# HEMIRUS HAND CLINIC GEL- alcohol gel GC US CORPORATION

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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# Active ingredients

Alcohol 62%

Antiseptic

#### Uses

- hand sanitizer to help decrease bacteria on the skin.
- when water, soap & tower are not available
- recommended for repeated use

## Warnings

Flammable. Keep away from fire or flame

For external use only.

Do not apply around eyes. Do not use

in ears & mouth

# When using this product

avoid contact with eyes.

In case of contact flush eyes with water

### Stop use and ask a doctor if

redness or irritation develop and persist for more than 72 hours.

#### Keep out of reach of children.

Children must be supervised in use of this product.

#### Directions

- pump as needed into your palms and throughly spread on both hands.
- rub into skin until dry.

#### **Other Information**

- Store at 20 °C (68 ° to 77 °F).
- may discolor fabrics.

#### **Inactive Ingredients**

Water, Phenoxyethanol, Trolamine, Carbomer homopolymer type C (Allyl pentaerythritol crosslinked) Methylparaben, Orange, Isopropyl myristate, D&C Orange No. 4

#### Questions?

• side effects occur. You may report side effects to FDA at 1-800-FDA-1088.

**Instant Hand Sanitizer!** 

Kills 99.9% of Most Common Germs!

Net Wt. 16.91 fl oz. (500ml)

Manufactured by Woosincosmetics Inc.

33, Sudo-ro, 125beon-gil, Bucheon-si, Gyeonggi-do, Korea 14491

500mL NDC: 71454-3039-1



KFDA Approved

# неmirus Hand Clinic Gel



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#### **Drug Facts**

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Purpose Antiseption

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MADE IN KOREA



#### HEMIRUS HAND CLINIC GEL

alcohol gel

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71454-3039

Route of Administration TOPICAL

## Active Ingredient/Active Moiety

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Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

<b>Inactive Ingredients</b>
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<b>Ingredient Nam</b>	e
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Strength

WATER (UNII: 059QF0KO0R)

PHENO XYETHANOL (UNII: HIE492ZZ3T)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
ORANGE (UNII: 5EVU04N5QU)	
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)	
D&C ORANGE NO. 4 (UNII: Q1LIY3BO0U)	

l	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:71454-3039-	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/31/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	03/31/2020		

# Labeler - GC US CORPORATION (080669481)

Establishment			
Name	Address	ID/FEI	Business Operations
WOOSIN COSMETICS CO.,LTD		688227829	manufacture(71454-3039)

Revised: 3/2020 GC US CORPORATION