

PURILAB- citric acid spray
Seoul Food Research & Development 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

Citric acid 0.026% v/v

INACTIVE INGREDIENTS

Ascorbic acid, Citrus extract, Citric extractives, DL-Malic acid, Ethanol, Lactic acid, Saccharum officinarum extract, Water

PURPOSE

Antimicrobial

WARNINGS

For external use only

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

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

- Disinfectant to help reduce bacteria on the skin and surfaces
- Recommended for repeated use

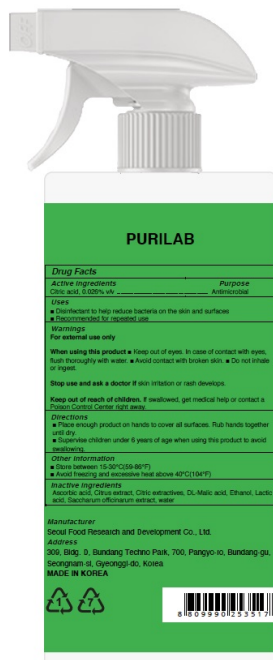
Directions

- Place 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 15-30°(59-86°)
- Avoid freezing and excessive heat above 40°(104°)

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



PURILAB

citric acid spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80642-060
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP) (ANHYDROUS CITRIC ACID - UNII:XF417D3PSL)	ANHYDROUS CITRIC ACID	0.13 g in 500 mL

Inactive Ingredients

Ingredient Name	Strength
Ascorbic acid (UNII: PQ6CK8PD0R)	
MALIC ACID (UNII: 817L1N4CKP)	
ALCOHOL (UNII: 3K9958V90M)	
Lactic acid (UNII: 33X04XA5AT)	
SACCHARUM OFFICINARUM WHOLE (UNII: 3Z20C92XNB)	
Water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80642-060-01	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/01/2020	

Labeler - Seoul Food Research & Development Co., Ltd. (688787758)

Registrant - Seoul Food Research & Development Co., Ltd. (688787758)

Establishment

Name	Address	ID/FEI	Business Operations
Seoul Food Research & Development Co., Ltd.		688787758	manufacture(80642-060)

Revised: 10/2020

Seoul Food Research & Development Co., Ltd.