ZEROQ- potassium carbonate liquid
JISANG 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.

---

potassium carbonate

e-polylysine, dextrin, 1,2-propanediol, citric acid, tea catechin, hydroxypropylmethyl cellulose, water

Sanitizer to help decrease bacteria on skin

keep out of reach of the children

Spray on skin as needed

For external use only.

When using this product

Avoid contact with eyes. Discontinue use if signs of irritation or rashes appear.

Keep out of reach of children.

Store at room temperature.

for external use only

Drug Facts

Active Ingredient — Purpose

Potassium carbonate (0.1%) ---------- antibacterial

Uses

Sanitizer to help decrease bacteria on skin.

Warnings

For external use only

When using this product

Avoid contact with eyes. Discontinue use if signs of irritation or rashes appear.

Keep out of reach of children.

Store at room temperature

Directions

Spray on skin as needed.

Inactive Ingredients

ε-Polylysine.Dextrin.1,2-Propanediol.Citric Acid.Tea Catechin. Hydroxypropylmethyl Cellulose. Distilled water.

Manufactured by

JISANG Co,.Ltd.
903-4, Manghyang-ro, Seonggeo-eup, Seobuk-gu, Cheonan-si, Chungcheongnam-do,Republic of Korea

Question or Comments

Call +82-41-622-0080

ZEROQ

potassium carbonate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80643-0002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POTASSIUM CARBONATE (UNII: BQN1B9B9HA) (CARBONATE ION - UNII:7UJQ5OPE7D)	POTASSIUM CARBONATE	0.5 g in 500 mL

Inactive Ingredients	
Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80643-0002-1	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/21/2020	

Labeler - JISANG Co., Ltd. (694802124)

Registrant - JISANG Co., Ltd. (694802124)

Establishment

Name	Address	ID/FEI	Business Operations
JISANG Co., Ltd.		694802124	manufacture(80643-0002)

Revised: 9/2020 JISANG Co., Ltd.