DAENG GI MEO RI KI GOLD PREMIUM HAND SANITIZER GEL- alcohol gel
Doori 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.

DAENG GI MEO RI KI GOLD PREMIUM HAND SANITIZER GEL

Active ingredient

Ethyl alcohol 70%

Purpose

Antimicrobial

Use

- Hand sanitizer to help reduce bacteria on the skin

Warnings

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 Poison Control Center right away.

Directions

- Put nough produt in your palm to cover hands and rub hands together briskly until dry.
- Children under 6 years of age should be supervised when using this product

Other information

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces

Inactive ingredients

Water, Butylene Glycol, Carbomer, Aminomethyl Propanol, Fragrance, Glycerin, Paeonia Suffruticosa Root Extract, Camellia Sinensis Leaf Extract, Aloe Ferox Leaf Extract

DAENG GI MEO RI KI GOLD PREMIUM HAND SANITIZER GEL

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50375-001
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
GLYCERIN (UNII: PDC6A3C0OX)	
PAEONIA X SUFFRUTICOSA ROOT (UNII: 7M7E9A2C8J)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
ALOE FEROX LEAF (UNII: 0D145J8EME)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50375-001-01	50 mL in 1 TUBE; Type 0: Not a Combination Product	04/21/2020	
2	NDC:50375-001-02	250 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/21/2020	
3	NDC:50375-001-03	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/21/2020	
4	NDC:50375-001-04	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/21/2020	
5	NDC:50375-001-05	3800 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/21/2020	

Labeler - Doori Cosmetics Co., Ltd. (688227465)

Registrant - Doori Cosmetics Co., Ltd. (688227465)

Establishment

Name	Address	ID/FEI	Business Operations
Doori Cosmetics Co., Ltd.		688227465	manufacture(50375-001)

Revised: 4/2020 Doori Cosmetics Co., Ltd.