MULTI PURPOSE STERILIZATION HAND SANITIZER- alcohol spray
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.

ACTIVE INGREDIENT

Active ingredients: Ethyl alcohol 83.0% v/v

INACTIVE INGREDIENT

Inactive ingredients:

Water

PURPOSE

Purpose: ANTISEPTIC

WARNINGS

Warnings:

For external use only-hands

Flammable. Keep away from heat and flame.

When using this product ■ Keep out of eyes. In case of contact with eyes, flush thoroughly with water. ■ Avoid contact with broken skin. ■ Do not inhale or ingest.

Stop use and ask a doctor if skin irritation or rash develops.

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Uses

Uses:

■ Hand sanitizer to decrease bacteria on the skin that could cause disease
■ Disinfection of the skin at the surgical site

Directions

Directions:

■ Spray the product on skin or other place where disinfection is required.
■ Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

Other information
• Store between 15-30C (59-86F)
• Avoid freezing and excessive heat above 40C (104F)

Questions

Questions
http://www.glpharm.co.kr or TEL : +82-43-537-7001

Package Label : Multi-purpose sterilization hand sanitizer 100mL

Package Label : Multi-purpose sterilization hand sanitizer 300mL

Package Label : Multi-purpose sterilization hand sanitizer 250mL

MULTI PURPOSE STERILIZATION HAND SANITIZER

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74713-0510
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	83 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:74713-0510-1	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	
2	NDC:74713-0510-2	300 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	
3	NDC:74713-0510-3	250 mL in 1 BOTTLE, SPRAY; 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	part333E	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(74713-0510)

Revised: 5/2020 GL Pharm. Co., Ltd.