

EPU HAND SANITIZER 946ML- ethyle alcohol gel
E.CIS Cosmetic 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.

EPU Hand Sanitizer 946ml

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic

E-Pu

**INSTANT
HAND SANITIZER**
70% Ethyl Alcohol as recommended

Help reduce bacteria on the skin

- For all day use
- Non-Dry, Non-Sticky
- Deep Moisturizing

32 fl.oz. (946 mL)

Drug Facts	
Active Ingredient(s) Ethyl Alcohol 70% v/v	Purpose Antiseptic
Use(s) Health care personnel hand rub to help reduce bacteria that potentially can cause disease.	
Warnings For external use only. Flammable. Keep away from heat or flame.	
Do not use <input type="checkbox"/> in children less than 2 months of age <input type="checkbox"/> 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.	
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Other Information <input type="checkbox"/> Store between 15-30°C (59-86°F) <input type="checkbox"/> Avoid freezing and excessive heat above 40°C (104°F)	
Inactive Ingredients Water, Glycerin, Carbomer, Anisomethyl Propanol, Mentha Arvensis Leaf Oil, Tocopheryl Acetate, Panthenol, Sodium Hyaluronate, Aloe Barbadosensis Leaf Extract	
Questions or Comments? 1-800-819-4159	

Manufactured by ECIS COSMETIC CO., LTD.
179, Gongdan 1-dong, Sihyeung-si,
Gyeonggi-do, Republic of Korea

MADE IN KOREA

8 809597 416469

Purpose

Antiseptic, Hand Sanitizer

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Questions or Comments? 1-800-619-4159	

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179, Gongdan 1-daeon, Sicheung-st,
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Use

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Stop use and ask a doctor

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Package Label - Principal Display Panel

EPU INSTANT HAND SANITIZER
70% Ethyl Alcohol as recommended
Help Reduce Bacteria On The Skin.
For all day use

Non-Dry, Non-Sticky

Deep Moisturizing,
32 fl.oz (946ml)

EPU HAND SANITIZER 946ML

ethyle alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73862-217
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	

PANTHENOL (UNII: WV9CM0O67Z)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73862-217-11	12 in 1 CASE	03/30/2020	
1	NDC:73862-217-10	946 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - E.CIS Cosmetic Co.,Ltd (689846270)

Registrant - E.CIS Cosmetic Co.,Ltd (689846270)

Establishment

Name	Address	ID/FEI	Business Operations
E.CIS Cosmetic Co.,Ltd		689846270	manufacture(73862-217)

Revised: 6/2020

E.CIS Cosmetic Co.,Ltd