G SOL SANITIZING- silver spray
GP&E Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

ACTIVE INGREDIENT

Silver 0.005%

INACTIVE INGREDIENTS

Water, Titanium Dioxide, Citric Acid, Polysorbate 20

PURPOSE

Sanitizer

WARNINGS

For external use only.

Do not use
• in children less than 2 months of age
• 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.

KEEP OUT OF REACH OF CHILDREN

If swallowed, get medical help or contact a Poison Control Center right away.

Use(s)

Hand sanitizer to help reduce bacteria that potentially can cause disease.

Directions

• Spray enough product on hands to cover all surfaces. Rub hands together until dry.
• Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

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

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

G SOL SANITIZING

silver spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81042-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

Silver (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)	Silver	0.005 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Titanium Dioxide (UNII: 15FIX9V2JP)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
Polysorbate 20 (UNII: 7T1F30V5YH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81042-010-01	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/01/2020	

Labeler - GP&E Co., Ltd. (689816478)

Registrant - GP&E Co., Ltd. (689816478)

Establishment

Name	Address	ID/FEI	Business Operations
GP&E Co., Ltd.		689816478	manufacture(81042-010)

Revised: 9/2022 GP&E Co., Ltd.