ZEROQ FACE BODY- 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

Distributed by LULU Co.,Ltd.

505-28, Seocho-daero 335-gil, Seocho-gu, Seoul, Republic of Korea

Question or Comments

Call +82-2-6207-7888

ZEROQ FACE BODY

potassium carbonate liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:80643-0003

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POTASSIUM CARBONATE (UNII: BQN1B9B9HA) (CARBONATE ION - UNII:7UJQ50PE7D)	POTASSIUM CARBONATE	0.1 g in 100 mL
ONIL/OJQ3OFE/D)	CARDUNATE	III 100 IIIL

Inactive Ingredients			
Ingredient Name	Strength		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
CITRIC ACID MONO HYDRATE (UNII: 2968 PHW8 QP)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
WATER (UNII: 059QF0KO0R)			

Packaging				
# Item Code Package Description		Marketing Start Date	Marketing End Date	
1 NDC:80643- 0003-1	100 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(80643-0003)

Revised: 9/2020 JISANG Co., Ltd.