DR.GERM HAND SANITIZER GEL- hand sanitizer gel
Oneskin Cosmetics 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.

Flammable. Keep away from fire or flame.

For external use only

When using this product do not use in or near the eyes.

In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts.

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

Ethyl Alcohol 70%

Antimicrobial

- Hand sanitizer to help reduce bacteria on the skin.

Water, Glycerin, Carbomer, Tetrahydroxypropyl Ethylenediamine, Orange Oil, Peppermint Oil, Sodium Hyaluronate, Allantoin, Mugwort Extract, Green Tea Extract.

Put enough product in your palm to cover hands and rub hands together briskly until dry.

Childern under 6 years of age should be supervised when using Dr. germ Hand Sanitizer.

Store below 110oF (43oC).
May discolor certain fabrics or surfaces.

Dr.germ Hand Sanitizer Gel

DR.GERM HAND SANITIZER GEL

hand sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70889-500
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
EDETOL (UNII: Q4R969U9FR)	
ORANGE OIL (UNII: AKN3KSD11B)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
ALLANTOIN (UNII: 344S277G0Z)	
ARTEMISIA PRINCEPS LEAF (UNII: SY077EW02G)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color		**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70889-500-01	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/19/2020	
2	NDC:70889-500-02	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/19/2020	
3	NDC:70889-500-03	1 in 1 BOX	03/19/2020	
3		50 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/19/2020	

Labeler - Oneskin Cosmetics Co., Ltd. (689846630)

Registrant - Oneskin Cosmetics Co., Ltd. (689846630)

Establishment

Name	Address	ID/FEI	Business Operations
Oneskin Cosmetics Co., Ltd.		689846630	manufacture(70889-500)

Revised: 4/2020

Oneskin Cosmetics Co., Ltd.