# DR. SAM PREMIUM HAND SANITIZER (500ML)- ethyl alcohol gel G.E.O MARKETING 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.

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### Dr. Sam Premium Hand Sanitizer (500mL)

### **Active ingedient**

Ethanol (ethyl alcohol) (62%)

#### **Purpose**

Antibacterial

#### Uses

Hand sanitizer for decreasing bacteria on skin

### Warnings

#### For external use only

Do not use if you are allergic to any of the ingredients

When using this product avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water. avoid contact with broken, irritated, or itching skin. Do not puncture or incinerate.

Stop use and ask a doctor if irritation or redness develops and condition persists for more than 72 hours.

If pregnant or breast-feeding, ask a health professional before use

### Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

#### **Directions**

Apply the adequate amount on hands

### **Inactive ingredients**

Purified water, Glycerin, L-menthol, Carbomer, Aminomethyl Propanol, Aloe extract

#### Dr. Sam Premium Hand Sanitizer (500mL)



Kills 99.9% of germs

Hand disinfectant for infection prevention

500 ml/16.9 fl. oz.



Contains moisturizing ingredients Moisturizing effect considering skin health Available outdoor and indoor anytime, anywhere Various diseases prevention and sterilization effect Safer with hand sanitizer Check of healthy ingredients

#### Drug Facts

wnen soap an **Warnings** 

al use only. Flammable. Keep away from heat or fla

Do not use

if you are allergic to any of the ingredients

in children less than 2 months of age

on open skin wounds

When using this product avoid contact with eyes, ears and mouth. In case of contact

Strength

## DR. SAM PREMIUM HAND SANITIZER (500ML)

ethyl alcohol gel

#### **Product Information**

HUMAN OTC DRUG Item Code (Source) NDC:78012-101 Product Type

TOPICAL **Route of Administration** 

#### **Active Ingredient/Active Moiety**

**Ingredient Name Basis of Strength** Strength 62 g in 100 mLALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL

### **Inactive Ingredients Ingredient Name**

GLYCERIN (UNII: PDC6A3C0OX)

LEVOMENTHOL (UNII: BZ1R15MTK7) AMINOMETHYLPROPANOL (UNII: LU49E6626Q)

ALOE (UNII: V5VD430YW9)

WATER (UNII: 059QF0KO0R)

#### **Packaging**

ı	# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ı	1 NDC:78012-101-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/25/2020	

### **Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/25/2020	

# Labeler - G.E.O MARKETING CORP (080740733)

Establishment									
Name	Address	ID/FEI	Business Operations						
Korea Medicare Co.,Ltd.		695832135	manufacture (78012-101)						

Revised: 6/2020 G.E.O MARKETING CORP