

**MEDIPLUS HAND SANITIZER- benzalkonium chloride liquid**  
**GL Pharm 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.*

-----

**Drug Facts**

benzalkonium chloride 0.66%

polyoxyethylene lauryl ether, ethanol, purified water

hand sanitizer

**KEEP OUT OF REACH OF THE CHILDREN**

Apply to clean, dry hands. Apply sufficient amount to thoroughly wet all surfaces of hands and fingers. Rub onto hands until dry.

Supervise children in the use of this product.

- Flammable. Keep away from fire or flame.
- For external use only.
- Do not use in eyes.
- If swallowed, get medical help promptly.
- Stop use, ask doctor If irritation occurs.
- Keep out of reach of children.

for external use only

**MEDIPLUS Hand Sanitizer**  
- Drug Facts -

**Active Ingredient:** Benzalkonium chloride 0.66% **Purpose:** Antiseptic

**Uses:** Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

**Warnings:**  
For external use only.  
Do not use 1) in children less than 2 months of age, 2) on open skin wounds.  
When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.  
Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.  
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.


**Directions:**  
1) Place enough product on hands to cover all surfaces. Rub hands together until dry. 2) Supervise children under 6 years of age when using this product to avoid swallowing.

**Other Information:** 1) Store between 1-30C (33.8-86F) 2) Avoid freezing and excessive heat above 40C (104F)

**Inactive Ingredients:** Polyoxyethylene lauryl ether, Ethanol, Purified water

**Manufacturer:** GL Pharm Co., Ltd 130, Hansan-ro, Deoksan-eup, Jincheon-gun, Chungcheongbuk-do, Korea

Net 10.14 fl. oz. TEL. 82.2.556.1793



**MEDIPLUS Hand Sanitizer**  
- Drug Facts -

**Active Ingredient:** Benzalkonium chloride 0.66% **Purpose:** Antiseptic

**Uses:** Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

**Warnings:**  
For external use only.  
Do not use 1) in children less than 2 months of age, 2) on open skin wounds.  
When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.  
Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.  
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.


**Directions:**  
1) Place enough product on hands to cover all surfaces. Rub hands together until dry. 2) Supervise children under 6 years of age when using this product to avoid swallowing.

**Other Information:** 1) Store between 1-30C (33.8-86F) 2) Avoid freezing and excessive heat above 40C (104F)

**Inactive Ingredients:** Polyoxyethylene lauryl ether, Ethanol, Purified water

**Manufacturer:** GL Pharm Co., Ltd 130, Hansan-ro, Deoksan-eup, Jincheon-gun, Chungcheongbuk-do, Korea

Net 3.38 fl. oz. TEL. 82.2.556.1793



## MEDIPLUS HAND SANITIZER

benzalkonium chloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:74713-0001
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.66 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74713-0001-1	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020	
2	NDC:74713-0001-2	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2020	

**Labeler** - GL Pharm Co Ltd (694506955)

**Registrant** - GL Pharm Co Ltd (694506955)

## Establishment

Name	Address	ID/FEI	Business Operations
GL Pharm Co Ltd		694506955	manufacture(74713-0001)

Revised: 4/2020

GL Pharm Co Ltd