

**HANNA HAND SANITIZER GEL- alcohol gel**  
**Rainbow Beauty Cosmetic 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.*

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**Drug Facts**

alcohol

Water, Glycerin, Carbomer, Triethanolamine, Fragrance, Camellia Sinensis Leaf Extract, Butylene Glycol, Aloe Barbadensis Leaf Extract, Portulaca Oleracea Extract, Centella Asiatica Extract, Pinus Densiflora Extract, CI 19140, CI 42090

Sterilization of hands and skin

**KEEP OUT OF REACH OF THE CHILDREN**

Apply an appropriate amount on your hands and rub well to dry.

1. Do not use on the following body parts. A wide range of body parts and damaged skin around the eyes and ears, in the oral cavity (may have irritating effects)
2. If the following symptoms appear, stop using them immediately and consult a doctor or pharmacist.
  - 1) Hypersensitivity symptoms such as rash, erythema, itching, and edema
  - 2) Skin irritation symptoms
3. Other precautions
  - 1) For external use only (do not underwear).
  - 2) Be careful not to get into your eyes, and if so, rinse well with clean water and consult a doctor or pharmacist.
  - 3) Be careful not to inhale the vapor when using it extensively or for a long period of time (irritation to the mucous membranes, headaches, etc. may occur if ethanol vapor is consumed in large quantities or repeatedly).
  - 4) If repeated use on the same site, be careful as the skin may become rough due to degreasing.
  - 5) Do not use sealed bandages, cast bandages, packs, etc., as irritation may occur.
  - 6) Do not use this medicine for anal or vaginal compresses as it may cause irritation or chemical burns.
  - 7) Do not use for any other purpose.
4. Precautions for storage
  - 1) Avoid shading and keep in shading.
  - 2) Keep it out of reach of children, and if a child swallows it, go to the hospital right away.
  - 3) After use, close the product completely with a lid to prevent the product from drying out or entering foreign Objects.

for external use only

**HANNA**  
HAND SANITIZER  
REFRESHING GEL  
ETHYL ALCOHOL 70%

10.14 fl.oz  
300ml

Locally Owned Company  
HANNA ENTERPRISES LLC

Reduce More Than  
**99.99%**  
of Germs

Manufacturer: Hanhwa Beauty Cosmetic Co., Ltd.  
21 Seoncheon-ro, Seongbuk-gu, Gyeonggi-do, Seoul, Korea, South Korea  
Distributed by: Hanna Enterprises LLC  
103 Bloor West Street, Stamford, Connecticut 06903

MADE IN KOREA

8 809317 968932

**HANNA HAND SANITIZER GEL**

**Drug Facts**

| Active Ingredients | Purpose       |
|--------------------|---------------|
| Ethyl alcohol 70%  | Antimicrobial |

**Use**  
Hand sanitizer to help reduce bacteria on skin & surface

**Warning**  
**Flammable. Keep away from fire or flame.**  
**For external use only**  
When using this product avoid contact with eyes. If contact occurs, 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**  
• Place enough product in your palm to thoroughly cover your hands  
• Rub hands together briskly until dry  
• No rinsing required • No towels needed.

**Other information**  
• Store at 110°F (43°C) • May discolor certain fabrics or surfaces.

**Inactive ingredients**  
Alcohol (70.00%), Water (28.129889%), Glycerin (1.00%), Carbomer (0.40%), Triethanolamine (0.369%), Fragrance (0.05%), Camellia Sinensis Leaf Extract (0.01%), Butylene Glycol (0.01%), Aloe Barbadensis Leaf Extract (0.01%), Portulaca Oleracea Extract (0.01%), Centella Asiatica Extract (0.01%), Pinus densiflora Extract (0.01%), CI 19140 (0.0001%), CI 42090 (0.0002%)

**Questions** hannaenterkorea@gmail.com

**HANNA**  
HAND SANITIZER  
REFRESHING GEL  
ETHYL ALCOHOL 70%

16.90 fl.oz  
500ml

Locally Owned Company  
HANNA ENTERPRISES LLC

Reduce More Than  
**99.99%**  
of Germs

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## HANNA HAND SANITIZER GEL

alcohol gel

### Product Information

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:71193-101 |
| <b>Route of Administration</b> | TOPICAL        |                           |               |

### Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength | Strength        |
|--|-------------------|-----------------|
| ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M) | ALCOHOL           | 70 mL in 100 mL |

## Inactive Ingredients

| Ingredient Name                          | Strength |
|--|----------|
| GREEN TEA LEAF (UNII: W2ZU1RY8B0)        |          |
| BUTYLENE GLYCOL (UNII: 3XUS85K0RA)       |          |
| PURSLANE (UNII: M6S840WXG5)              |          |
| TROLAMINE (UNII: 9O3K93S3TK)             |          |
| PINUS DENSIFLORA BARK (UNII: U68X322T49) |          |
| FD&C YELLOW NO. 5 (UNII: I753WB2F1M)     |          |
| ALOE VERA LEAF (UNII: ZY81Z83H0X)        |          |
| CENTELLA ASIATICA (UNII: 7M867G6T1U)     |          |
| GLYCERIN (UNII: PDC6A3C0OX)              |          |
| HYALURONATE SODIUM (UNII: YSE9PPT4TH)    |          |
| CARBOMER 940 (UNII: 4Q93RCW27E)          |          |
| WATER (UNII: 059QF0K00R)                 |          |
| D&C BLUE NO. 4 (UNII: 0KSY80VYS3)        |          |

## Packaging

| # | Item Code        | Package Description                                   | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:71193-101-01 | 500 mL in 1 BOTTLE; Type 0: Not a Combination Product | 09/03/2020           |                    |
| 2 | NDC:71193-101-02 | 300 mL in 1 BOTTLE; Type 0: Not a Combination Product | 09/03/2020           |                    |

## Marketing Information

| Marketing Category      | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part333A                                 | 09/03/2020           |                    |

**Labeler** - Rainbow Beauty Cosmetic Co., Ltd. (695684820)

**Registrant** - Rainbow Beauty Cosmetic Co., Ltd. (695684820)

## Establishment

| Name                              | Address | ID/FEI    | Business Operations                       |
|-----------------------------------|---------|-----------|---|
| Rainbow Beauty Cosmetic Co., Ltd. |         | 695684820 | manufacture(71193-101) , label(71193-101) |

Revised: 9/2020

Rainbow Beauty Cosmetic Co., Ltd.