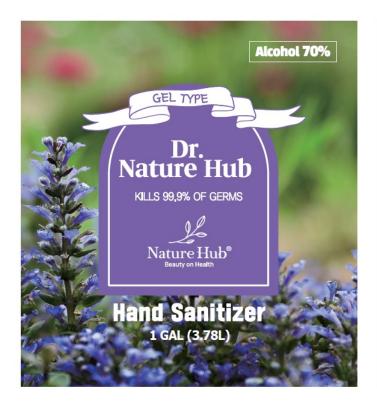
Uses

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

Directions

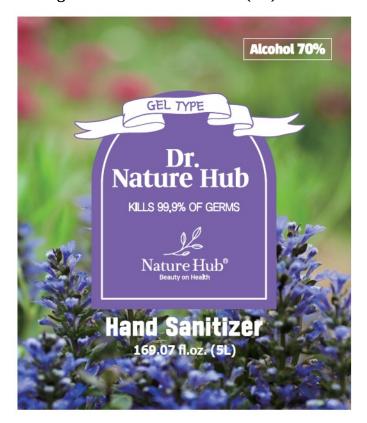
- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Package Label: Dr. Nature Hub (1gal)





Package Label: Dr. Nature Hub (5L)





DR. NATURE HUB

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76901-020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Alcohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	Alcohol	0.7 L in 1 L	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)		
Glycerin (UNII: PDC6A3C0OX)		
TROLAMINE (UNII: 903K93S3TK)		
Allantoin (UNII: 344S277G0Z)		
Dexpanthenol (UNII: 106C93RI7Z)		
GLYCYRRHIZINATE DIPO TASSIUM (UNII: CA2Y0 FE3FX)		
LAVENDER OIL (UNII: ZBP1YXW0H8)		
ASIAN GINSENG (UNII: CUQ3A77YXI)		
Orange Oil (UNII: AKN3KSD11B)		
LIMO NENE, (+)- (UNII: GFD7C86Q1W)		

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:76901-020- 01	3.78 L in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2020		
2	NDC:76901-020- 02	5 L 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(76901-020)	

Revised: 6/2020 SHIMIZU CO., LTD.