

ZEROTH PREMIUM HAND SANITIZER- alcohol gel

Biocleansingcompany 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

ALCOHOL 70% v/v

INACTIVE INGREDIENTS

Water, Carbomer, Tetrahydroxypropyl Ethylenediamine, Tocopheryl Acetate, Lavandula Angustifolia (Lavender) Oil, Fragaria Chiloensis (Strawberry) Fruit Extract, Sophora Angustifolia Root Extract, Morus Alba Bark Extract, Glycyrrhiza Uralensis (Licorice) Root Extract, Citrus Paradisi (Grapefruit) Fruit Extract, Camellia Sinensis Leaf Extract, Cnidium Officinatum Rhizome Extract

Purpose

Hand Sanitizer

Warnings

For external use only. Flammable. Keep away from heat or flame

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.

Uses

- instant hand antiseptic to decrease bacteria on the skin

Directions


- Apply to clean, dry hands. Apply sufficient amount to thoroughly wet all surfaces of hands and fingers. Rub onto hands until dry.
- Supervise children in the use of this product.

Other information

- Store between 1-30C (39-86F)

- Avoid freezing and excessive heat above 40C (104F)

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



제로스
500 ML / 16.91 FL OZ
MADE IN KOREA

Kills 99.9% of Germs

ZEROth

**PREMIUM
HAND
SANITIZER**

ZEROth

ETHANOL 70%
KFDA Approved
FDA OTC Registered

Drug Facts

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Distributor: Biocleansingcompany Co., Ltd
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Manufacturer: Dongbang Cosmetics Co., Ltd.
39 Gajaeul-ro 32beon-gil, Seo-gu, Incheon, S. Korea

ZEROth PREMIUM HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78393-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	350 mL in 500 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
EDETOL (UNII: Q4R969U9FR)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
BEACH STRAWBERRY (UNII: 98542F2TQS)	
SOPHORA FLAVESCENS ROOT (UNII: IYR6K8KQ5K)	
MORUS ALBA BARK (UNII: 7O71A48NDP)	
GLYCYRRHIZA URALENSIS ROOT (UNII: 42B5YD8F0K)	
GRAPEFRUIT (UNII: O82C39RR8C)	

GREEN TEA LEAF (UNII: W2ZU1RY8B0)

CNIDIUM OFFICINALE ROOT (UNII: 8S3OZD358J)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78393-010-01	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/01/2020	

Labeler - Biocleansingcompany Co., Ltd (695538805)

Registrant - Biocleansingcompany Co., Ltd (695538805)

Establishment

Name	Address	ID/FEI	Business Operations
DongbangCosmetics Co., ltd.		694452564	manufacture(78393-010)

Revised: 5/2020

Biocleansingcompany Co., Ltd